Unveiling the Interlayer Spacing Dependence of Zn-Ion Storage Performance in Layered Vanadium Phosphates

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Abstract

Layered vanadium phosphate (VOPO$_4$·2H$_2$O) is reported as a promising cathode material for rechargeable aqueous Zn$^{2+}$ batteries (ZIBs) owing to its unique layered framework and high discharge plateau. However, its sluggish Zn$^{2+}$ diffusion kinetics, the low specific capacity and poor electrochemical stability remains a major issue in battery application. In this work, a group of phenylamine (PA)-intercalated VOPO$_4$·2H$_2$O with varied interlayer spacing (14.8, 15.6 and 16.5 Å) is synthesized respectively via a solvothermal method for cathode of aqueous ZIBs. The specific capacity is quite dependent on d-spacing in PA-VOPO$_4$·2H$_2$O system followed by an approximate linear tendency, and the maximum interlayer spacing (16.5 Å phase) results in a discharge capacity of 268.2 mAh·g$^{-1}$ at 0.1 A·g$^{-1}$ with a high discharge plateau of ~ 1.3 V and an energy density of 328.5 Wh·Kg$^{-1}$. Both of the experimental data and DFT calculation identify that the optimal 16.5 Å spacing can boost fast Zinc-ion diffusion with an ultrahigh diffusion coefficient of ~ $5.7 \times 10^{-8}$ cm$^2$·s$^{-1}$. The intercalation of PA molecules also significantly increases the hydrophobility in the aqueous electrolyte, resulting in the inhibiting the decomposition / dissolution of VOPO$_4$·2H$_2$O and remarkably improved cycling stability over 2000 cycles at 5.0 A·g$^{-1}$ with a capacity retention of ~200 mAh·g$^{-1}$. Our study provides a feasible solution on the sluggish Zn$^{2+}$ diffusion kinetics and poor cyclstability, and also shows a clear understanding on the interlayer chemistry of layered phosphates towards aqueous Zinc-ion storage.

Keywords:

phenylamine-intercalation; VOPO$_4$·2H$_2$O; interlayer spacing; diffusion kinetics; electrochemical stability; aqueous Zinc-ion battery
Nowadays, although rechargeable lithium-ion batteries (LIBs) are widely used for various portable electronics and electric vehicles, limited Li sources and severe safety issue are two main drawbacks of LIBs. With the rapid development of modern electronics and advanced equipments, beyond-lithium-ion (Na\(^+\), K\(^+\), Mg\(^{2+}\), Ca\(^{2+}\), Zn\(^{2+}\), Al\(^{3+}\)) batteries have recently attracted growing attention for energy storage systems.\(^1\) Among them, Zinc is abundant metallic elements in the earth crust far exceeding Li. More importantly, rechargeable aqueous Zinc-ion batteries (ZIBs) are based on aqueous electrolyte, resulting in much superior safety than that of alkali metal (Li\(^+\), Na\(^+\), K\(^+\)) based secondary batteries. Consequently, ZIBs have been considered as very promising candidates for next-generation battery systems.\(^2,3\)

Nevertheless, the revolution of aqueous ZIBs is still at the early stage and far from the prospective applications, which is mainly limited by the absence of suitable cathode materials especially with high energy density and sufficient long lifespan.\(^4\) The origin of this issue should be ascribed to the intrinsic difference between univalent lithium ion and divalent zinc ion. Since the similar radius of Li\(^+\) (0.076 nm) and Zn\(^{2+}\) (0.074 nm), the surface charge density of divalent Zn\(^{2+}\) is more than the twice of Li\(^+\), and the much stronger electrostatic interaction between Zn\(^{2+}\) and lattice framework of the electrode materials thus leads to sluggish diffusion kinetics and even structure deterioration.\(^5,6\)

