Seed-mediated atomic-scale reconstruction of silver manganate nanoplates for oxygen reduction towards high-energy aluminum-air flow batteries

Jae Chan Ryu, Haeseong Jang, Joohyuk Park, Youngshin Yoo, Minjoon Park & Jaephil Cho

铝空气电池是下一代高能量密度储能的有前途的候选者，但存在的固有限制阻碍了它们的实用化。在这里，我们展示了用银纳米粒子介导的银锰酸盐纳米片作为碱性介质中氧还原的高活性和化学稳定催化剂。通过原子分辨透射电子显微镜，我们发现银锰酸盐纳米片表面的条纹图案起源于银和锰的锯齿状原子排列，形成高密度位错的晶格。这种结构能够提供高电导率，低电极电阻和丰富的离子吸附活性位点。该催化剂在流式铝空气电池中表现出杰出的性能，证明了在100 mA cm⁻²下的高加权和体积能量密度为~2552 Wh kg⁻¹和~6890 Wh l⁻¹，以及在机械充电过程中的高稳定性。
Metal-air cells have been highlighted as next-generation batteries, providing the possibilities of low cost and high-energy storage. Aqueous aluminum–air batteries (AABs) are a promising candidate for efficient power delivery in the field of transportation and uninterrupted power supply because of its high-energy density (theoretically, ~813 Wh kg\textsubscript{Al}\textsuperscript{-1} and ~21,954 Wh L Al\textsuperscript{-1}, see Supplementary Note 1) with abundant reserves, low cost, and light weight of aluminum (26.98 g mol\textsuperscript{-1}). In particular, aluminum has the highest volumetric capacity (~8.04 Ah cm\textsuperscript{-3}) among the electrode materials (e.g., Li: ~2.06, Mg: ~3.83, and Zn: ~5.85 A h cm\textsuperscript{-3}). However, there are several hurdles that need to be overcome for the practical application of the AABs, such as high corrosion rate of aluminum anodes, rapid self-discharge, and precipitation of solid by-products (Al(OH)\textsubscript{3} and Al\textsubscript{2}O\textsubscript{3}) on the surface of electrodes. Thus, technical progress has been mainly limited to the mismatch between theoretical and practical performances (<1000 Wh kg\textsubscript{Al}\textsuperscript{-1}). To date, intensive efforts have been devoted to overcome the above-mentioned issues by developing a proper aluminum alloy as anode\textsuperscript{10–12}, electrolytes\textsuperscript{13–15} and cell designs\textsuperscript{16,17}. Recently, AABs with Cu/Cu\textsubscript{2}O nanoparticles and Cu–N–C with the Ketjenblack supports were reported by using a sacrificial Cu-based metal–organic framework, showing a stable discharge of 1.53 V at 40 mA cm\textsuperscript{-2} with the aid of additives\textsuperscript{18}. In terms of the cell design, AABs containing two different electrolytes were also reported\textsuperscript{19} and demonstrated a specific discharge capacity of ~1810 mAh g\textsubscript{Al}\textsuperscript{-1} and energy of ~2081 Wh kg\textsubscript{Al}\textsuperscript{-1} at a current density of 20 mA cm\textsuperscript{-2}. In addition, Shim et al. fabricated deformable AABs with the high degree levels, which exhibited a specific capacity of 496 mAh g\textsubscript{Al}\textsuperscript{-1} at a current density of 5 µA cm\textsuperscript{-2}. However, its poor rate capability with low current density still limited the practical use. Thus, it is urgent to develop highly efficient, feasible alternative approaches. There is an additional intrinsic problem in AABs, which is a sluggish rate of the oxygen reduction reaction (ORR). Research into electrocatalysts for ORR can be categorized into metal- and non-metal elements. This result implies that as growth proceeded during heat treatment, the silver atom could move to the available site on the manganese substrate by creating the orthorhombic crystal structure of Ag\textsubscript{x}Mn\textsubscript{2}O\textsubscript{4}. Figure 1b summarized the catalytic effect of the SMNP in terms of the electrochemical properties, which will be discussed in detail.

