A fluorescent nanosensor paint detects dopamine release at axonal varicosities with high spatiotemporal resolution

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The neurotransmitter dopamine (DA) controls multiple behaviors and is perturbed in several major brain diseases. DA is released from large populations of specialized structures called axon varicosities. Determining the DA release mechanisms at such varicosities is essential for a detailed understanding of DA biology and pathobiology but has been limited by the low spatial resolution of DA detection methods. We used a near-infrared fluorescent DA sensor, AndromeDA, to detect DA secretion from cultured murine dopaminergic neurons with high spatial and temporal resolution. We found that AndromeDA detects discrete DA release events and extracellular DA diffusion and observed that DA release varies across varicosities. To systematically detect DA release hotspots, we developed a machine learning–based analysis tool. AndromeDA permitted the simultaneous visualization of DA release for up to 100 dopaminergic varicosities, showing that DA release hotspots are heterogeneous and occur at only ~17% of all varicosities, indicating that many varicosities are functionally silent. Using AndromeDA, we determined that DA release requires Munc13-type vesicle priming proteins, validating the utility of AndromeDA as a tool to study the molecular and cellular mechanism of DA secretion.

dopamine | neurotransmission | chemical signaling | sensors | imaging

The modulatory neurotransmitter dopamine (DA) controls a range of brain processes, from motor function to reward and motivation (1, 2). Aberrant DA signaling, on the other hand, is causally involved in multiple brain disorders, including Parkinson’s disease, schizophrenia, and addiction (3–5). Despite this eclectic functional relevance, little is known about the molecular and cellular mechanisms that mediate and control DA release.

Fast neurotransmitters, such as glutamate, operate with high spatial and temporal accuracy. Our understanding of fast neurotransmission has been greatly advanced through methods that allow the study of fast neurotransmission with high precision, including imaging of live presynaptic boutons. Contrasting with fast neurotransmitters, dopaminergic (DAergic) neurons form multiple types of axonal release sites (6, 7) called varicosities, many of which are nonsynaptic and even release DA from somata and dendrites (8). DA typically diffuses considerable distances from release sites, and signal transduction by metabotropic DA receptors is much slower than signaling by fast-acting neurotransmitters via ionotropic receptors.

The distinct mode of DAergic signaling is of key functional significance and hence likely underpinned by cellular and molecular specializations of its release machinery that differ from fast-acting systems, e.g., with regard to the organization, priming, fusion, and dynamics of synaptic vesicles (SVs). Indeed, the few known examples of proteins involved in the control of DA release indicate that DAergic varicosities not only differ from typical fast-acting synapses but are also very heterogeneous in structure, function, and molecular composition (6, 9–12). To assess the function of this heterogeneous system, DA release must be examined with a spatial resolution that discriminates individual varicosities. Common electrochemical DA detection methods cannot resolve large numbers of varicosities in parallel due to their low spatial resolution, determined by electrode number, type, and position (13–15). Furthermore, genetically encoded DA sensors (16–19) have not yet been used to examine DA release from discrete varicosities with subcellular precision, though recent data indicate DA detection at discrete dendritic release sites in substantia nigra (20). We thus developed a DA detection method based on single-walled carbon nanotubes (SWCNTs), which emit photostable (21) structure-dependent fluorescence in near infrared (NIR) (870 nm to 2,400 nm) ideal for biological imaging (22). Functionalized SWCNTs are highly effective nanosensors (23–29), and they have been used as high-affinity DA sensors that detect DA release from cell lines (30) or in brain slices (31). However, the potential gain in spatial resolution provided by nanosensor-based approaches, e.g., to examine the spatiotemporal complexity of DA release, has not been explored, mainly because other methods are still required to gain in spatial resolution provided by nanosensor-based approaches.

Significance

The neurotransmitter dopamine controls normal behavior and dopaminergic dysfunction is prevalent in multiple brain diseases. To reach a detailed understanding of how dopamine release and signaling are regulated at the subcellular level, we developed a near infrared fluorescent dopamine nanosensor ‘paint’ (AndromeDA) to directly image dopamine release and its spatiotemporal characteristics. With AndromeDA, we can ascribe discrete DA release events to defined axonal varicosities, directly assess the heterogeneity of DA release events across such release sites, and determine the molecular components of the DA release machinery. AndromeDA thus provides a new method for gaining fundamental insights into the core mechanisms of dopamine release, which with greatly benefit our knowledge of dopamine biology and pathobiology.
primary DAergic neurons have to be differentiated in culture over several weeks for such purposes and because cultivation and adhesion on a nanosensor layer affect sensor sensitivity and cell viability. By using a sensor paint on top of mature neuronal cultures, we circumvented these challenges and now present a SWCNT-based DA detection method—adsorbed nanosensors detecting release of dopamine (AndromeDA)—that optically discriminates DA release from large numbers of neuronal release sites simultaneously.

Results

AndromeDA Detects Distinct, Heterogeneous DA Release Events from Large Populations of Discrete Varicosities. NIR fluorescent SWCNTs were functionalized with single-stranded deoxyribonucleic acid, consistent of 10 thymine and guanine repeats ([GT]10-ssDNA) to create nanosensors that increase in fluorescence emission with reversible DA binding (SI Appendix, Figs. S1 and S2). To generate a two-dimensional (2D) nanosensor layer capable of high spatiotemporal resolution, we applied a concentrated solution of SWCNT-(GT)10-ssDNA nanosensors to poly-L-lysine (PLL)—coated coverslips, resulting in a painted nanosensor surface on the glass, i.e., AndromeDA. NIR fluorescence imaging used to characterize the responsiveness of AndromeDA to DA showed fluorescence increases of up to 40% upon reversible DA binding in a concentration-dependent manner, with an half maximal effective concentration (EC50) of 299 nM (SI Appendix, Fig. S2).

To study DA release from varicosities, we initially tried culturing neurons directly on nanosensor-coated coverslips, similar to previous experiments with PC12 cells (30), but this approach compromised neuron viability and nanosensor sensitivity. Instead, cocultured neurons from mouse hippocampus and ventral midbrain were grown on coverslips for at least 6 wk, and nanosensors were applied immediately prior to imaging. This resulted in a uniform layer on the glass surrounding the cells (Fig. 1A), as nanosensors did not adhere to cells (SI Appendix, Fig. S3). A schematic of a single nanosensor is shown in Fig. 1B. Atomic force microscopy (AFM) demonstrated the shape of single nanosensors and the uniformity of the nanosensor layer on a PLL-coated surface (Fig. 1C). Using neurons from neonatal mice, in which most DAergic neurons express enhanced green fluorescent protein (EGFP) under the tyrosine hydroxylase (TH) promoter (TH-EGFP mice) (SI Appendix, Fig. S4 A–C), we identified DAergic axons surrounded by AndromeDA using a custom-built optical setup to simultaneously image light in the visible and NIR range (SI Appendix, Fig. S5). By visualizing DAergic varicosities and the location of DA in time and space, AndromeDA allows the detection and separate analysis of DA release and diffusion (Fig. 1D).

