

ARTICLE

In-materio Reservoir Working at Low Frequencies in a Ag₂S-island Network

Received 00th January 20xx,
Accepted 00th January 20xx

DOI: 10.1039/x0xx00000x

Motoharu Nakajima,^a Kazuki Minegishi,^a Yosuke Shimizu,^a Yuki Usami,^b Hirofumi Tanaka,^b and Tsuyoshi Hasegawa*^a

The abstract should be a single paragraph that summarises the content of the article A Ag₂S-island network is fabricated with surrounding electrodes to enable it to be used as a reservoir for unconventional computing. Local conductance change occurs due to the growth/shrinkage of Ag filaments from/into each Ag₂S island in the reservoir. The growth/shrinkage of Ag filaments is caused by the drift of Ag⁺ cations in each Ag₂S island, which results in a unique non-linear response as a reservoir, especially at lower frequencies. The response of the reservoir is shown to depend on the frequency and amplitude of the input signals. So as to evaluate its capability as a reservoir, logical operations were performed using the subject Ag₂S-island network, with the results showing an accuracy of greater than 99%.

Introduction

Artificial intelligence, such as Deep Learning, has become a widely used conventional technique in our present-day information society. The remarkable progress of artificial intelligence, which has been achieved mainly by software-based systems running on von-Neumann computers,¹ has enabled functions such as image recognition and voice recognition.² However, there is some concern that von-Neumann computing is approaching its performance limits,^{3,4} which will in turn limit further progress in artificial intelligence. The development of hardware-based neuromorphic computers is one of the solutions to this problem.⁵

Recently, much attention is being directed at reservoir computing (RC); an emerging computing method based on biological recurrent neural networks.⁶⁻⁸ Information processing in RC systems can be divided into two parts. In the first part, time-series signals are input into a reservoir, in which input signals are non-linearly transformed and output. In the second part, the output signals from a reservoir are linearly combined by multiplying a weight to give a computing result. In RC, what should be optimized by learning are the weights connecting a reservoir and the output layer, which is much different

from deep learning where weights between all neighboring layers have to be optimized.⁹⁻¹¹ In addition, research into materials-based RC systems is accelerating as a result of the ability of various physical and chemical systems to work as reservoirs,¹² including the use of nanoparticles of Au,¹³⁻¹⁶ SnOx^{17,18} and Ag-Ag₂S (core-shell).^{19,20} Nanowires have also been used to form reservoirs, as has been demonstrated by the use of networks of Ag-Ag₂S nanowires^{9,21-23} and molecule-adsorbed carbon nanotubes.²⁴⁻²⁶ In previous studies, several systems^{9,19-23} have utilized atomic switch technology, which controls the growth and shrinkage of a Ag filament from Ag₂S.²⁷ In conventional atomic switch operation, a Ag filament grows in a gap between a Ag₂S electrode and a counter metal electrode. By using the gap size and the Ag₂S electrode size as the parameters, operating characteristics such as switching speed,²⁸ volatile/nonvolatile operation²⁹ and short-term plasticity/long-term potentiation³⁰ can be controlled. For example, when being used as a single device, small gaps on the order of 1 nm enable faster nonvolatile switching, while large gaps on the order of 100 nm cause slower volatile/nonvolatile selective switching.³¹

In previous studies, Ag₂S-based reservoirs used smaller gaps of molecules surrounding Ag₂S nano-particles^{19,20} or an electrolyte between Ag₂S nanowires.²¹⁻²³ The frequency of input signals was on the order of 10 Hz or higher, depending on the gap size and the materials used in a given gap. When we use a reservoir as an edge-computing device, such as would be used for environment monitoring, it should respond to frequencies specific to the phenomenon to be detected. For instance, earthquakes are characterized by frequencies of 10 Hz or lower. It has recently been

^a Department of Pure and Applied Physics, Graduate School of Advanced Science and Engineering, Waseda University, 3-4-1 Okubo, Shinjuku-ku, Tokyo 169-8555, Japan.

^b Department of Human Intelligence Systems, Graduate School of Life Science and Systems Engineering, Kyushu Institute of Technology, 2-4 Hibikino, Wakamatsu, Kitakyushu 808-0196, Japan.