The scanning electron microscopy (SEM) of the BM shows that the particle shape was bulky agglomerates with a size of ~15 µm because of the lack of additives for the morphology control (Fig. 1c), and X-ray diffraction (XRD) pattern indicated that it contained different phases of α-Mn\textsubscript{3}O\textsubscript{4}, β-Mn\textsubscript{2}O\textsubscript{3} and Mn\textsubscript{3}O\textsubscript{4} (Supplementary Fig. 5), and XRD analysis also exhibited four different phases of α-Mn\textsubscript{3}O\textsubscript{4} and β-Mn\textsubscript{2}O\textsubscript{3} (Supplementary Fig. 2a). In contrast, the PM showed nanosized polycrystalline structure with a size of ~100 nm (Fig. 1d), and consisted of mixed Mn\textsubscript{2}O\textsubscript{4} and Mn\textsubscript{3}O\textsubscript{4} phases (Supplementary Fig. 2b). We used a crystalline silver metal seed having an average size of ~100 nm, which is typically enclosed by four facets, for example, (111), (200), (220), and (311), as shown in Supplementary Fig. 2c. By using this metal seed, the plate-like configuration of the SMNP was obtained with distributed silver nanoparticles on the surface (Fig. 1e), showing a thickness of 30–40 nm (Supplementary Figs. 3, 4). Energy-dispersive X-ray spectroscopy (EDX) mapping revealed that the SMNP was composed of the silver, manganese, and oxygen atoms (Supplementary Fig. 5), and XRD analysis also exhibited four different phases of Ag, Ag\textsubscript{2}O, Ag\textsubscript{2}Mn\textsubscript{2}O\textsubscript{4}, and Mn\textsubscript{2}O\textsubscript{4} (Supplementary Fig. 2d). This result was consistent with the chemical position data from X-ray photoelectron spectroscopy (XPS) analysis. The Ag 3d, Mn 2p, and O 1s core levels from the SMNP structure exhibited the pronounced peaks related to the Ag\textsubscript{x}Mn\textsubscript{2}O\textsubscript{4} at 367.5, 373.5, 529.7, and 641.7 eV (Supplementary Fig. 6). We also measured the specific surface areas of the BM, PM, and SMNP, showing the values of 4.97, 14.18, and 11.47 m\textsuperscript{2} g\textsuperscript{-1}, respectively, where the PM showed the

Results

Electrocatalyst preparation and characterization. Our unique strategy enables the formation of silver metal seed-mediated silver manganate nanophases (hereafter referred to as SMNP) by simple precipitation and heat-treatment methods. Briefly, the manganese precursor and silver nanoparticles functionalized with polyvinylpyrrolidone as a chelating agent were dispersed in ethanol solution, and then NaOH as a reducing agent was added under stirring. After the precipitation reaction was completed, the prepared manganese hydroxide with a silver nanoparticle was calcined at 450 °C for 15 min (see Methods). Figure 1a shows the schematic of different morphologies of each sample according to synthesis conditions. For comparison, bulk manganese oxide (BM) and polyhedron manganese oxide (PM) were prepared without a reducing agent or silver metal seed, respectively. One observation is that the incorporation of silver metal in a manganese hydroxide precursor enabled the growth of the plate-like SMNP, in which silver nanoparticles could act as a growth seed (Supplementary Fig. 1). The lower coordination of surface atoms (typical low-index (111) or (100)) was reported to induce the internal lattice strain to minimize the surface energy.\n
In our case, the silver nanoparticles in the SMNP, which has a low index of (111) and a particle size of <100 nm are expected to induce the similar structural disorder, thus generating surface strain in Mn\textsubscript{2}O\textsubscript{4}.

In addition, the unique striped patterns with intragranular defects were observed in the SMNP, which could be ascribed to the zigzag atomic arrangements of silver and manganese elements. This result implied that as growth proceeded during heat treatment, the silver atom could move to the available site on the manganese substrate by creating the orthorhombic crystal structure of Ag\textsubscript{x}Mn\textsubscript{2}O\textsubscript{4}. Figure 1b summarized the catalytic effect of the SMNP in terms of the electrochemical properties, which will be discussed in detail.