To validate AndromeDA in a biologically relevant setting, ventral midbrain neurons were stimulated to evoke DA release (Fig. 2). Electrical field stimulation (200 pulses, 20 Hz), which induces action potential firing (32), caused a rapid increase in the extracellular NIR fluorescence of neuronal cultures, corresponding to AndromeDA activation by evoked DA release and diffusion (Fig. 2, untreated neurons). Chemical depolarization of neurons with 90 mM KCl resulted in a relatively strong and sustained increase in extracellular NIR fluorescence (Fig. 2A and B). fluorescence quantification shown in SI Appendix, Fig. S6 A–D). AndromeDA activation throughout the extracellular space was widespread, reflecting the diffusion of DA away from release sites and the high sensitivity of the sensors (Fig. 2 A and B). To account for possible differences in the density of nanosensor layers between experiments, 100 μM DA was added to the extracellular space of the neurons, and the intracellular fluorescence was measured with AndromeDA.
the samples at the end of each experiment to maximally activate the nanosensor layer (Fig. 2 A and B), and fluorescence values from electrical or chemical stimulation were then expressed as a proportion of this maximal value. Some variation between experiments in the baseline (untreated) peak responses evoked by electrical or chemical stimulation was evident (Fig. 2 D and F and SI Appendix, Fig. S6 B and D), which can be attributed to variation in DA release between culture batches.

Electrical stimulation did not cause any fluorescence increase in the absence of DAergic neurons (SI Appendix, Fig. S7 A and C), indicating that AndromeDA specifically detects secreted DA. KCl application in the absence of neurons caused a modest fluorescence increase (mean ± standard error of mean [SEM]) for each stimulus method. Reserpine decreases the evoked peak AndromeDA fluorescence compared with untreated neurons. n = 10 untreated experiments per condition. Average AndromeDA fluorescence over time for neurons stimulated at 20 Hz with and without treatment with reserpine. Peak fluorescence (mean ± standard error of the mean [SEM]) for each stimulus method. L-DOPA increases the evoked peak AndromeDA fluorescence compared with untreated neurons. n = 5 (untreated) and 11 (L-DOPA) experiments (scale bars, 10 μm). In line graphs, solid lines represent mean values, and shaded areas represent SEM. Statistical comparisons used two-tailed Welch’s t test. ***p < 0.001. Red arrows indicate the direction of changes in the AndromeDA signal in treated neurons compared to untreated controls. Note that the intensities of DA-treated images in A and B are above the displayed fluorescence scale. F = Fluorescence intensity.

Fig. 2. Imaging of DA release and diffusion. (A) Images of normalized AndromeDA fluorescence (Left to Right) prior to stimulation (No stim), during 20-Hz electrical stimulation, during 90 mM KCl stimulation, and during application of 100 μM DA reveal DA release in untreated neurons (Upper). Treatment with reserpine (1 μM, 90 min) decreases stimulus-dependent activation of AndromeDA (Lower). AndromeDA fluorescence was quantified at ROIs centered on EGFP-positive varicosities (examples shown as red circles). (B) Neurons were either untreated or pretreated with L-DOPA (100 μM, 45 min), and experiments were performed as in A. (C) Average AndromeDA fluorescence over time for neurons stimulated at 20 Hz with and without treatment with reserpine. (D) Peak fluorescence (mean ± standard error of the mean [SEM]) for each stimulus method. Reserpine decreases the evoked peak AndromeDA fluorescence compared with untreated neurons. n = 10 untreated experiments per condition. (E) Average AndromeDA fluorescence over time for neurons stimulated at 20 Hz with and without L-DOPA pretreatment. (F) Peak fluorescence (mean ± SEM) for each stimulus method. L-DOPA increases the evoked peak AndromeDA fluorescence compared with untreated neurons. n = 5 (untreated) and 11 (L-DOPA) experiments (scale bars, 10 μm). In line graphs, solid lines represent mean values, and shaded areas represent SEM. Statistical comparisons used two-tailed Welch’s t test. ***p < 0.001. Red arrows indicate the direction of changes in the AndromeDA signal in treated neurons compared to untreated controls. Note that the intensities of DA-treated images in A and B are above the displayed fluorescence scale. F = Fluorescence intensity.

the hippocampal neurons were preincubated in L-3,4-dihydroxyphenylalanine (L-DOPA) (SI Appendix, Fig. S7E).

To further validate that AndromeDA specifically detects DA release, we assessed neurons treated with the vesicular monoamine transporter (VMAT) inhibitor reserpine (34), which depletes DA from SVs and abolishes DA release with prolonged incubation (35). Reserpine (1 μM, 90 min) decreased peak AndromeDA fluorescence at regions of interest (ROIs) around varicosities after electrical (88% reduction in peak fluorescence) and KCl (~68% reduction) stimulation (Fig. 2 A, C, and D and SI Appendix, Figs. S6 A and B and S8). Complementing this, we examined the effect of L-DOPA, which increases the DA content of SVs in DAergic neurons 10-fold (36). L-DOPA (100 μM, 45 min) significantly increased peak AndromeDA activation at ROIs around varicosities when neurons were depolarized (Fig. 2 B, E, and F and SI Appendix, Figs. S6 C and D and S8). Thus, altering vesicular DA content correspondingly altered the AndromeDA signal, confirming its DA specificity.

The detection of DA diffusion in the extracellular space reflects cumulative DA release from many varicosities and does not facilitate the assessment discrete varicosities. However, in...
addition to more diffuse AndromeDA activation, we observed brief, localized hotspots of increased fluorescence, generally during neuronal stimulation, above the surrounding fluorescence and generally located adjacent to varicosities (Fig. 3 A–C and Video S1). DA is stored in SVs in varicosities at very high concentrations, and DA release from a single varicosity causes a localized high DA concentration that rapidly dissipates as secreted DA diffuses into the surrounding extracellular space (8). Thus, fluorescence hotspots corresponded to the high, local DA concentration that occurs upon DA release events. Manual identification and analysis of hotspots lacked the necessary sensitivity to reliably detect small events. We thus developed a Fiji plug-in that uses machine learning to identify hotspots from NIR videos, the Dopamine Recognition Tool (DART). Imaging EGFP and NIR fluorescence in parallel, we observed up to 100 DAergic varicosities and 50 DART-identified hotspots.
within a single imaging field (12,800 μm²; Fig. 3 A and B). Hotspots were generally brief, usually between three and five imaging frames in duration (200 to 333 ms). DART automatically defined the region within each detected hotspot as a ROI (Fig. 3B). To isolate the fluorescence corresponding specifically to each hotspot, the mean response of the extracellular space was subtracted from the mean fluorescence within an identified hotspot, thus correcting for AndromeDA activation by DA diffusing from other release sites. After correction, hotspot traces demonstrated a characteristically fast rising phase and a slower decaying phase (Fig. 3C). While most varicosities did not exhibit nearby hotspots, varicosities were observed exhibiting single or multiple nearby hotspots during 20-Hz stimulation (Fig. 3B). Hotspots were also induced by KCl stimulation (SI Appendix, Fig. S9 and Video S2).