Electronic Supplementary Information (ESI) available. See DOI: 10.1039/x0xx00000x

pointed out that monitoring ultra-low-frequency vibrations of about 0.01Hz-0.1Hz, which are caused by the slow slip phenomenon at plate boundaries,^{32,33} is important in detecting the premonitory symptoms of large-scale earthquakes.³⁴ We have successfully developed a reservoir that can respond to the characteristic lower frequencies of earthquakes, which is just one example of the phenomena needing to be monitored.

In this study, a Ag₂S-island network is used as a reservoir, in which network conductive channels are formed by Ag filaments grown from Ag₂S islands,³⁵ as schematically shown in Fig. 1. Ag filament growth also causes a change in the conductance of each Ag₂S island, depending on the growth length of a given Ag filament and the size of a given Ag₂S island,³⁶ which also contributes to modify the conductance of each channel. Moreover, input signals induce gradients of Ag⁺ cations in Ag₂S islands, which results in the formation of an inner electric potential in each Ag₂S island. Namely, each Ag₂S island works as a nano-battery.³⁷ Due to these effects, Ag₂S island networks show unique characteristics as reservoirs.

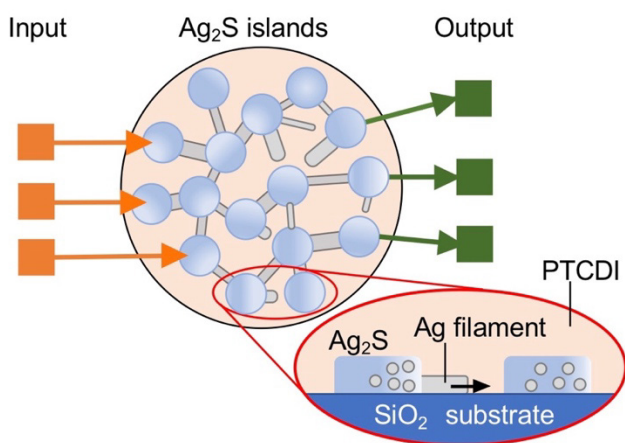


Fig. 1 Schematic of a Ag₂S island network-based reservoir. Ag filaments grown from Ag₂S islands make conductive channels in the network.

Experimental

In order to have a larger time constant in the reservoir operation, we made gaps larger than those used in previous studies.^{19,20} The Ag₂S reservoir used in this experiment was fabricated as follows.

First, surrounding Au electrodes were fabricated on a SiO₂ substrate by EB-lithography. Two types of electrodes were created. One has four electrodes surrounding a circular reservoir area with a diameter of about 1 μm (Fig. 2a), the other has sixteen electrodes surrounding a circular reservoir area with a diameter of 30 μm. The smaller four-electrode sample was used to evaluate the fundamental characteristics, while the sixteen-electrode sample was used to demonstrate the logic operations.

After the electrodes were made, Ag was thermally deposited onto the substrate annealed at 130 °C, while monitoring the current flowing between two of the electrodes with a bias application of 10 mV.^{38,39} When the monitored current increased to a few nA as a result of the tunneling current flowing among the Ag islands, the Ag deposition was stopped. The thickness of the Ag films deposited by this method was an average of 9.5 nm, which resulted in the formation of Ag islands 50 - 100 nm in size, with a spacing of a few nm between them.

Then, the Ag islands were sulfurized with sulfur vapor in a vacuum heating furnace at 70 °C for 20 min. During the process of crystallizing into Ag₂S, Ag islands change their morphology, which usually widens the spacing between the islands. Since the growth of Ag filaments requires a small number of electrons,^{40,41} we deposited molecules of N,N'-Dioctyl-3,4,9,10-perylenedicarboximide (PTCDI-C8) on the whole surface in order to ensure the presence of a small current, of less than 1 nA, to enable the Redox reaction (Ag⁺ + e⁻ ↔ Ag). The thickness of the said molecular film was 3 nm. Figure 2b shows an example of PTCDI-C8 covered Ag₂S islands observed by atomic force microscopy. Sulfurization seems to reduce the island size to 20 – 50 nm. Since atomic force microscopy could not resolve the narrow gaps between the Ag₂S islands, they appear to be connected to each other, although there are actually spaces between them.