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highest value because it consisted of the nanoparticles (Supplementary Table 1).

High-resolution transmission electron microscopy (HR-TEM) image of the SMNp clearly showed the striped patterns on the surface along with a wide range of dislocations and intragranular defects (Fig. 1f, g). The fast Fourier transform (FFT) image of the SMNp shows the orthorhombic phases on the basis of the unit spots of (−100) and (001) along the [010] zone, which are consistent with XRD results of Ag₂MnO₄ (space group: Pnma). This implies that the structure of nanoplate parts in the SMNp was Ag₂MnO₄ and not manganese oxide itself. To understand the complex structures of the SMNp, inverse FFT images were obtained by applying different masks on it along the [010] zone (Fig. 1h, i). Apparently, a high concentration of dislocations with crystal defects was discovered on the striped structures of the SMNp, and the other zone axis of [123] direction also showed similar results. Notably, from the dislocation edges, d-spacing value was increased from ~0.389 to ~1.124 nm, indicating continuously propagated defects (Supplementary Fig. 7). These dislocation edges could act as the nucleation sites, generating the intragranular cracks. The evolution of dislocation could be originated from the different atomic size of silver and manganese elements within the crystal lattice. The larger atomic size of the silver element than the manganese element could trigger a substantial lattice strain in the nanoplate, which significantly dislodges nearby atoms and generates an anisotropic defect structure in the SMNp. In terms of catalytic activity, the exposed defects may provide abundant active sites and ionic channels, reinforcing the ion transport kinetics. To clarify the morphology change and the concentration of dislocations of the SMNp, the control experiment with different amounts of precursors (silver nanoparticle, manganese source, and reducing agent) showed high conductivity and stability. Fig. 1 Formation of the stripe pattern and dislocation. a, b Schematic of the synthesis process (a) of the bulk manganese oxides (BM), polyhedron manganese oxide (PM), and silver manganate nanoplate (SMNp), and the summary of the catalytic effects (b) of the SMNp. c-e Scanning electronic microscopy (SEM) images of the BM (c), PM (d), and SMNp (e) showing plate-like structures with silver nanoparticles. f, g High-resolution transmission electron microscopy (HR-TEM) images of the SMNp (f) and high-magnified image (g) of the box in f, which shows zigzag atomic arrangements with intragranular cracking. Fast Fourier transform (FFT) image of g presents the lattice planes of (−100) and (001) along the [010] zone, indicating the orthorhombic structure of Ag₂MnO₄. h, i Inverse FFT images of applied masks at the blue circle (h) and red circle (i) in g, showing a wide range of dislocations with defects. The dislocations are shown by circles and edge dislocations are represented by a T-shaped symbol. The upper right insets in h and i indicate the applied masks and the lattice spacing of the (101) plane is denoted as the left inset in h. Scale bars, 5 μm (c), 500 nm (d), 200 nm (e), 100 nm (f), 4 nm (g), and 1 nm (h, i)
agent) was conducted (see Methods), which was summarized in Supplementary Figs. 8–10. As a result, increasing the amounts of silver nanoparticles reduced the concentration of dislocations with lower defects, showing high crystallinity of selected area electron diffraction (SAED) patterns (Supplementary Fig. 8). On the contrary, nanoplate morphology was not observed when both amounts of manganese and the reducing agent were controlled (Supplementary Figs. 9, 10). This implied that the seed-mediated growth for forming nanoplate morphology required appropriate contents of manganese source and the reducing agent.

To further confirm the origin of striped pattern on the SMNp, we performed atomically resolved scanning transmission electron microscopy (STEM) and electron energy loss spectroscopy (EELS). To clearly track the silver and manganese arrangements, we used high-angle annular dark-field STEM (HAADF-STEM) analysis and STEM–EDX mapping of the SMNp. As shown in Fig. 2a, individual atomic columns of silver and manganese were not observed when both amounts of manganese and the reducing agent were controlled (Supplementary Figs. 9, 10). This implied that the seed-mediated growth for forming nanoplate morphology required appropriate contents of manganese source and the reducing agent.