Inorganic layered compounds, based on a typical layered structure and rich intercalation chemistry, have been particularly studied in metal-ion batteries. Especially, layered vanadium phosphate, VOPO\(_4\)\(_2\)·2H\(_2\)O, has been reported as an important cathode material for ZIBs accompanied with a Zn\(^{2+}\) intercalation / de-intercalation mechanism in its interlayer galleries.\(^7,9\) Remarkably, taking advantage of the enhanced ionicity of V-O bonds with the existence of PO\(_4\)\(^3-\), layered VOPO\(_4\)\(_2\)·2H\(_2\)O exhibites a much higher discharge plateau (1.1 - 1.2 V) than other V-based cathode materials.\(^10-12\) It should be emphasized that energy density is quite dependent on the discharge plateau of cathode, and such a merit of layered VOPO\(_4\)\(_2\)·2H\(_2\)O is highly desirable to achieve high energy density of ZIBs.\(^10\) Unfortunately, the specific capacity, rate capability of VOPO\(_4\)\(_2\)·2H\(_2\)O-based ZIBs is still unsatisfied due to the strong electrostatic interactions between Zn\(^{2+}\) and VOPO\(_4\) host with sluggish Zn\(^{2+}\) diffusion kinetics. On the other hand, the dissolution / decomposion behavior of VOPO\(_4\)\(_2\)·2H\(_2\)O in mild aqueous electrolyte generally results in very fast capacity / voltage fading during long-term cycling with a rather poor cyclstability. Although the dissolution / decomposion issue has been addressed by PO\(_4\)\(^3-\) addition and high salt concentration of the aqueous electrolyte to the shift the decomposion equilibrium and prevent dissolution\(^9\), the as-reported capacity of VOPO\(_4\)·2H\(_2\)O cathode is still less
than 170 mAh·g⁻¹, which is much smaller than its theoretical specific capacity (>300 mAh·g⁻¹). It is still a challenge to realize the superior performances for layered VOPO₄·2H₂O combining high discharge capacity, high rate capability and long cycling stability toward aqueous ZIBs applications.

Recent progress on lithium-/sodium-/magnesium-ion batteries unveils the interlayer spacing of layered materials as natural two-dimensional (2D) ion transport/diffusion channels. In principle, increasing the interlayer spacing of layered structure facilitates the ion transport by creating a lower energy barrier. Superior battery performance should be explored from the layered materials with larger interlayer spacing. As well, it is of great interest to make a fundamental understanding of the dependence of Zinc-ion storage behaviour on interlayer spacing for layered VOPO₄·2H₂O. However, it is still no report on the interlayer spacing modulation in layered vanadium phosphate up to date.

Herein, we found that the interlayer spacing of layered VOPO₄·2H₂O (VOP, 7.4 Å) can be successfully tuned by a phenylamine (PA) intercalated strategy through a facile solvothermal route. The crystal water in the VOPO₄·2H₂O was extracted from the interlayer space accompanied with the PA molecules intercalation. Three PA-intercalated VOPO₄ (labeled as PA-VOP) phases with identical d-spacing (16.5 Å, 15.6 Å and 14.8 Å) has been respectively achieved (Fig. 1a). We revealed an approximate linear dependence of the capacity dependence on interlayer spacing, and the maximum interlayer spacing (16.5 Å phase) results in a discharge capacity of 268.2 mAh·g⁻¹ at 0.1 A·g⁻¹ and an energy density of 328.5 Wh·Kg⁻¹. More importantly, the PA intercalation also significantly increases the hydrophobility in the aqueous electrolyte, resulting in inhibition of the decomposition/dissolution of VOPO₄·2H₂O and remarkably improved cycling stability over 2000 cycles with a capacity retention of ~200 mAh·g⁻¹ at 5.0 A·g⁻¹.

**Results**

**Controllable PA intercalation.** Uniform VOPO₄·2H₂O (JCPDS: 36-1472, labeled as VOP) nanoplates were first obtained by a conventional refluxing method reported previously. After a rapid solvothermal treatment of the as-prepared VOPO₄·2H₂O nanoplates in PA-contained isopropanol solvent at 60 ºC for 2 h, the sharp (001) peak of pristine VOPO₄·2H₂O substantially shifted to a lower angle (5.3°) with much decreased intensity (Fig. 1b), corresponding to an extreme increase in basal spacing up to 16.5 Å with simultaneously reduced crystallinity during the intercalation. Fourier transform infrared spectroscopy (FTIR) reveals the presence of υ(N-H), υ(C-H), δ(C-H) vibration mode and featured peak of benzene ring (Fig. 1c), demonstrating the successful intercalation of phenylamine molecules. It is
noteworthy that the characteristic peaks of crystal water in precursor VOPO$_4$·2H$_2$O completely disappear in FTIR spectrum after PA-intercalation. Thermogravimetric analysis (TGA) (Supplementary Fig. 1) identifies a weight loss of 25.3 % of PA-VOP up to 400 °C, corresponding to 0.6 unit phenylamine molecule contained in one unit PA-VOP structure. These results imply the phenylamine molecules replace the crystal water in VOPO$_4$·2H$_2$O interlayer to well maintain the initial layered framework as follows:

$$\text{VOPO}_4 \cdot 2\text{H}_2\text{O} + 0.6 \text{PA} \rightarrow \text{PA}_{0.6} @ \text{VOPO}_4 + 2\text{H}_2\text{O}$$

Scanning electron microscope (SEM) observations show an edge-crimped plate like morphology with lateral size of 5 to 20 µm and increased thickness feature compared to precursor VOP (Fig. 1d, Supplementary Fig. 2). However, transmission electron microscope (TEM) and atomic force microscopy (AFM) characterizations suggest a much smaller thickness of ~ 30 nm for our PA-VOP sample (Supplementary Fig. 3, Fig. 4). We consider that such a thickness difference should be caused by the possible liquid-exfoliation during the ultrasonic treatment of the sample preparation for TEM/AFM observation. Especially, one can clearly distinguish an interlayer spacing of ~ 1.6 nm from the cross-sectional high-resolution transmission electron microscope (HRTEM) image (Supplementary Fig. 4), which is well agreed with the one detected from the XRD data in Fig. 1b. Energy dispersive spectrum (EDS)-mapping profiles further clarify the uniform distribution of V, P, O, C, N elements after PA intercalation (Supplementary Fig. 4).
Fig. 1 Controllable PA intercalation in layered VOPO$_4$·2H$_2$O. a Scheme of capacity regulation for Zinc-ion storage via PA-intercalation engineering. b Powder XRD patterns and c FTIR spectra of VOP and PA-VOP samples, respectively. d Typical SEM image of PA-VOP (16.5 Å phase).