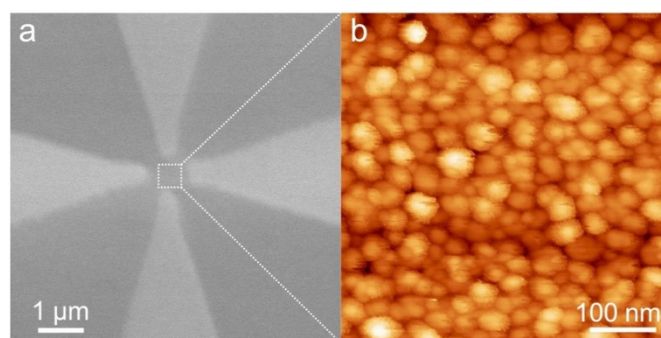


Fig. 2 (a) SEM image of four electrodes surrounding the Ag₂S island network. (b) AFM image of Ag₂S islands.

Results and Discussion

Basic characteristics of a Ag₂S-island network

Non-linear transformation of input signals is the most important function performed by a reservoir. Figure 3 shows the experimental results of non-linear transformation using the four electrode devices, where a sinusoidal wave V_{in} of 0.1 Hz was input with input amplitudes of (a) 4 V and (d) 6.2 V. The change in output signals V_{out} measured at one of the three electrodes are shown in Figs. 3b and 3e, respectively. Change in the resistance of the reservoir $R_{reservoir}$, which was calculated with a reference resistance R_0 of 1 MΩ and eq. (1) where we assumed the inner potential V_{inner} was zero, are shown in Figs. 3c and 3f, respectively.

$$R_{reservoir} = R_0 \cdot \left(\frac{V_{in} + V_{inner} - V_{out}}{V_{out}} \right) \quad (1)$$

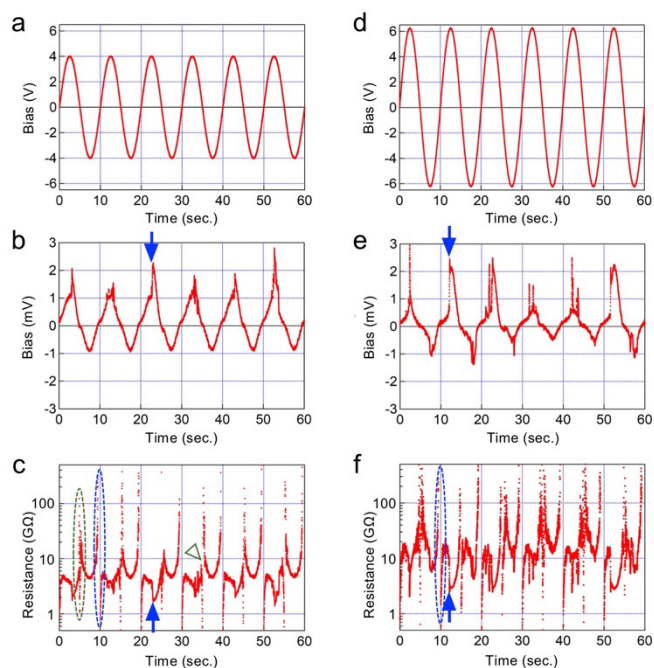


Fig. 3 Experimental results of a non-linear transformation following a sinusoidal input. The left-hand column shows (a) input, (b) output and (c) resistance when the amplitude of the input was 4 V. The right-hand column shows (d) input, (e) output and (f) resistance when the amplitude of the input was 6.2 V.

The shape of the output waves V_{out} completely changed from those of the input signals, indicating that the Ag_2S -island network has a non-linear transformation function. It can be seen that the larger amplitude (6.2 V) modified the input wave more than did the smaller amplitude (4 V). More interestingly, steep increases in output were frequently observed, such as those indicated by the blue arrows in Figs. 3b and 3f. These remarkable increases are assumed to be caused by the completion of the formation of Ag filaments that connect Ag_2S islands. As observed in the operation of a single atomic switch, they also resulted in a steep decrease in resistance in the Ag_2S -island network, as indicated by the blue arrows in Figs. 3c and 3g, where the resistance decreased (c) 3 GΩ and (f) 8 GΩ, respectively. Although the shape of the output signals changes a little in each period, non-linear transformations were repeatedly carried out, which is a requirement for time series prediction by RC.^{42,43}