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corresponding number of transferred electrons (obtained at a potential of 0.30 V in which strongly supported the formation of Ag$_2$MnO$_4$ structure evenly distributed in the plate-like structure (Supplementary spectrum, and this revealed that silver and manganese atoms were (vs. RHE) with 1600 r.p.m showing good stability of SMNp compared with Pt/C mentary Fig. 11). Furthermore, the ABF intensity pro
tetragonal crystal structure (Supplementary Fig. 13). microscopy and spectroscopy results of the PM showed a typical crystallographic models. In comparison to this, the electron microcopy and spectroscopy results of the PM showed a typical tetragonal crystal structure (Supplementary Fig. 13).

The detailed elemental composition and distribution of the SMNp was investigated by using STEM–EDX mapping and point spectrum, and this revealed that silver and manganese atoms were evenly distributed in the plate-like structure (Supplementary Figs. 14 and 15). Interestingly, the average atomic ratio of silver and manganese elements on the SMNp was approximately 2, which strongly supported the formation of Ag$_2$MnO$_4$ structure (Supplementary Table 2). A series of EEL spectra were randomly collected from the STEM image (Fig. 2f), where all sites showed the identical Ag M$_{4,5}$-edge, O K-edge, and Mn L$_{2,3}$-edge (Fig. 2g). This implies that the plate-like region of the SMNp consists of the Ag$_2$MnO$_4$ structure.

**Electrocatalyst activity and stability.** To investigate the ORR activity of the SMNp, we measured linear sweep voltammetry (LSV) using a rotating ring–disk electrode (RRDE) in O$_2$-saturated electrolyte solution of 0.1 M KOH solution at 24 °C (Fig. 3a). For comparison, the ORR performances of Ag, BM, PM, and commercial 20 wt% Pt/C were also measured under the same condition. The pristine Ag nanoparticle showed the poor catalytic activity with respect to both onset potential and limiting current density. The catalytic activity of the BM and PM was higher than that of Ag nanoparticle, showing positively shifted onset potentials. Although the PM had the highest specific surface area (14.18 m$^2$ g$^{-1}$), the ORR activity was limited because it contained the Mn$_3$O$_8$ phase having low electrical conductivity$^{35}$. For the SMNp, we observed a much higher onset and half-wave potentials of 0.90 ± 0.01 and 0.80 ± 0.01 V vs. RHE (reversible hydrogen electrode) than the other samples, which was little smaller than the onset (1.05 ± 0.01 V vs. RHE) and half-wave potential (0.85 ± 0.01 V vs. RHE) of the Pt/C. It was clear that the existence of silver nanoparticles in the SMNp could significantly decrease the overpotential due to its high electrical conductivity, promoting electron transfer processes. The fluctuating current density of the SMNp at low-voltage range may result from capacitance correction because of large non-faradaic capacitance (Supplementary Fig. 16). The ORR activities of the control samples with different contents of precursors were shown in Supplementary Fig. 17. We found that the SMNp with an optimized content of silver nanoparticle (0.10 g), manganese source (0.07 g), and reducing agent (0.6 mL) showed the best electrochemical performance. We also conducted the rotation speed-dependent polarization curves for the SMNp to estimate the reaction pathways for the ORR (Supplementary Fig. 18). The Koutecky–Levich (K–L) plots of the
SMNp showed the nearly parallel linear plots, which was the first-order reaction with respect to oxygen.

To elucidate the reaction pathway for the ORR, we calculated the value of electron transfer number per oxygen molecule (n). The number of electrons for the SMNp was ~3.9, which was obviously higher than that for Ag, BM, and PM (Fig. 3b), implying that the direct four-electron pathway was governed during the ORR process. Further, we confirmed the excellent ORR activity of the SMNp from the mass transfer-corrected Tafel plots (Fig. 3c). The slope of the SMNp at a low overpotential region showed the lowest value of 54 mV dec⁻¹, which was even lower than Pt/C (60 mV dec⁻¹). This result also indicated that the rate-determining step of both the SMNp and Pt/C was the formation of superoxides by one electron transfer. In addition, a larger kinetic current density of the SMNp (1.13 mA cm⁻² at 0.85 V vs. RHE) indicated a higher ORR activity, in comparison to the Ag nanoparticle, BM, and PM (Supplementary Fig. 19). Such improvements of the SMNp can be explained by the intragranular cracks and abundant dislocations in nanoplate structures, providing a wide range of oxygen adsorption and desorption sites for facile ion transport.