The areas and peak intensities of hotspots were highly variable, indicating a striking heterogeneity in DA release events at the level of single varicosities (Fig. 4 A and B). To further validate that hotspots represent discrete DA release events, we performed additional spatiotemporal analyses of images. An example image showing DAergic axons and the ROIs occupied by hotspots observed while imaging these axons is shown in SI Appendix, Fig. S10, demonstrating hotspot clustering close to axons. As DA release occurs at varicosities upon action potential firing, we hypothesized that hotspots should occur close to varicosities and be triggered by electrical stimulation. To verify the close spatial association of hotspots to DAergic varicosities, we took advantage of our high spatial resolution and measured the distance from hotspot centers to the center of the nearest varicosity as identified in the corresponding EGFP image (Fig. 4 C and D and SI Appendix, Fig. S11A). The median hotspot-varicosity distance was 1.2 μm, with ~86% of hotspots within a 3-μm radius of a varicosity. Remaining hotspots are likely due to DA release from EGFP-negative DAergic neurons (~30% of all TH-positive neurons; SI Appendix, Fig. S4). We also measured hotspot-varicosity distance using NIR and EGFP images from noncorresponding experiments, providing a randomized negative control (Fig. 4E and SI Appendix, Fig. S11B). In this case, hotspot-varicosity distances were significantly larger than corresponding NIR and EGFP images, verifying that hotspots generally occur close to varicosities (Fig. 4E and SI Appendix, Fig. S11B).

To verify the dependency of hotspots on neuronal depolarization, we first examined whether they corresponded in time with neuronal stimulation. Most hotspots (~82%) occurred during electrical stimulation, ~4% occurred prior to stimulation, and ~14% occurred after stimulation had ceased (Fig. 4F). Thus, a large majority of hotspots occurred during exogenous neuronal stimulation (Fig. 4G), which we referred to as evoked hotspots. Evoked and nonevoked hotspots were quantitatively similar (Fig. 4H). In samples stimulated with KCl, hotspots corresponded in time with the application of KCl (SI Appendix, Fig. S11C). We next considered the cause of nonevoked hotspots. They were not the result of glutamatergic signaling in the culture, which could activate DAergic neurons, as experiments were performed in the presence of glutamate receptor antagonists. We tested the effect of 300 nM tetrodotoxin (TTX), which blocks action potentials, in the absence of electrical stimulation. Hotspots were not detected in the presence of TTX, but nonevoked hotspots appeared upon TTX removal (Fig. 4F and SI Appendix, Fig. S12 A–C). After TTX removal, neurons exhibited normal evoked DA release (SI Appendix, Fig. S13). These data indicate that nonevoked hotspots are caused by the spontaneous electrical activity of DAergic neurons, so that all hotspots are evoked by neuronal firing and subsequent electrically evoked DA release events. To quantify when each hotspot occurred in time, we used the time at which the hotspot peak intensity was reached. A cumulative frequency histogram plotting of the timing of hotspot occurrence in L-DOPA–treated neurons showed that most hotspots appeared within the first 4 s (80 pulses) of electrical stimulation (SI Appendix, Fig. S14), with hotspots becoming less frequent as 20–Hz stimulation continued. This decrease in detectable discrete DA release events with repeated stimulation may indicate a depletion of DA at varicosities. These data therefore demonstrate the utility of AndromeDA in studying in DA release dynamics across populations of individual varicosities.

Together, these data demonstrate that AndromeDA can detect and discriminate i) summed DA release in the extracellular space, representing diffusing DA from many local release sites (Fig. 2 and Video S2), and ii) discrete DA release events at single varicosities (Figs. 3 and 4). Importantly, AndromeDA detects many release events from large numbers of varicosities with subcellular resolution, a significant methodological advance for studying DA release from large numbers of sites in parallel.

To assess whether DA dynamics in our system are affected by the dopamine transporter (DAT), we simulated the contributions of DA diffusion and uptake to extracellular DA levels (SI Appendix, DA Diffusion Simulation and Figs. S15 and S16) (37). The simulation shows that the open 2.5-dimensional geometry and the fast on rate of DA binding by nanosensors (30, 38, 39) render the impact of DAT activity on our data in this experimental setting negligible.

AndromeDA Differentiates Active and Silent Varicosities. DAergic varicosities are filled with SVs (40). Interestingly, previous studies have indicated that DA release occurs only in a subpopulation of varicosities (6, 9), although DA release was measured indirectly with fluorescence-based exocytosis assays. To investigate this using a direct method of DA detection, we examined the proportion of total varicosities that exhibited spatially associated AndromeDA hotspots, which indicate local DA release events. In our analyses, the median hotspot area was 5.5 μm², and assuming that hotspots are approximately circular, this gives a diameter of 2.6 μm. We thus set a threshold distance of ≤3 μm for defining that a given hotspot was spatially associated with a varicosity and examined the proportion of varicosities that show discrete DA release events. In untreated neurons, hotspots were relatively infrequent, observed in only 25% of all experiments and adjacent to 13% of all varicosities on average (Fig. 4I). L-DOPA treatment increased hotspot frequency, with hotspots observed in all experiments and adjacent to 17% of all varicosities. L-DOPA increases the quanta size of DA release (35, 36), and thus the increase in hotspot frequency following L-DOPA treatment is likely due to increased quanta size, allowing discrete release events to register above background. While L-DOPA treatment increased hotspot frequency (Fig. 4I), it did not alter hotspot peak intensity or area (SI Appendix, Fig. S9 A and B). Similar to electrical stimulation data, only a subpopulation of varicosities showed hotspots upon KCl-mediated depolarization (SI Appendix, Fig. S17). These data support the notion that DA is released from only a subset of active varicosities by demonstrating the phenomenon via direct DA detection.