In the calculation of $R_{reservoir}$, we assumed that the inner potential V_{inner} is zero, which results in disconnection of the curves in Figs. 3c and 3f, as indicated by the dotted blue ellipses. Since the inner potential should be not so large, it was expected from eq. (1) that the disconnection caused by the neglect of V_{inner} happens only when V_{in} and V_{out} become small, e.g., every five sec. However, it was observed every ten sec. For instance, at 5 sec. in Fig. 3c, disconnection is not so visible, as indicated by the dotted green ellipse. These differences can be understood to be caused by the remaining Ag^+ cations in a Ag_2S island. When the polarity of V_{in} changes from positive to negative, such as at 5 sec. in Figs. 3a and 3d, Ag filaments will shrink. In other words, any Ag filament that still exists on a Ag_2S island at

that time causes the number of Ag^+ cations in the Ag_2S island to be smaller than it was before the growth of said Ag filament. Such small number of Ag^+ cations can only form a smaller inner potential. On the other hand, when the polarity of V_{in} changes from negative to positive, such as at 10 sec. in Figs. 3a and 3d, Ag filaments will grow, meaning that almost all the Ag^+ cations are still in a Ag_2S island, and with a higher density at the side exhibiting growing. The larger number of Ag^+ cations in such distribution can form a larger inner potential. By smoothly connecting the curves, it is expected that an inner potential, with a maximum of about 1 V, will be formed in the Ag_2S -island network. When repeating the bias application, disconnection is also observed at a timing of every 5 sec., as indicated by the green arrow in Fig. 3c. As can be seen in Fig. 3f, this becomes more visible when using a larger input bias, which can make a larger slope in the distribution of Ag^+ cations in a Ag_2S island.

Dependence on amplitude and frequency of input signals

We measured the change in resistance ΔR when a steep increase in the output bias was observed, as shown in Figs. 3b and 3e, because ΔR indicates the degree of Ag filament growth in a Ag_2S -island network. Figure 4a shows the change in ΔR as a function of input frequency. With both input amplitudes of 4V and 6.2 V, ΔR decreases with increase in the input frequency, and becomes almost zero at around 0.5 Hz. This frequency range is much different from those of previous studies, in which frequencies of around 10 Hz were used,¹⁹⁻²³ suggesting that the spacings between the Ag_2S islands in this study were much larger than observed in those used in previous studies.

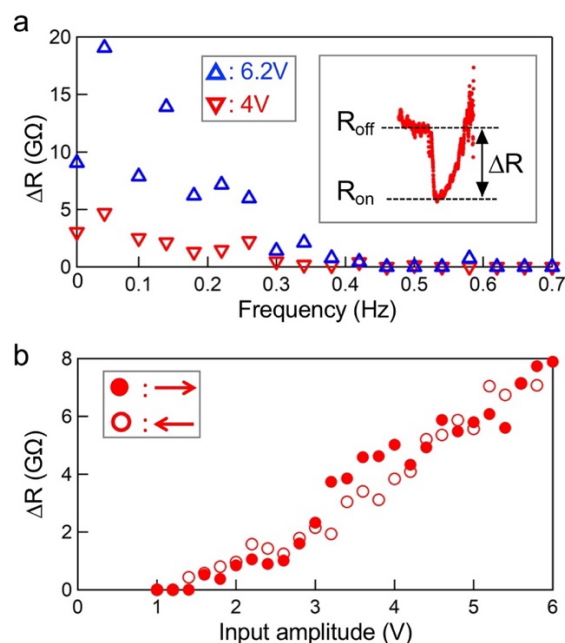


Fig. 4 Dependence of ΔR on the frequency (a) and on the input amplitude (b). The definition of ΔR is shown in the inset.

Figure 4b shows the dependence of ΔR on the amplitude of a sinusoidal input wave of 0.1 Hz. As can be seen, ΔR increases with increase in the input amplitude, indicating that Ag filaments grow longer and/or thicker at larger input wave amplitudes. The measurements were done both from 1 V to 6 V and vice versa. Because no dependence on the measuring sequence was observed, we were able to conclude that Ag filament growth was stably repeated, at least in the bias range up to 6 V. Here, it should be noted that ΔR is almost zero when the input amplitude is less than 1.4 V or the input frequency is higher than 0.5 Hz. This should not be taken to mean that Ag₂S-island networks function only with restricted conditions, which is not the case. Non-linear transformation is carried out even when ΔR is almost zero, as we were able to show by demonstrating logic operations in such condition (see Supplemental Information).