**Interlayer spacing dependence of Zn-ion storage.** Interestingly, we found the interlayer spacing of the PA-intercalated samples can be carefully tuned by solvothermal time. Three PA-VOP samples with different interlayer spacing of 14.8, 15.6 and 16.5 Å can be obtained under well-controlled solvothermal time of 60, 90 and 120 min, respectively (Fig. 2a). A further increase of solvothermal time compromises the interlayer spacing, indicating a saturated intercalation reaction at this threshold time (Fig. 2b). TGA curves suggests the weight loss of PA-VOP (14.8 Å), PA-VOP (15.6 Å) and PA-VOP (16.5 Å) is 19.1 %, 21.3 % and 24.5 % from 140 to 380 °C, respectively (Supplementary Fig. 5). Thus, it is rational that the increase of interlayer spacing is attributed to higher content of intercalated phenylamine molecules upon prolonging reaction time. Note that the as-obtained interlayer spacing of 16.5 Å in our work is the maximal one among all of the reported layered cathodes for aqueous ZIBs such as layered δ-MnO$_2$, PA-MnO$_2$ and Zn$_{0.25}$V$_2$O$_5$·nH$_2$O (Fig. 2c). SEM observation demonstrates the analogous plate-like morphology of the three PA-VOP samples other than a remarkably increased thickness caused by phenylamine insertion (Fig. 2d-f, Fig. 1d).
CR2032 coin cells were then assembled employing these PA-VOP samples as cathode material, Zn foil as anode and 2M \( \text{Zn}(	ext{CF}_3\text{SO}_3)_2 \) solution as aqueous electrolyte, respectively. Typical cyclic voltammetry (CV) curves of Zn // PA-VOP (16.5 Å phase) battery reveals a general active process in the first scan cycle, while the curves of the stabilized second and third cycle gives a pair of oxidation / reduction peak located at around 1.6 V and 1.3 V, respectively (Supplementary Fig. 6). It is should be noteworthy that PA molecules is generally inactive in mild aqueous electrolyte, and the electrochemical activity and capacity contribution should be mostly provided from VOPO₄ framework.²⁹-³¹ The galvanostatic charge discharge (GCD) curves also exhibit steady charge and discharge plateaus with well agreement with the voltage range in CV curves (Supplementary Fig. 6, 7). It has been reported that non-metal proton (H⁺) has also been recognized as charge carrier ions for some aqueous batteries.³² To clarify this, we compare the electrochemical performance on both aqueous and non-aqueous system (0.5 M \( \text{ZnSO}_4 \) / acetonitrile, Supplementary Fig. 8). No apparent difference has been observed between the CV peaks, thereby ruling out the possible contribution from proton intercalation in our system. \textit{Ex-situ} XRD at different voltages in the second cycle reveals a revisable Zn²⁺ intercalation / de-intercalation mechanism of the PA-VOP cathode (Supplementary Fig. 9). We found the specific capacity of the PA-VOP cathodes is quite dependent on its interlayer spacing. As shown in Fig. 2g, the pristine VOP (7.4 Å) and the PA-VOP samples (14.8 Å phase, 15.6 Å phase and 16.5 Å phase) delivers a discharge capacity of 138.4 mA·g⁻¹, 226.5 mAh·g⁻¹, 243.7 mAh·g⁻¹, and 268.2 mAh·g⁻¹, respectively. Similar enhancement tendency has been also observed for rate-capacity of series layered phosphates as shown in Fig. 2h. The resultant 16.5 Å phase delivers the maximum rate capability (Fig. 2i): a high specific capacity of 268.2, 254.8, 236.4, 220.7, 207.2, 187.5 mAh·g⁻¹ has been respectively realized at the current density of 0.1, 0.5, 1.0, 2.0, 5.0, 10.0 A·g⁻¹ with a recovered capacity of 253.3 mAh·g⁻¹ when the applied current density turned back to 0.1 A·g⁻¹.
Fig. 2 Interlayer spacing dependent battery performance. a XRD patterns of various PA-VOP samples synthesized under different temperature and solvent. b Dependence of interlayer spacing and solvothermal time. c Typical layered materials served as cathode of aqueous ZIBs. d-f SEM image of VOP, PA-VOP (14.8 Å phase) and PA-VOP (15.6 Å phase) samples, respectively. g GCD curves of VOP based ZIB and PA-VOP based ZIBs with adjustable interlayer spacing, respectively (14.8, 15.6 and 16.5 Å). h Cycle test at varied current density of VOP and PA-VOP cathodes with different interlayer spacing, severally. i GCD curves of VOP and PA-VOP cathode with different interlayer spacing at a current density of 5 A·g⁻¹, respectively.

We reveal the capacity-interlayer spacing dependence could be followed by an approximate linear trend as summarized in Fig. 3a. The specific capacity of 16.5 Å phase (268.2 mAh·g⁻¹) at the current density of 0.1 A·g⁻¹ is about two times higher than that of pristine VOP (138.4 mAh·g⁻¹) and also well above that of layered PPy-VOPO₄ cathode (~ 80 mAh·g⁻¹) reported before. Meanwhile, one can see a voltage plateau versus specific capacity diagram for mainstream cathodes of aqueous ZIBs reported in
recent years (V-based oxides, Mn-based compounds, Prussian Blue analogs, polyanion compounds, organics and other chalcogenides) shown in Fig. 3b. Benefited from \(d\)-spacing dependent performance as shown above, it is evident that rate-capacity of our optimal PA-VOP sample (16.5 Å phase) surpasses most traditional materials including Mn-based oxides (\(\alpha\)-MnO\(_2\), Mn\(_2\)O\(_3\)), V-based compounds (Na\(_3\)V\(_2\)(PO\(_4\))\(_3\), Zn\(_3\)V\(_2\)O\(_7\)(OH)\(_2\)·2H\(_2\)O\(_2\)) and Co\(_3\)O\(_4\) reported recently (Fig. 3c). Taking advantage of its high specific capacity of 268.2 mA·g\(^{-1}\) and prominent discharge plateau (~ 1.3 V), the 16.5 Å-phase PA-VOP renders a high energy density of 328.5 Wh·Kg\(^{-1}\) at a power density of 122.6 W·Kg\(^{-1}\), having apparent advantages in contrast with various popular cathode materials such as V\(_3\)O\(_7\)·nH\(_2\)O, \(\alpha\)-MnO\(_2\), \(\delta\)-MnO\(_2\), ZnMn\(_{1.86}\)O\(_4\), ZnHCF\(_{43}\) and CuHCF (Fig. 3d). This excellent energy and powder densities promote the application in flexible and wearable electronic devices (Insert in Fig. 3d).