In terms of the manganese oxidation state, the mixed valence state (Mn³⁺/⁴⁺) of the SMNp confirmed by the XPS measurement could promote the oxygen bond cleavage. In addition to the ORR activity, the electrochemical stability of the catalysts is another important factor, thus, we measured its chronoamperometric (i-t) response for the ORR (Fig. 3d). Interestingly, the durability of the SMNp showed ~17.4% higher than that of the Pt/C, which could be ascribed to ~200 mV higher equilibrium potential of Ag/Ag₂O than that of Pt/PtO in alkaline solution.

Aluminum-air flow battery performance. The electrochemical performance of the SMNp catalyst was evaluated by aluminum-air flow batteries (AAFBs). As shown in Fig. 4a, we designed a flow-type aluminum-air cell, and this cell assembly consists of an aluminum metal anode, catalyst-loaded air electrode, and alkaline electrolyte tank (Supplementary Fig. 20). By supplying fresh electrolyte, we can avoid the precipitation of by-products, making the surface of electrodes clean. In addition, self-discharge problem can be prevented by draining off the electrolyte, ensuring a long shelf life of the AAFBs. In comparison to a conventional static cell, the AABF demonstrated improved peak power density of ~71.2 mW cm⁻² at 100 mA cm⁻², and the discharge duration of the AAFBs was 11 times longer than the static cell at 50 mA cm⁻² (Supplementary Fig. 21a, b). In terms of the by-products precipitation, the surface of aluminum anode in a static cell was covered with a solid Al(OH)₃ precipitate, thus causing severe cell degradation with low discharge capacity (Supplementary Fig. 21c, d).

To further increase the battery performance, we used catalysts loaded with air electrodes (Ag nanoparticle, BM, SMNp, and Pt/C) in the AAFBs. The discharge process was initiated after 10 min of rest with a circulating electrolyte to activate aluminum anode by eliminating the surface oxide phase. As shown in Fig. 4b and Supplementary Fig. 22, the SMNp demonstrated significantly enhanced peak power density of ~105.2 mW cm⁻² compared to the pristine air electrode. This value was comparable to that of the Pt/C (~104.0 mW cm⁻²), and the maximum current density of the SMNp (~237 mA cm⁻²) was even higher than that of the Pt/C (~226 mA cm⁻²). Power density results of other samples were summarized in Supplementary Table 3. The effects of the ZnO additive in 6 M KOH electrolytes are shown in Supplementary
Fig. 23, which increases the discharge voltage of ~0.11 V. However, we conducted flow cell tests without any additives to clearly demonstrate the electrolyte flowing system. The galvanostatic discharge tests were conducted at different constant currents of 50 and 100 mA cm$^{-2}$. The specific capacity of the SMNp was ~2642 A h kgAl$^{-1}$ at 50 mA cm$^{-2}$, which was calculated on the basis of aluminum weight (Supplementary Fig. 24). When the current density increased to 100 mA cm$^{-2}$, the SMNp showed discharge time of 7.3 h corresponding to the specific capacity of ~2843 A h kgAl$^{-1}$, approaching the theoretical specific capacity of 2978 A h kgAl$^{-1}$ (Fig. 4c and see Supplementary Note 1). Notably, the gravimetric and volumetric energy densities of the SMNp were ~2552 W h kg$^{-1}$ and ~6890 W h L$^{-1}$, which was summarized in Supplementary Table 4. When the cells were subjected to the constant voltage test at 1 V (Supplementary Fig. 25), the SMNp exhibited the maximum current density of ~87 mA cm$^{-2}$, which was higher than that of the Pt/C (~82 mA cm$^{-2}$). This outstanding electrochemical performance could be originated from the synergistic effects of silver nanoparticle with high electrical conductivity and plate-like, striped Ag$_2$MnO$_4$ structure with abundant dislocations.