Power Spectral Density

In the human brain, spike firing occurs with a temporal correlation of several hundred milliseconds,^{44,45} which is characterized by $1/f$ noise.⁴⁶ Similar spatiotemporal fluctuations have been observed in various other systems, including physical reservoirs.²¹⁻²³ Analysis of power spectral density (PSD) often reveals the characteristics of the phenomena that occurred inside. For instance, PSD analysis on a network of Ag nanowires coated with polyvinylpyrrolidone (PVP) revealed the existence of two mechanisms that cause switching; one being low-frequency switching due to changes in connectivity across domains, and the other high-frequency switching due to Ag filament growth in the PVP film at individual junctions.⁴⁷

In this study, the PSD of a Ag₂S-island network was measured by applying a constant voltage of 4 V between two electrodes at opposite sites, the result of which measurement is shown in Fig. 5. Fitting the spectrum to $f^{-\alpha}$ gives $\alpha = 1.44$, indicating the existence of interactive change in resistance in the Ag₂S-island network. Namely, one Ag filament growth induces another Ag filament growth/shrinkage, in what amounts to a chain reaction, across the Ag₂S-island network. When Ag filaments bridge Ag₂S islands, the distribution of an electric potential in the Ag₂S-island network can change remarkably, causing a highly interactive phenomena characterized by α larger than 1. No significant Ag filament growth is observed at frequencies higher than 10 Hz. As a result, resistance change in one area does not affect other areas. Then, α becomes small due to independent random switching in the network, such as 0.76 in this measurement. Because of such characteristics, Ag₂S-island networks are more responsive to signals with lower frequencies.

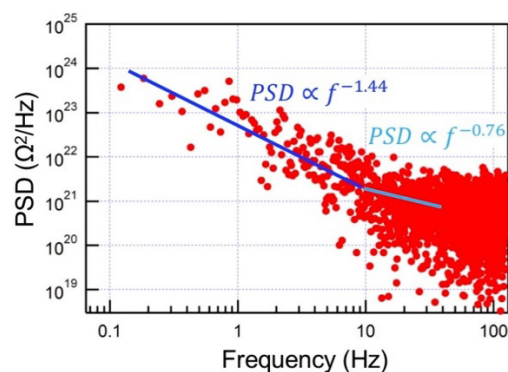


Fig. 5 Power spectral density (PSD) measured by applying a constant bias of 4V to a Ag₂S island network.

Logic operation

Logic operation is demonstrated using a Ag₂S-island network with 16 surrounding electrodes, as schematically shown in Fig. 6(a). The area of the Ag₂S-island reservoir was expanded to 30 μm in diameter in order to arrange the 16 electrodes with a certain number of Ag₂S islands, even in between adjacent electrodes. The 30- μm network also showed characteristics similar to those of a 1 μm -network, e.g., frequency and amplitude of input dependent non-linear transformation, as described in Supplemental Information. We input the two binary signals V_{in1} and V_{in2} , as shown in Fig. 6(b), to two electrodes located apart from each other, and the output signals, $O_1 - O_{14}$, were measured at the remaining fourteen electrodes. Figure 6(c) shows the fourteen measured output signals. Several output signals gradually changed over a few seconds, after the change in input signals, indicating the larger time constant of the Ag₂S island network.

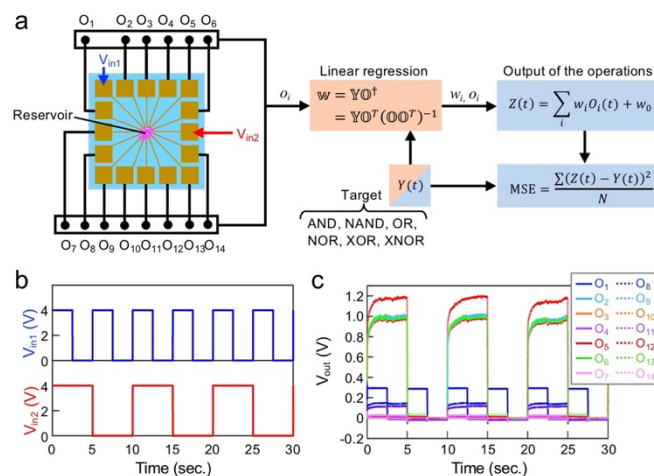


Fig. 6 (a) Schematic of a reservoir used for logic operations. (b) Input signals. (c) Output signals.