**Fig. 3** Performance comparison. a Dependence of specific capacity and interlayer spacing of various VOPO\(_4\)-based cathodes at the current density of 0.1 A·g\(^{-1}\). b Voltage plateau versus specific capacity for various mainstream cathodes recently reported and our PA-VOP (16.5 Å) cathode. c Comparison of the rate-capacity between our PA-VOP cathode and the other recently reported cathodes.
for aqueous ZIBs.\textsuperscript{25,40,46,53-54} d Ragone plot of PA-VOP cathode and a series of other cathode materials reported previously.\textsuperscript{27,35,41,43,55-56} Insert is a solid-state battery constructed from the PA-VOP @ CFC cathode driving a group of LEDs.

**Reaction/diffusion kinetics mechanism.** In order to clarify the aforementioned $d$-spacing dependent battery-performance, corresponding study on electronic conductivity and Zinc-ion diffusion kinetics is further conducted. Four-probe conductivity test shows a 2 ~ 7 times higher electronic conductivity of PA-VOP (16.5 Å) in contrast to that of pristine VOP with pressure increased, which is mainly attributed to the intrinsic high conductivity of phenylamine (Fig. 4a). CV curves at varied scan rates are employed to distinguish the capacitive / ion diffusion contribution by analyzing the dependence of the logarithm of peak current density and the scan rate (Supplementary Fig. 10). Theoretically, the peak current ($i$) and sweep rate ($v$) in CV curve follows the rule as:\textsuperscript{57,58}

$$i = a v^b$$  \hspace{1cm} (1)

Where $a$ and $b$ are adjustable values. $b$ value of 0.5 indicates a semi-infinite diffusion mechanism, while $b$ value of 1.0 suggests capacitive behavior. In our case, $b$ value for the predominant peaks of VOP, PA-VOP (14.8 Å phase) and PA-VOP (16.5 Å phase) cathode is determined to be 0.63, 0.73 and 0.76, respectively (Supplementary Fig. 10). Apparently, the capacitive contribution in PA-VOP (16.5 Å phase) cathode is 44.8 %, 49.6 %, 56.7 %, 60.4 %, and 65.9 % at the scan rate of 0.1, 0.2, 0.5, 0.8, and 1.0 mV·s$^{-1}$, respectively. These values is much higher than that in VOP cathode (28.4 %, 33.2 %, 38.9 %, 42.8 %, and 47.5 % at 0.1, 0.2, 0.5, 0.8, and 1.0 mV·s$^{-1}$, respectively) (Fig. 4b, c).

On the other hand, Zinc-ion diffusion kinetics is further identified by galvanostatic intermittent titration technique (GITT) (Fig. 4d). Strikingly, an ultrahigh Zinc-ion diffusion coefficient of $\sim 5.7 \times 10^8$ cm$^2$·s$^{-1}$ has been detected in 16.5 Å phase, nearly 5 orders of magnitude higher than that of pristine VOP cathode ($6.2 \times 10^{-13}$ cm$^2$·s$^{-1}$) and also much higher than that of conventional cathode materials reported recently including V$_2$O$_5$, ZnMn$_2$O$_4$, Zn$_3$V$_2$O$_7$(OH)$_2$·2H$_2$O, and K$_2$V$_8$O$_{25,37,41,59}$ (Fig. 4e). This drastic increase demonstrates a great improvement in terms of Zn$^{2+}$ intercalation / de-intercalation kinetics by an enlarged 2D interlayer channel as illustrated in Fig. 4f. In principle, large Zinc-ion diffusion coefficient is generally observed in ultrathin samples due to the shortened effective diffusion path and larger activated surface, which drastically facilitate Zn$^{2+}$ diffusion and charge transfer.\textsuperscript{28,60,61} Our previous study also confirms fast Zn-ion diffusion ability in 6.0-nanometer ultrafine spinel oxide
nanodots. However, in the present study, the as-obtained 16.5 Å phase show a significant increase on thickness compared with the pristine VOP sample (Fig 2d-f). Therefore, the thick samples display much faster kinetics than that of thin VOP nanoplates unusually. Such an phenomenon should indicate the Zinc-ion transport in our layered phosphate series is dominated by interlayer diffusion on account of the enlarged interlayer spacing would provide high-efficiency 2D ion diffusion / hopping channels.