Local AndromeDA Responses Diminish with Distance from DAergic Axons. As axons are the source of secreted DA, AndromeDA signals are expected to be stronger closer to axons. Exploiting
Fig. 4. Hotspots are heterogeneous sites of activity-dependent DA secretion. Hotspots observed in untreated (black dots) and L-DOPA-treated (100 μM, 45 min, green dots) electrically stimulated neurons are shown. (A) Area of activated AndromeDA within single hotspots. (B) Peak fluorescence within single hotspots. (C) Scheme illustrating how distance and hotspot area are defined. (D) Distances between the center of individual hotspots and the nearest varicosity. (E) Varicosity-hotspot distance was larger when noncorresponding EGFP images were used (randomized). (F) Time points at which individual hotspots occur. Electrical stimulation from 10 to 20 s is highlighted (red). (G) Hotspots occur primarily during electrical stimulation. (H) Hotspot peak fluorescence is no different from that of hotspots observed in the absence of stimulation. Only hotspots from neurons treated with L-DOPA were analyzed due to the large number of hotspots available for analysis compared with untreated neurons. (I) Hotspots are the result of spontaneous neuronal firing. The representative trace shows AndromeDA fluorescence over time from an EGFP-positive varicosity within the ROI indicated (white arrowhead). During TTX application, no hotspots are evident, but washout of TTX results in the appearance of hotspots (black arrowheads). (J) Only a subpopulation of varicosities exhibits closely adjacent (<3 μm) hotspots. L-DOPA treatment causes a significant increase in the proportion of varicosities with adjacent hotspots. In E, dots represent the median of each experiment, and column height is the mean of all experiments (±SEM). In G, H, and J, dots represent the median of each experiment, and column height is the median of all experiments. In A, B, D, and F, hotspots are organized on the y axis into individual experiments (lines), and the hatched area shows the frequency distribution of all hotspots. In A–H, data represent 68 hotspots from n = 4 experiments (untreated) and 110 hotspots from n = 10 experiments (L-DOPA). Data were compared using either two-tailed Welch’s t test (E) or the Mann-Whitney test (G, H, and J). *P < 0.05, **P < 0.01. Hotspot traces were generated by subtracting the mean signal of the extracellular space (ΔF/Fl) from the mean signal within the hotspot ROI (ΔF/Fl) for each time point (scale bar, 10 μm).

the spatial resolution of AndromeDA, we quantified the local change in fluorescence during neuronal stimulation at different distances from axons (Fig. 5A). To do this, the background fluorescence due to DA diffusion in a region distant from the axons was subtracted from the fluorescence within each ROI around the axon, thus giving the local change in fluorescence over time. ROIs close to axons showed the greatest relative change in fluorescence, with the amplitudes of stimulus-evoked fluorescence signals decreasing with distance from the axon (Fig. 5B). These data highlight the rapid rise in relative fluorescence adjacent to axons that occurs with electrical stimulation, followed by a decline in fluorescence that precedes the cessation of electrical stimulation. Together with our observation that hotspots decrease in frequency during 20-Hz stimulation (SI Appendix, Fig. S14), these data indicate that DA release occurs rapidly following the onset of stimulation and decreases with time, indicating DA depletion at varicosities. In line with this, DA depletion has been observed using amperometry recordings in striatal slices (10–12), and repeated stimulation of DAergic neurons at 20 Hz results in depletion of SVs when assayed using synaptophosphorylin (41).
neurons lacking Munc13-1 and Munc13-2 (Munc13 double knock-out, DKO) (Fig. 6A). The breeding scheme we used also provided neurons lacking only Munc13-1 (heterozygous for Munc13-2 expression) and ones that were heterozygous for the Munc13-1 deletion so that we were able to examine the impact of reducing Munc13-1 expression specifically (SI Appendix, Fig. S21). Electrically stimulated Munc13 DKO neurons showed no signal above noise level of fluorescence imaging and no hot-spots, as opposed to neurons from littermate control mice (Fig. 6 A–C and SI Appendix, Fig. S21 A and B). Of the genotypes analyzed, only Munc13 DKO resulted in a significant reduction in AndromeDA activation evoked by 20-Hz stimulation (SI Appendix, Fig. S21 C–E). These data show that Munc13 proteins are required for evoked DA release and demonstrate the utility of AndromeDA in studying the molecular and cellular underpinnings of DA secretion.

Discussion

Optical DA sensing technology has recently made major progress, with the development of SWCNT-based DA nanosensors (23, 24, 30, 38) rapidly followed by genetically encoded fluorescent DA sensors (17, 18, 53, 54). Compared with electrophysiology, these methods provide many advantages, particularly in that they allow for an unbiased approach to study DA signals in situ.

Validation of AndromeDA in SV Priming-Deficient Neurons.

SV priming by Munc13 proteins, which dock SVs to the presynaptic membrane and render them fusion competent (42), is absolutely required in essentially every synapse type tested so far, with Munc13 deletion causing near-complete or complete loss of neurotransmitter release (43, 44). To determine whether this also applies to synaptic DA secretion, we used AndromeDA to assess the Munc13 dependence of DA release from cultured neurons and also performed a collaborative analysis of Munc13 function in DAergic synapses using amperometry in striatal slices (12).

We used immunolabeling to assess Munc13 expression in DAergic neurons. The dominant Munc13-1 isoform was readily detected at DAergic active zones as defined by anti-bassoon colabeling (SI Appendix, Fig. S18). Due to the lack of effective antibodies against Munc13-2 and Munc13-3, we analyzed previously characterized knock-in (KI) mice that express Munc13-2–enhanced yellow fluorescent protein (EYFP) or Munc13-3–EGFP fusion proteins from the endogenous genetic loci (45). EYFP or EGFP immunodetection revealed no Munc13-2–EYFP–positive puncta in TH-positive DAergic axons, whereas cultured hippocampal neurons, as Munc13-2–positive controls (46, 47), showed clear signals (SI Appendix, Fig. S19). Similar observations were made with D1 and D2 DAergic Munc13-3–EGFP neurons and cultured cerebellar granule neurons as positive controls (48, 49) (SI Appendix, Fig. S20). These data show that Munc13-1 is the dominant Munc13 in cultured murine DAergic neurons and that Munc13-2 and Munc13-3 are absent or expressed at very low levels, consistent with findings showing predominant Munc13-3 expression in hindbrain (48, 50, 51). On the other hand, Munc13-2 mRNA has been shown to be expressed in at least some mouse DAergic neurons (52). Thus, to maximize the likelihood of detecting effects of Munc13 deletion on DA release, we studied TH-EGFP

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**Fig. 5.** AndromeDA activation decreases with distance from axons. (A) To examine how the local changes in AndromeDA fluorescence were influenced by distance from axons, AndromeDA fluorescence was quantified at different distances from axons. The first ROI was defined by thresholding the EGFP signal (white) of the DAergic axon and tracing it (ROI 1). The line defining the boundary of this ROI was then moved outwards by 2 µm from the axon to create ROI 2. This process was repeated to create concentric ROIs, as indicated in green and identified using numbers. (B) The change in the local normalized fluorescence within each ROI compared with a distant region containing no axons was quantified (Materials and Methods). This gave the local increase in AndromeDA fluorescence within each ROI. ROI 1, which is closest to the axon, shows a rapid, high-amplitude increase in fluorescence with electrical stimulation. As the local changes in AndromeDA fluorescence are analyzed farther from the axon, the amplitude of the response is decreased. This demonstrates that local AndromeDA activation is greatest immediately adjacent to axons and that the local DA level rises rapidly with stimulation. Traces in B represent average normalized fluorescence over time. ROI fluorescence values, which are colored according to the ROI they represent, were generated by subtracting the mean normalized (ΔF/F0) signal of a selection of the extracellular space from the mean signal within the ROI (ΔF/F0) for each time point. Data represent n = 7 independent coverslips.