A linear combination $Z(t)$ was made by eq. (2) using the fourteen outputs, the weights, $w_1 - w_{14}$, and the bias, w_0 , where the weights and the bias were calculated by linear regression. The mean square error (MSE) between $Z(t)$ and the target $Y(t)$ was calculated by eq. (3), where N is 15,000 in this experiment. We demonstrated six logic

operations, i.e., AND, NAND, OR, NOR, XOR, and XNOR. The accuracy of each logic operation was calculated by eq. (4).⁴⁸

$$Z(t) = \sum_i w_i O_i(t) + w_0 \quad (2)$$

$$\text{MSE} = \frac{\sum(Z(t) - Y(t))^2}{N} \quad (3)$$

$$\text{Accuracy} = ((1 - \text{MSE}) \times 100)\% \quad (4)$$

Figure 7 shows the results of linear combinations to synthesize the six logic operations, with the target signals indicated by a solid black line in each panel. Including XOR and XNOR, which are difficult to synthesize due to the requirement of higher non-linearity in the transformation,^{17,38} all logic operations are carried out with a very high accuracy of over 99%. The fourteen outputs of $O_1 - O_{14}$ are individually shown in Supplemental Information. The optimized weights of $w_1 - w_{14}$ for each logic operation are also found in Supplemental Information. The results clearly suggest that Ag_2S -island networks act as reservoirs capable of outputting high-dimensional representations of inputs.

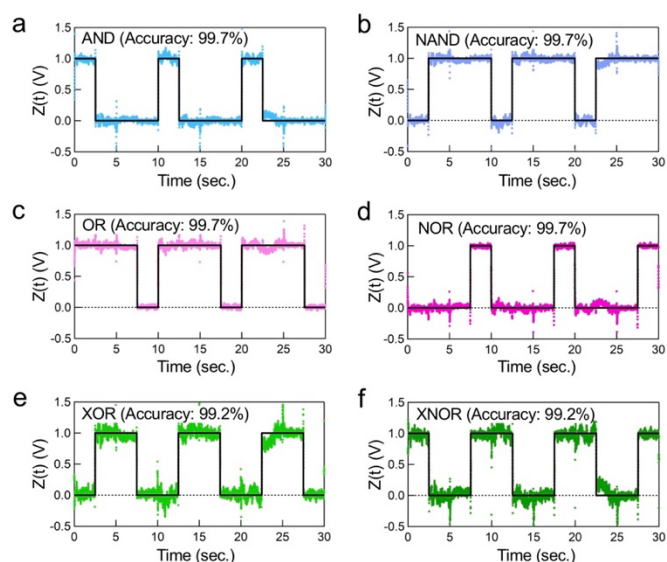


Fig. 7 Results of the logic operations. Synthesized curves are indicated by colored lines in each panel. The target in each logic operation is indicated by a solid black line.

Conclusions

In this study, we developed an RC device using a Ag_2S -island network as a reservoir, in which Ag filament growth from Ag_2S islands, as a result of electrochemical reactions, causes a change in resistance between electrodes. Due to the longer time constant of the Ag filament growth compared to electron movement, a Ag_2S -island network reservoir is more sensitive to frequencies lower than 0.5 Hz. The subject Ag_2S -island network reservoir shows a non-linear transformation that depends on the frequency and amplitude of the input signals. Power spectral density analysis clearly indicates that the Ag filament growth induces interactive responses in a Ag_2S -island

network at said lower frequencies, while local resistance change independently occurs at higher frequencies. However, regardless of the frequencies, the Ag_2S -island network reservoir non-linearly transforms input signals, as confirmed by the OR, AND, XOR, NOR, NAND, and XNOR logic operations at an high accuracy of over 99%. There are several phenomena characterized by lower frequencies, including low-frequency earthquakes, which it would be extremely beneficial to detect using edge devices in actual environments. The technique demonstrated in this study would contribute to develop of such systems.