Remarkably, the AAFB with the SMNp showed excellent cycle stability upon a mechanically charging process, where the electrolyte and aluminum plate were replaced after fully discharging at 25 mA cm$^{-2}$ (Fig. 4d). At the initial stage, the Pt/C showed relatively lower overpotentials than the SMNp, showing a discharge potential of ~1.24 V. However, from the second cycle, the significant potential drop was observed in the Pt/C because of degradation of platinum element in concentrated alkaline solution. The SMNp presented stable discharge curves during prolonged cycles, suggesting its chemical stability, which was agreed with the chronoamperometric response in Fig. 3d. We also confirmed that the SMNp catalyst was stably maintained after battery cycling by SEM–EDX analysis (Supplementary Fig. 26). Therefore, the choice of the silver-based materials offers the large potentials of developing high-energy AAFBs with a minimal cell degradation.

**Discussion**

The origin of the improvement of cell potential and stability by the SMNp catalyst can be understood with respect to the electrical conductivity of the electrode and the degree of metal dissolution. First, we measured electrical conductivity of the catalyst-loaded air electrodes by using a through-plane
connection can be overcome by the silver element.53 As shown in Fig. 5a, the electrical conductivity of the Ag-loaded air electrode (5.05 × 10−2 S cm−1) was higher than that of the Pt/C-loaded one (4.01 × 10−2 S cm−1) because of lower electrical resistivity of silver than platinum. Interestingly, the SMnP revealed the highest electrical conductivities of 5.18 × 10−2 S cm−1, which could be ascribed to the silver nanoparticle-distributed nanoplate configuration. This also implied that the intrinsic drawback of dislocation having poor grain-to-grain connection can be overcome by the silver element.53 The electrical properties of all samples are summarized in Supplementary Table 5.

At a current density of < 100 mA cm−2, we confirmed that the discharge voltage of the Pt/C was substantially higher than that of the SMnP in polarization curves (Supplementary Fig. 22), which was well matched with the results from LSV curves (Fig. 3a). However, the discharge voltage of the SMnP has started to be comparable with that of the Pt/C at a current density of > 100 mA cm−2, which eventually exceeded the discharge voltage of Pt/C. These phenomena were also confirmed by the individual discharge curves of the SMnP and Pt/C with different current densities of 25 (Fig. 4d), 50 (Supplementary Fig. 24), and 100 mA cm−2 (Fig. 4c). When we carefully observed the discharge curves in relatively low current densities of 25 and 50 mA cm−2, the overpotentials for the SMnP were higher than Pt/C, thus lowering the discharge voltage. However, the SMnP showed a comparable voltage profile to Pt/C at a high current density of 100 mA cm−2, which could be ascribed to the superior electrical conductivity of the SMnP. This was confirmed by measuring the electrical conductivity of catalysts-loaded air electrode itself (Fig. 5a and Supplementary Fig. 27). Therefore, under the current density of > 100 mA cm−2, the electrical conductivity of the electrode plays a key role for enhancing the overall electrochemical performance of aluminum–air flow battery.

In terms of the chemical stability, we analyzed the dissolution of precious metal after cycling by using inductively coupled plasma mass spectrometry (ICP-MS). The electrolytes were collected after the AAFB was discharged for 6 h at 50 mA cm−2 (Fig. 5b). The cell equipped with the Pt/C showed a concentration of platinum dissolution of 33.5 × 10−2 ppm. For the SMnP, we observed that the concentration of silver dissolution was 18 times lower than that of platinum (1.8 × 10−2 ppm). This exceptional chemical stability of silver element in the SMnP can reasonably support the results of chronoamperometric response and mechanical recharging test.