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**Fig. 6.** Munc13-1 and Munc13-2 are required for evoked DA release. (A) Representative images of AndromeDA-painted ventral midbrain TH-EGFP neurons lacking expression of Munc13-1 and Munc13-2 (Munc13 DKO) compared with neurons from littermate control mice. Images show normalized AndromeDA fluorescence prior to stimulation, during 20-Hz electrical field stimulation, and during application of 100 µM DA. ROIs were defined around varicosities of EGFP-positive axons (red circles). (B) AndromeDA signal within ROIs over time for control and Munc13 DKO neurons with electrical stimulation. Munc13 DKO neurons exhibit no AndromeDA activation in response to electrical stimulation (red arrow indicates the loss of the AndromeDA signal in Munc13 DKO neurons). (C) Average maximal fluorescence peak during 20-Hz electrical field stimulation in control and Munc13 DKO neurons. For control neurons, n = 6, and for Munc13 DKO neurons, n = 4, where n represents independent experiments. In B, solid lines represent the mean and shaded areas represent SEM. In C, mean ± SEM is shown. Data were compared using two-tailed Welch’s t test. *P < 0.05. Note that the intensities of DA-treated images in A are above the displayed fluorescence scale.
higher spatial resolution and multiplexing (54). While immobi-
lized nanosensors had previously been used to detect DA release
from PC12 cells (30), AndromeDA is compatible with neurons,
which are of far greater biological relevance. Moreover, our
combination of an immobilized sensor layer added to a mature
neuron culture of EGFP-positive DAergic neurons allows the
detection of DA with high spatial resolution and the correlation
of DA release with subcellular structures. Nanosensors have
been used previously with brain slices, but no attempt was
made to ascribe DA release to particular subcellular structures
(31). However, studying discrete axonal varicosities is essential
to understand their functional properties and control by pre-
synaptic proteins, whose levels vary across varicosity popula-
tions (6, 10–12). Compared with brain tissue, the low density
of varicosities in neuronal culture makes the routine analysis
of single varicosities and their function feasible. Such analysis
of discrete DA release events in neuronal cultures has been possible
with amperometry but has been restricted to a single varico-
sity at a time (35). Instead, AndromeDA permits the analysis
of large populations of varicosities in parallel. The DA affinity
of AndromeDA is similar to that of DA receptors (55), thus
providing a spatiotemporal readout of DA in the biologically
relevant concentration range. Additionally, the ability to tune
the properties of SWCNT-based DA nanosensors expands their
potential utility in addressing different aspects of DA release
(38). AndromeDA can be applied to mature neuronal cultures with no detectable impact on cell viability. As such, Androm-
eDA is readily applicable to in vitro cellular models of human
disease in which varicosity function and DA release are sus-
ppected to be abnormal, such as neurons derived from Parkinson’s
disease patients. Our development of a machine learning–based tool, DART, for the detection of DA release events also greatly enhances the ability of our method to iden-
tify discrete sites of release.

Multiple tools have been developed to study DA exocytosis at the level of single varicosities, including styryl dyes (6), a false fluorescent neurotransmitter (9), and pHluorin-based
probes (56). While these have provided important insights into
diagnostic biology, they do not report DA release itself or the move-
ment of DA after release. Still, such exocytosis assays in neuronal
culture (6) and striatal slices (9) indicated high functional heterogeneity of DAergic varicosities, with many appearing to be functionally silent and the remaining active ones exhibiting
different functional properties. Using AndromeDA, we now
provide direct proof that most DAergic varicosities do not release DA. We found that after treating neurons with L-DOPA to boost hotspot detection, only ~17% of all varicosi-
ties exhibited hotspots. Validating our method, this is strikingly
similar to the proportion of DAergic varicosities that exhibit
electrically evoked release of the false fluorescent neurotransmitter
FFN200 in the striatum (~17%) (9). In essence, our data based on the direct detection of DA confirm that silent varicosi-
ties are an inherent property of DAergic neurons. We observed
no morphological differences between varicosities that exhibit
by nearby adjacent hotspots and those that lack them. Given
the sparse distribution of molecular components necessary for DA
release (6, 10–12), we presume that differential DA release
between varicosities is determined by differential configuration
of the molecular SV fusion machinery in these varicosities.
Moreover, individual AndromeDA hotspots were entirely
driven by action potentials and varied considerably in their
timing, amplitude, and area, directly demonstrating the heter-
genocity of DA release. Our data further show that spontaneous
action potential firing in DAergic neurons drives sufficient DA
release to be visualized as discrete hotspots. Such spontaneous
pacemaker firing is an inherent property of DAergic neurons
(8, 57). Importantly, these data could only be provided by a
method capable of examining DA release events from large
numbers of varicosities in parallel, highlighting the utility of
AndromeDA for interrogating fundamental aspects of DA
release. The biological role of the functional heterogeneity of
DAergic varicosities is unknown, but it is likely that the spatial
and temporal patterns of DA release arising due to DA release
from a functionally diverse varicosity population are important
in DA signaling and information processing.

Stimulation of DAergic neurons resulted in a steep rise in extracellular AndromeDA fluorescence, consistent with rapid
evoked DA release, and hotspots were observed primarily
within the first few seconds of repeated stimulation. However,
the rise in extracellular fluorescence around axons (Figs. 2 and
5) occurs faster than the emergence of hotspots in time, which
do not begin to appear until after 1 s of stimulation (SI
Appendix, Fig. S14). This indicates that hotspots do not repres-
ent every DA release event. Hotspots cross the threshold of
detection presumably due to more DA being released, as
highlighted by the increase in hotspots seen when neurons are
handled with L-DOPA. DART is highly stringent in hotspot
detection, so that some bona fide DA release events may not be
detected as a result. The reason why certain release events are
large enough to be detected as hotspots is unclear. These events
may correspond to several release events occurring within a
short time frame or to so-called complex DA release events,
which result in more DA release from SVs (58). It is unlikely
that hotspots correspond to DA release from large dense-core
vesicles, given that DA release occurs almost entirely from SVs
(35, 58, 59). Moreover, the AndromeDA DA optical detection
method reports discrete DA release events with subcellular reso-
novation, allowing a characterization of the functional properties
of individual DAergic varicosities.