Author Contributions

M. Nakajima and T. Hasegawa designed the whole part of the experiments and wrote the manuscript. M. Nakajima performed all the experiments. T. Hasegawa supervised the whole part of this study. K. Minegishi and Y. Shimizu analysed the experimental data. Y. Usami and H. Tanaka designed the experiments of logic operation. All authors reviewed and contributed to the manuscript.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

Part of this work was supported by JST/CREST.

References

- W. Yi, K. K. Tsang, S. K. Lam, X. Bai, J. A. Crowell and E. A. Flores, *Nat. Commun.*, 2018, **9**, 4661.
- Y. LeCun, Y. Bengio and G. Hinton, *Nature*, 2015, **521**, 436-444.
- D. J. Frank, *IBM J. Res. Dev.*, 2002, **46**, 235-244.
- A. Barabasi, *Science*, 2009, **325**, 412-413.
- L. A. Camunas-Mesa, B. Linares-Barranco and T. Serrano-Gotarredona, *Materials*, 2019, **12**, 2745.
- M. Inubushi and K. Yoshimura, *Sci. Rep.*, 2017, **7**, 10199.
- W. Maass, T. Natschläger and H. Markram, *Neural Comput.*, 2002, **14**, 2531-2560.
- C. Gallicchio, A. Micheli and L. Pedrelli, *Neurocomputing*, 2017, **268**, 87-99.
- H. O. Sillin, R. Aguilera, H.-H. Shieh, A. V. Avizienis, M. Aono, A. Z. Stieg and J. K. Gimzewski, *Nanotechnology*, 2013, **24**, 384004.
- C. Du, F. Cai, M. A. Zidan, W. Ma, S. H. Lee and W.D. Lu, *Nat. Commun.*, 2017, **8**, 2204.
- M. Lukoševičius and H. Jaeger, *Comput. Sci. Rev.*, 2009, **3**, 127-149.
- K. Vandoorne, P. Mechet, T. V. Vaerenbergh, M. Fiers, G. Morthier, D. Verstraeten, B. Schrauwen, J. Dambre and P. Bienstman, *Nat. Commun.*, 2014, **5**, 3541.
- C. Minnai, A. Bellacicca, S. A. Brown and P. Milani, *Sci. Rep.*, 2017, **7**, 7955.
- M. Mirigliano, F. Borghi, A. Podestà, A. Antidormi, L. Colombo and P. Milani, *Nanoscale Adv.*, 2019, **1**, 3119.
- M. Mirigliano, D. Decastri, D. Dellasega, A. Casu, A. Falqui and P. Milani, *Nanotechnology*, 2020, **31**, 234001.
- Y. Viero, D. Guérin, A. Vladyka, F. Alibart, S. Lenfant, M. Calame and D. Vuillaume, *Adv. Funct. Mater.*, 2018, **28**, 1801506.