We compared the gravimetric energy density of our AAFB with gasoline59 (Fig. 5c). It was reported that the theoretical energy density of gasoline was ~13,000 W h kg−1 (~9500 W h L−1). However, the practical energy density of gasoline is ~1700 W h kg−1 (~1200 W h L−1) considering the average tank-to-wheel efficiency of 12.6%.60 Compared to this, the gravimetric energy density of our AAFBs with SMnP was ~2552 W h kg−1, which showed large potentials for the aluminum as an alternative energy source. The volumetric energy density of each system was also summarized in Supplementary Fig. 28. Finally, we showed that the SMnP catalyst can be also used in zinc–air flow batteries (ZAFBs) and compared with AAFBs (Fig. 5d, Supplementary Fig. 29 and Table 6). The ZAFBs with SMnP showed the discharge capacity of ~558 mA h and gravimetric energy density of ~704 W h kg−1 at a current density of 100 mA cm−2, which still lags behind the performance of the AAFBs. Notably, the amount of consumed metal per discharge capacity in the AAFBs was ~0.35 g h−1, in comparison to the value of ~1.24 g h−1 in the ZAFBs, indicating three times higher conversion efficiency of the AAFBs.

In conclusion, we have prepared a silver nanoparticle seed-mediated silver manganese nanoplate architecture for ORR. We discovered that the silver atom can migrate into the available crystal lattice and rearrange manganese oxide structure, thus creating abundant surface dislocations. The activity and durability of this material outperformed the platinum-based electrocatalyst in concentrated alkaline solution. Furthermore, we have shown that flow-based aluminum–air battery with our catalyst not only offers a route to stable, robust energy conversion device, but also brings unprecedented gravimetric and volumetric energy densities of ~2507 W h kg−1 and ~6890 W h L−1, as summarized in Supplementary Table 7. This innovative strategy prevented the precipitation of solid by-product in the cell and dissolution of a precious metal in air electrode. We believe that our AAFB system has the potential for a cost-effective and safe next-generation energy conversion system.

Methods

Catalyst synthesis. The silver nanoparticle-mediated silver manganese nanoplate (SMnP) was synthesized by the simple precipitation and heat-treatment process. First, 0.07 g of manganese(II) nitrate hydrate (≥98%, Sigma–Aldrich) and 0.1 g of silver nanoparticle (polynaplyrolidone functionalized, particle size ≤100 nm, ≥99.5%, Sigma–Aldrich) were subsequently added into 2 mL of ethanol solution under vigorous stirring with a magnetic bar. Next, to form a uniform coating manganese layer on silver nanoparticles, 0.6 mL of 8 M NaOH solution as a reducing agent was prepared by sodium hydroxide beads (≥98%, SAMCHUN), and then poured into the above-blended solution. After filtered into the above-blended solution, the solution was centrifuged at 4000 r.p.m. for ~20 min. then poured into the above-blended solution. After filtering into the above-blended solution, the solution was centrifuged at 4000 r.p.m. for ~20 min. and 200 µL of 5 wt% NaOH solution (0.15 and 2.40 mL) were conducted with the same procedure above.

Structural characterization. The morphology of the synthesized materials was examined using scanning electron microscopy (SEM, VERIOS 460, FEI) operating at 10 kV and high-resolution transmission electron microscope (HR-TEM, ARM300, JEOL) operating at 300 kV. Elemental mapping was attained using energy-dispersive X-ray spectroscopy (EDX) equipped with the SEM and TEM. The crystal structure analysis was performed using an X-ray diffractometer (XRD, D/Max2000, Rigaku) with Cu-Kα radiation. Surface chemistry was analyzed using X-ray photoelectron spectroscopy (XPS, Thermo Scientific Ka spectrometer, 1486.6 eV).

Preparation of catalyst inks and working electrode. The catalyst ink was prepared using a mixture of 16 mg of the catalyst powder, 4 mg of a Ketjenblack, 800 µL of ethanol, and 200 µL of 5 wt% Nafion solution (Sigma–Aldrich). The Pt/C catalyst ink was made by adding 10 mg of Pt/C based on Vulcan XC 72 (20 wt%, Premetek Co.) powder into 800 µL of ethanol and 200 µL of 5 wt% Nafion solution. The catalyst inks were homogeneously mixed by ultrasonication, and then 5 µL of the ink was loaded on the surface of the polished glassy carbon of the rotation ring–disk electrode (RRDE) as a working electrode. After loading ink, the working electrode was dehydrated by the vacuum chamber for 30 min. As a result, the loading levels of the synthesized catalysts and Pt/C were 0.637 mg cm−2 and 0.398 mg cm−2, respectively (79.6 mg cm−2 for the pure Pt amount).