The speed and fidelity of signaling by fast-acting neurotrans-
mitters is ensured by a specialized molecular machinery that
regulates synaptic exocytosis, endocytosis, and SV dynamics.
Given the unique features of DAergic neurotransmission, it has
been assumed that specific molecular determinants define the
DA release machinery. Indeed, recent studies showed that
synaptotagmin-1 and Rab3-interacting molecules (RIMs) are
required for DA release (10, 11), as is the case for the fast-
acting neurotransmitter glutamate, while the active zone
protein ELKS is not required (10). We found that the genetic
deletion of Munc13-family SV priming proteins abolishes
evoked DA release. In accord with the well-defined, essential
role of Munc13s in vesicle priming in multiple other neuronal
and endocrine systems, this documents an essential SV priming
function of Munc13s in DAergic synapses. Similar observations
were made in a parallel study in striatal slices, leading to the
conclusion that DAergic synapses require SV priming and
release-site scaffolding by Munc13s, RIMs, and Liprins,
whereas multiple other presynaptic components are dispensable
(12). Most importantly, in the context of the method-
development focus of our study, our present analysis of
Munc13-deficient neurons documents the major utility of
AndromeDA as a tool to study the molecular and cellular
underpinnings of DA secretion. In the future, it will facilitate
detailed analysis of how molecules shape the heterogeneity of
DA release across populations of individual varicosities.

By allowing high-resolution imaging of DA release from
many release sites, AndromeDA paves the way for a deeper
understanding of DAergic signaling at a subcellular level.
AndromeDA represents a tool that will provide fundamental insights into the basic properties of DA secretion, into the heterogeneity and plasticity of DA release, and into the detailed molecular processes that shape DA release. In principle, our painting approach to generating 2D sensor layers such as AndromeDA can also be applied to other neurotransmitter nanosensors, such as our recently developed serotonin sensor (25). Such high-resolution approaches are essential to decipher the complex physiological and pathophysiological functions of DA and other modulatory neurotransmitters.

Materials and Methods

Animals. Mouse lines, genotyping, housing, and ethics are provided in SI Appendix, Extended Materials and Methods.

Preparation of Primary Neuron Cocultures. To prepare neurons intended for AndromeDA imaging, a coculture protocol was established by combining dissociated hippocampal primary neurons with ventral midbrain neurons in a single-cell culture system. Hippocampal neurons, cultured 5 to 7 days prior to the addition of ventral midbrain neurons, promoted DAergic neuron survival for at least 6 wk. This period of survival is comparable to the classical ventral midbrain neuronal preparation grown on glial cells (60-62), and it provided the areas of cell-free glass necessary for AndromeDA application.

For neuron culture, glass coverslips (25 mm, #1 Menzel Gl) were coated with 0.0008% (wt/vol) PLL (Sigma-Aldrich, P4707) in phosphate-buffered saline (PBS). Hippocampal neurons were then prepared as previously described (63), seeded at 1.5 × 10^4 neurons per coverslip in neuron medium (2% fetal bovine serum [FBS], 2% B-27 [Gibco, 17504-044], penicillin [200 U/mL streptomycin [200 μg/mL] [Gibco, 15140-130], 2 mM GlutaMAX [Gibco, 35050-038] in Neurobasal-A), and incubated at 37°C with 5% CO2. The next day, the medium was replaced with neuron medium supplemented with 67 μg/mL 5-fluoro-oxyuridine (Sigma-Aldrich) and 165 μg/mL uridine (Sigma-Aldrich) to inhibit cell proliferation.

After 7 d, ventral midbrain neurons were prepared using a protocol based on two methods described previously (6, 62). Ventral midbrain neurons of mice at postnatal day 0 (P0) were dissected and digested with papain at 34°C for 45 min, followed by 15 min of incubation in trituration medium (10% FBS in Neurobasal-A medium [NBA]). After trituration, cell suspensions were centrifuged (5 min, 1,500 rpm). Cells were resuspended in neuron medium and seeded (1.5 × 10^5 cells/coverslip) on top of the hippocampal neurons. Human glial-derived neurotrophic factor (GDNF; Alomone Labs, G-240) was added to the wells (10 ng/mL).

The next day, the medium was exchanged with preconditioned neuron medium containing 67 μg/mL 5-fluoro-oxyuridine, 165 μg/mL uridine, and 10 ng/mL GDNF. Neuron medium was preconditioned by incubation for 24 h on primary cultures of cortical glial cells (63). Ventral midbrain neuron cultures were from Munc13 DKO mice and their littermate controls were used to prepare neurons intended for imaging buffer (136 mM NaCl, 2.5 mM KCl, 2 mM CaCl₂, 1.3 mM MgCl₂, 10 mM HEPES, and 10 mM D-glucose, pH 7.4). Imaging buffer was prepared and reused for >1000 imaging sessions. Neurons were electrically stimulated by field stimulation, delivering 200 μs pulses at 20 Hz (2-ms duration, 16 V) from a Stimulus Generator 4000 (Multichannel Systems) through parallel electrodes. For chemical depolarization, 3 M KCl was added to the chamber and rapidly mixed by pipetting (final K⁺ concentration = 90 mM).

Synthesis of (GT)_10-Functionalized SWCNTs. Chirality-enriched (5,6)-SWCNTs (Sigma-Aldrich, 704148) were suspended in water (9 mg/mL). Approximately three flakes of SWCNTs were then added to 500 μL of D₂O PBS, prepared with heavy water (Sigma-Aldrich, 368407) and containing 200 pmol/μL (GT)_10 oligonucleotides. Heavy water improves the separation of solubilized single DNA-SWCNT complexes during centrifugation. Oligonucleotides were synthesized by the DNA Core Facility of the Max Planck Institute for Multidisciplinary Sciences, Göttingen, Germany, using a medium-throughput oligo synthesizer (Dr. Oligo 48, Biolytic Lab Performance Inc.) and purified by reverse-phase, high-pressure liquid chromatography (PerSeptive Biosystems).

A stable dispersion of SWCNTs was achieved based on previously published protocols (30, 38) through tip sonication for 1 h at 30% power at 4°C (Fischer Scientific Model 120 Sonic Dismembrator, Fisherbrand Probe 12921181, tip size 2 mm). SWCNT-(GT)_10 complexes were separated from nonfunctionalized SWCNTs by centrifugation (16,000 rpm for 30 min at 4°C). The supernatant containing SWCNT-(GT)_10 complexes was collected; this centrifugation enrichment step was repeated twice, and the final supernatant was used as a nanosensor stock.