17. S. K. Bose, J. B. Mallinson, R. M. Gazoni and S. A. Brown, *IEEE Trans. Electron Devices*, 2017, **64**, 5194.
18. P. Y. Le, B. J. Murdoch, A. J. Barlow, A. S. Holland, D. G. McCulloch, C. F. McConville and J. G. Partridge, *Adv. Electron. Mater.*, 2020, **6**, 2000081.
19. Hadiyawarman, M. Eguchi and H. Tanaka, *Jpn. J. Appl. Phys.*, 2020, **59**, 015001.
20. Hadiyawarman, Y. Usami, T. Kotooka, S. Azhari, M. Eguchi and H. Tanaka, *Jpn. J. Appl. Phys.*, 2021, **60**, SCCF02.
21. A. V. Avizienis, H. O. Sillin, C. Martin-Olmos, H.-H. Shieh, M. Aono, A. Z. Stieg and J. K. Gimzewski, *PLoS One*, 2012, **7**, e42772.
22. E. C. Demis, R. Aguilera, K. Scharnhorst, M. Aono, A. Z. Stieg and J. K. Gimzewski, *Jpn. J. Appl. Phys.*, 2016, **55**, 1102B2.
23. K. S. Scharnhorst, J. P. Carbajal, R. C. Aguilera, E. J. Sandouk, M. Aono, A. Z. Stieg and J. K. Gimzewski, *Jpn. J. Appl. Phys.*, 2018, **57**, 03ED02.
24. H. Tanaka, M. Akai-Kasaya, A. T. Yousefi, L. Hong, L. Fu, H. Tamukoh, D. Tanaka, T. Asai and T. Ogawa, *Nature Commun.* 2018, **9**, 2693.
25. Y. Usami, B. van de Ven, D. G. Mathew, T. Chen, T. Kotooka, Y. Kawashima, Y. Tanaka, Y. Otsuka, H. Ohoyama, H. Tamukoh, H. Tanaka, W. G. van der Wiel and T. Matsumoto, *Adv. Mater.*, 2021, **33**, 2102688.
26. D. Banerjee, T. Kotooka, S. Azhari, Y. Usami, T. Ogawa, J. K. Gimzewski, H. Tamukoh and H. Tanaka, *Adv. Intell. Syst.*, 2022, 2100145.
27. K. Terabe, T. Hasegawa, T. Nakayama and M. Aono, *Nature*, 2005, **433**, 47.
28. A. Nayak, T. Tsuruoka, K. Terabe, T. Hasegawa and M. Aono, *Nanotechnology* 2011, **22**, 235201.
29. T. Hino, T. Hasegawa, H. Tanaka, T. Tsuruoka, K. Terabe, T. Ogawa and M. Aono, *Nanotechnology* 2013, **24**, 384006.
30. T. Ohno, T. Hasegawa, T. Tsuruoka, K. Terabe, J. K. Gimzewski and M. Aono, *Nat. Mater.*, 2011, **10**, 591.
31. M. Araki and T. Hasegawa, *Jpn. J. Appl. Phys.*, 2020, **59**, 040605.
32. N. Poiata, J.-P. Vilotte, N. M. Shapiro, M. Supino and K. Obara, *J. Geophys. Res. Solid Earth*, 2021, DOI:10.1029/2021JB022138.
33. K. Matsuda, S. Ide, K. Ohta and T. Matsuzawa, *Earth, Planets and Space*, 2020, DOI:10.1186/s40623-020-01172-8.
34. K. Obara and A. Kato, *Science*, 2016, **353**, 253-257.
35. K. Ojima, T. Hasegawa, Y. Naitoh, H. Shima and H. Akinaga, *Jpn. J. Appl. Phys.*, 2020, **59**, SN1011.
36. N. Mishima, T. Tsuruoka and T. Hasegawa, *Jpn. J. Appl. Phys.* 2021, **60**, SE1001.
37. I. Valov, E. Linn, S. Tappertzhofen, S. Schmelzer, J. van den Hurk, F. Lentz and R. Waser, *Nature Commun.* 2013, **4**, 1771.
38. Y. Naitoh, K. Albrecht, Q. Wei, K. Yamamoto, H. Shima and T. Ishida, *RSC Adv.*, 2017, **7**, 53503.
39. K. Shinotsuka, L. Shen, K. Dai, H. Tokuno, S. Miura, T. Segawa, M. Kondo and Y. Hatta, *Jpn. J. Appl. Phys.*, 2019, **58**, 032001.
40. T. Hasegawa, K. Terabe, T. Tsuruoka and M. Aono, *Adv. Mater.*, 2012, **24**, 252-267.
41. K. Terabe, T. Nakayama, T. Hasegawa and M. Aono, *J. Appl. Phys.*, 2002, **91**, 10110.
42. E. Bollt, *Chaos*, 2021, **31**, 013108.
43. D. J. Gauthier, E. Bollt, A. Griffith and W. A. S. Barbosa, *Nat. Commun.*, 2021, **12**, 5564.
44. J. R. Manning, J. Jacobs, I. Fried and M. J. Kahana, *J. Neurosci.*, 2009, **29**, 13613-13620.
45. K. J. Miller, L. B. Sorensen, J. G. Ojemann and M. den Nijs., *PLoS Comput. Biol.*, 2009, **5**, e1000609.
46. A. Mazzoni, F. D. Broccard, E. Garcia-Perez, P. Bonifazi, M.E. Ruaro and V. Torre, *PLoS One*, 2007, **2**, e439.
47. A. Diaz-Alvarez, R. Higuchi, P. Sanz-Leon, I. Marcus, Y. Shingaya, A. Z. Stieg, J. K. Gimzewski, Z. Kuncic and T. Nakayama, *Sci. Rep.*, 2019, **9**, 14920.
48. D. Banerjee, S. Azhari, Y. Usami and H. Tanaka, *Appl. Phys. Express*, 2021, **14**, 105003.