Electrochemical measurements. The rotating ring-disc electrode (RRDE, ALS Co., Ltd.) measurement was conducted using the catalyst films that were deposited on a glassy carbon electrode in 0.1 M KOH electrolyte saturated with oxygen at 1600 r.p.m. A Pt wire and Hg/HgO were used as a counter and a reference electrode, respectively. Electrochemical characterization was carried out using a bipotentiostat (IViumStat) with ohmic drop correction. The ORR current was measured after the current had saturated by using a cyclic voltammetry (CV) measurement that was made for voltages of 0.20 to ~0.67 V (vs. Hg/HgO) at a scan rate of 10 mV s−1. The capacitive current was corrected by measuring a CV measurement in 0.1 M KOH electrolyte saturated with argon. The measured potentials (vs. Hg/HgO) were calibrated with respect to a reversible hydrogen electrode (RHE) value by adding 0.929 V to the measured potential (vs. Hg/HgO). The number of transferred electrons (n) was calculated using the equation below where the collection efficiency (N) was determined using 10 mM K[Fe(CN)6] electrolyte under argon atmosphere, which was around 0.41. This value is similar to the
theoretical value of 0.42. \( I_d \) and \( I_r \) represent disk and ring currents, respectively.

\[
n = \frac{4}{3} \frac{I_d}{I_d + I_r/N}
\]

The electrochemical properties of the AAFBs were evaluated by using a potentialstat/galvanostat (WBC3000, WonAtech) with a voltage cutoff of 0.4 V for a primary flow cell test and 10 h of discharge per cycle for a mechanically charging process, respectively.

**Kinetic current calculation.** Kinetic current of electrocatalysts was obtained from mass transfer correction using Levich equation.

\[
n_{\text{measured}} = I_k + I_d \quad \text{where} \quad n_{\text{measured}} \text{ is the measured oxygen reduction current density, } I_k \text{ is the kinetic current density, and } I_d \text{ is the limiting current density.}
\]

**Aluminum-air flow cell tests.** For aluminum-air flow cell test based on the SMNP catalyst, a flow cell kit was fabricated in laboratory scale. Aluminum plate (alloy 6061, 97.93%, Alfa-Aesar) was used as an anode and 40 ml of 6 M KOH as an electrolyte. No separator was used and the air electrode was prepared via a homogenizer to form homogeneous gas diffusion layer (GDL). First, activated charcoal (Darco G60A, Sigma-Aldrich) and catalysts (including Pt/C) were mixed using a homogenizer to form homogeneous gas diffusion layer (GDL). First, activated charcoal was used as an anode and 40 ml of 6 M KOH as an electrolyte. The nickel foam was pasted on the nickel foam using roll press machine with a thickness of ~700 \( \mu \text{m} \). The pristine air electrode was made by the same procedure at a weight ratio of 7:3 for activated charcoal and Pt/C binder. The GDL was hot-pressed at 120 °C for 5 min attached with a PTFE film to ensure proper gas distribution and avoid leakage problems. For a mechanically rechargeable test, the aluminum plate and electrolyte were replaced after discharging at 25 mA cm\(^{-2}\) for 10 h. The air electrodes were continuously used without replacement to confirm the cycle stability. The flow rate of electrolytes for flow cell test was 100 mL min\(^{-1}\).

**Data availability**

The data that support the findings of this study are available from the corresponding author upon request.

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Author contributions
J.R., H.J. and M.P. designed the experimental plans. J.R. synthesized the electrocatalysts, conducted the RRDE measurements, flow cell experiments, structural analysis, crystallographic modeling, and wrote the initial draft of the manuscript. H.J. carried out the synthesis of electrocatalysts and RRDE measurements. J.P. participated in the calculation of the number of transferred electrons and FFT patterns. J.R. and Y.Y. planned and conducted HR-TEM experiments and analysis. J.R., M.P. and J.C. collaborated in writing the final manuscript.

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