UV-Vis-NIR Absorption and NIR Fluorescence Spectroscopy. The nanosensor stock concentration was determined by measuring the absorption spectrum in PBS (SI Appendix, Fig. S22) with an ultraviolet-visible (UV-Vis)-NIR spectrometer (JASCO V-670, Spectra Manager Software) and integrating the area below the (5, 6) peak using a molar extinction coefficient of 991 nm and an estimated length of 200 nm (65). The concentration was adjusted to 300 nM, and the stock was stored at 4°C for up to 3 wk. NIR fluorescence spectra (SI Appendix, Fig. S1) of nanosensors were acquired with a Shamrock 193i spectrometer (Andor Technology) on an iX53 microscope (Olympus). Then, 200 μL of nanosensor suspension in a 96-well plate was excited through a monochromator, at 560 nm, connected to a LSE341 light source (LOT-Quantum Design). NIR fluorescence spectra were taken before and after the addition of 100 μM DA.

Microscope Configuration. A custom-built system based on an Olympus IX53 microscope was used for imaging, using a 100× oil-immersion objective (UPLSAPO100XS, Olympus). The system used two cameras for simultaneous imaging of NIR and visible light. Visible light fluorescence was excited using an xCite 120Q (Excellis Technologies) and imaged using an Andor Zyla 5.5 camera (Andor Technology). NIR fluorescence was excited using a 561-nm laser (Cobolt Jive, Cobolt AB; Pmax. = 500 mW) and imaged using a Xenics Cheetah-640-TE1 InGaAs camera (Xenics). NIR imaging (>900 nm) was performed at 15 frames/s for all live-cell experiments. A schematic describing the optical components and configuration of the system is provided (SI Appendix, Fig. S5).

Application of AndromeDA and Live-Cell Image Acquisition. For imaging, a coverslip containing cultured neurons was washed with imaging buffer (2×10 mM NaCl, 2.5 mM KCl, 2 mM CaCl2, 1.3 mM MgCl2, 10 mM HEPES, and 10 mM D-glucose, pH 7.4). Imaging buffer was removed and 10 μL of SWCNT-(GT)_10 nanosensor stock was applied to the coverslip. Nanosensors immediately adsorb to the coverslip glass around the neurons, so the coverslip was washed immediately to remove unbound nanosensors. We refer to the 2D layer of nanosensor paint created through this process as AndromeDA.

The coverslip was mounted in an open-bath imaging chamber with integrated parallel-field stimulation electrodes (RC21-BRFS, Warner Instruments), facilitating the rapid addition of KCl and DA to the chamber during imaging. Experiments were performed at 21°C. Imaging buffer was supplemented with 10 μM 2,3-dihydroxy-6-nitro-7-sulfamoyl-benzo[f]quinoxaline (an α-amino-3-hydroxy-5-methyl-4-isoxazolepropionic acid [AMPA]/kainate receptor antagonist), 50 μM (2R)-α-amino-5-phosphonopentanoate (an N-methyl-D-aspartate [NMDA] receptor antagonist), and 10 μM sulpiride (a dopamine D2 receptor antagonist), ensuring that the DA release was not a product of circuit activity from glutamatergic neurons in the culture. Sulpiride was included to prevent autoinhibition of DA release.

Neurons were electrically stimulated by field stimulation, delivering 200 square-wave biphasic pulses at 20 Hz (2-ms duration, 16 V) from a Stimulus Generator 4000 (Multichannel Systems) through parallel electrodes. For chemical depolarization, 3 M KCl was added to the chamber and rapidly mixed by pipetting (final K⁺ concentration = 90 mM). Experiments typically followed this format: imaging with field stimulation, wash, imaging with KCl stimulation, wash, and then imaging with 100 μM DA addition. For 100 μM DA treatment, 1 mM DA (freshly dissolved) was added to the chamber to a final chamber concentration of 100 μM after mixing. 100 μM DA provided a measurement of the maximal AndromeDA response (SI Appendix, Fig. S4).
Appendix, Fig. S3), and normalization against this value allowed for correction for variations in maximum fluorescence among individual nanosensor batches. Protocols were performed only once for each coverslip. If not stated otherwise, neurons were compared with littermates only.

For imaging experiments with reserpine (Tocris Bioscience), reserpine was added to the medium (1 μM) of midbrain neurons and incubated for 90 min before experimentation (37 °C, 5% CO2) (66). AndromeDA was then applied and imaging experiments were conducted in the presence of fresh 1 μM reserpine. For L-DOPA (Tocris Bioscience) treatment, L-DOPA was added to the culture medium (100 μM) and the cells were incubated for 45 min (37 °C, 5% CO2) (35, 58), after which the cells were switched to fresh imaging buffer before AndromeDA application. Imaging was conducted in the absence of L-DOPA because the nanosensors respond to L-DOPA (23). In some experiments, action potential firing was inhibited using 300 nM TTX. 300 nM TTX is routinely used in our laboratory to inhibit action potential firing in neurons, and it washes out within 5 s. Imaging was carried out for 5 min in the presence of TTX in imaging buffer, followed by washing of the coverslip, replacement of the buffer with fresh imaging buffer, and imaging of stimulated DA release as described above.

To minimize data variability across experiments, we recommend checking the nanosensor stock concentration based on the UV-vis-NIR absorption spectra to ensure that a consistent 300 nM concentration is applied to coverslips. Nanosensors should be stored at 4 °C and kept for no longer than a week before preparing a fresh stock or centrifuging to remove aggregates. Nanosensors should be applied to the coverslip briefly, and for a consistent time (~3 s), before thorough washing to remove unbound nanosensors. We further recommend counting the midbrain neurons prior to plating and mixing well before the neurons settle on the coverslip to help generate a relatively consistent neuronal density on the coverslips to minimize biological variability in experiments.

DA Dose-Response Experiments. The dose dependency of the AndromeDA response to DA was determined by applying increasing concentrations of DA (0.1 nM, 1 nM, 10 nM, 100 nM, 1 μM, 10 μM, and 100 μM) to AndromeDA-coated glass coverslips. Coverslips were mounted in a closed imaging chamber and perfused with PBS (0.5 mL/min). DA solutions were prepared fresh and only used for up to 15 min to minimize oxidation. While imaging, a coverslip was perfused with each DA concentration in ascending order, with 10 min of washing by perfusion in between each DA concentration. The imaging region was not changed during each experiment. NIR images were acquired at 1 frame/s and normalized as described below, and pixel mean intensity values were collected from the entire image using the measure stack function in Fiji. The AndromeDA EC_50 dose-response curve (SI Appendix, Fig. S3) was calculated by plotting the peak NIR fluorescence response to each DA concentration and fitting a sigmoidal curve using nonlinear regression in GraphPad Prism.

Image Processing: Normalization. NIR images were acquired using Fiji (2.0.0-rc69/1.52u2/Java 1.8.0_172, 64 bit) with custom-written plugins (available from the authors). NIR image sequences were converted to 16-bit image stacks. A background image, acquired by capturing a frame of the buffer-filled imaging chamber without cells, was subtracted from all images to adjust for variations in ambient light on different days of imaging. The initial fluorescence of each pixel, Fc, was calculated for each NIR image sequence as the intensity of the projected mean image of the first 10 frames. For each pixel, the fluorescence intensity was normalized as follows. For a given image in the image sequence Fm, the normalized fluorescence change for every pixel is calculated:

\[
F_n = \frac{(F - F_0)}{F_0},
\]

where F is the intensity value of the pixel for the image being considered and F0 is the initial fluorescence. Normalized image sequences were then used for analysis.

Image Processing: Selection of Areas for AndromeDA Imaging. Areas of the coverslip containing no AndromeDA due to coverage by neurons (SI Appendix, Fig. S2) were excluded from analysis. The camera used to image EGFP (complementary metal oxide semiconductor, CMOS) has a larger sensor than the camera used to image NIR fluorescence (InGaAs). EGFP images were thus cropped and scaled to overlay with the NIR field of view, ensuring that EGFP and NIR images corresponded in space.

A single microscopic field of view was imaged in an experiment. To account for variations in axonal density, we analyzed NIR fluorescence only within 2 μm (10 pixels) around morphologically defined varicosities. Average fluorescence intensity within the ROIs was measured in each normalized image series calculated and plotted over time. A curve was then fit to the first 10 s of the resulting time trace, which represented the AndromeDA fluorescence baseline. This baseline curve was subtracted from the entire plot of fluorescence vs. time to account for any decline in AndromeDA fluorescence intensity over time.

DART: A Machine-Learning-Based Tool to Detect Hotspots of AndromeDA Activation. Hotspots were defined as localized, transient increases in AndromeDA fluorescence above the surrounding fluorescence signal. We developed a semiautomated, unbiased approach to their detection. DART is a Java object-oriented language algorithm, which runs as a plugin in Fiji and uses a hybrid approach combining computer vision and machine learning in hotspot detection (67). DART attempts to correct for fluorescence loss and flickering for enhanced detection of events. DART then uses a background subtraction approach for foreground detection of bright events in the image sequence (68, 69).

To discriminate bright pixels arising due to noise from pixels associated with true hotspots, DART attempts to reduce image noise via a machine learning-based calculation of the threshold for the subtracted image for the subtraction processes described above. Noise reduction is carried out through a series of machine-based decision-making steps that involves thresholding, median filtering, and eroding. As a result, a binary image is formed consisting of true events as larger white pixel areas and false events as smaller white pixel areas (resulting from noise) over a black background. To separate between true and false events, an erosion operation takes place with a kernel dimension [3 × 3], forming a binary image that is noise free. The algorithm then uses the eroded binary image to detect, segment, and label the white areas as individual events (hotspots). The segmentation process executes using an algorithm similar to the seed fill algorithm (70), called a wave spread function (WSF). This recursive WSF scans the image horizontally from left to right and then repeats the scan until it reaches the end of the image. Detected white pixels get a specific region identity (ID) number. Upon storing a pixel, the algorithm spreads the same process to the next neighboring connected pixels. All connected pixels get the same region ID number. The spread process is organized according to a pixel compass of an 8-point connection to identify separate event locations and durations over time. The algorithm also attempts to compensate for mechanical drifts, noise, and segmentation error by applying a user-defined search radius to search through previous white regions’ center coordinates and determine whether they belong to the same events or new events. Event ID numbers are then assigned based on this process. DART is available on request from the authors.

Image Processing: Analysis of NIR Nanosensor Fluorescence at Discrete Release Hotspots. DART was used to define ROIs containing AndromeDA hotspots. DART ROIs were then applied to normalized NIR movies. DART ROIs were manually verified to ensure that the ROIs contained bona fide hotspots of localized, transient AndromeDA activation. To correct for the activation of AndromeDA in the extracellular space by DA diffusion, the mean fluorescence intensity was measured in a region containing no axons. For each frame, the hotspot fluorescence was calculated as follows:

\[
\Delta F/F_0 = \left( \frac{F_{\text{Hotspot}} - F_0}{F_0} \right)
\]

\[
\Delta F/F_0 = \left( \frac{F_{\text{Hotspot}} - F_{\text{Extracellular space}}}{F_0} \right)
\]

The resulting fluorescence values for each DART ROI were plotted as a time sequence describing the relative increase in NIR fluorescence above AndromeDA activation due to diffusing DA. This method highlights local, transient increases in nanosensor fluorescence that would result from DA release events. For representative images of hotspots (Fig. 3 and SI Appendix, Fig. S12), a Gaussian filter was applied to images to improve image clarity.

Immunocytochemistry. Detailed immunocytochemistry materials and methods are provided in SI Appendix, Extended Materials and Methods.

AFM. AFM was conducted in intermittent-contact mode (scan rate = 0.5 Hz, 512 lines) using an Asylum Research MFP-3D Origin instrument equipped with rectangular cantilevers (Opus, MikroMasch Europe; Al coating, tetrahedral tip, νres = 300 kHz, k = 26 N m⁻⁻¹). Freshly cleaved muscovite mica was coated with PLL, treated briefly with nanosensors, washed, and dried using a N2 stream before sample measurement. Sample analysis was performed using Gwyddion.
Statistical Analysis. All statistical analyses were performed using Prism 8 (GraphPad). Sample size and independence are described in the figure legends. Where appropriate, data were analyzed for normality of distribution using the Shapiro-Wilk normality test. For normally distributed data, student t-tests were compared using unpaired two-tailed Welch’s t test or a Brown-Forsythe and Welch ANOVA depending on sample size. For nonparametric samples, were compared using unpaired two-tailed Welch’s t test or a Brown-Forsythe and Welch ANOVA depending on sample size. For nonparametric data, the Mann-Whitney or the Kruskal-Wallis test was used depending on sample size. The significance level was set at P < 0.05. An experiment (i.e., a given coverslip) was excluded from analysis if no induced increase in nanosensor fluorescence was observed during KCl stimulation or the final addition of 100 μM DA to each coverslip at the end of the experiment. NIR image sequences that were obscured through x/y drift or other obstacles such as air bubbles in the imaging chamber were excluded from analysis.

Data Availability. The Java archive file for the DART plugin for Fiji (DOI: 10.5281/zenodo.6406536 (67), as well as the code for the DA diffusion simulation and the data for the corresponding figures (DOI: 10.5281/zenodo.6406318 (37)), can be accessed on Zenodo. Raw data (images) are available from the corresponding authors upon reasonable request. All other study data are included in the article and/or supporting information.

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