Besides, density functional theory (DFT) based first-principle simulation is conducted for a deeper understanding of the greatly improved diffusion kinetics. Climbing-image nudged elastic band (CI-NEB) method is adopted to reveal the possible diffusion paths and the corresponding energy barrier of zinc-ion. The interlayer site on the top of the corner-shared oxygen atom of [PO₄] tetrahedron and [VO₆] octahedron gives an optimal absorption energy of 0.07 eV, which can be determined as the potential adsorption site (called as C site) for zinc-ion diffusion. The as-optimized diffusion path denotes hopping process through the right above site of [VO₆] octahedron and [PO₄] tetrahedron between the adjacent C sites (Fig. 4g, h). Subsequently, an interlayer-dependent activation energy decrease tendency is derived based on such a diffusion path. The energy barrier for zinc-ion diffusion in VOP bulk (7.4 Å) is calculated to be 0.13 eV, and dramatically decreases to a much lower value (~0.03 eV) until a threshold interlayer spacing of 9.7 Å. Further increase of spacing gives rise to a slight change of energy barrier, and it finally decrease to ~ 2.3 × 10⁻⁴ eV for PA-VOP (16.5 Å phase) sample. Consequently, the greatly enhanced diffusion kinetics, predominant capacitive contribution and high electronic conductivity in PA-VOP samples provide the essential insight for the interlayer spacing dependent Zn²⁺ storage performance.
Fig. 4 Reaction/diffusion kinetics mechanism. 

a Electronic conductivity test under different pressure.

b Capacitive contribution of VOP, PA-VOP (14.8 Å phase) and PA-VOP (16.5 Å phase), respectively. 

c CV curve containing the schematic of capacitive and diffusion contribution of PA-VOP (16.5 Å phase) cathode at the scan rate of 0.1 mV·s⁻¹ 

d GITT curve and e the calculated diffusion coefficient of VOP, PA-VOP (14.8 Å phase) and PA-VOP (16.5 Å phase), respectively. 

f Schematic illustration of Zn²⁺ diffusion kinetics in PA-VOP and VOP cathode. 

h Zinc-ion diffusion path scheme in top view and side view mode. 

i Calculated energy barrier curve of the PA-VOP samples with different interlayer spacing.
**Cyclability improvement.** It is known that layered VOPO$_4$$\cdot$$n$H$_2$O cathode undergoes severe capacity/voltage fading with rather poor cyclability originated from the decomposition/dissolution of VOPO$_4$$\cdot$$n$H$_2$O in mild aqueous electrolyte. Interestingly, we found the chemical stability and electrochemical reversibility was significantly enhanced after the PA intercalation. Compared to aqueous Zn(SO$_4$)$_2$ electrolyte, superior lifespan was observed in Zn(CF$_3$SO$_3$)$_2$ electrolyte due to the better desolvation effect of bulky (CF$_3$SO$_3$)$^-$(versus (SO$_4$)$_2$)$^-$(Supplementary Fig. 11). One can see the Zn // 2M Zn(CF$_3$SO$_3$)$_2$ // PA-VOP (16.5 Å phase) battery delivers an initial specific capacity of 200.9 mAh·g$^{-1}$ and a steady cycle stability of 92.3 % capacity retention after 2,000 cycles under a high current density of 5.0 A·g$^{-1}$. In contrast, the VOP based battery just exhibits an inferior capacity of ~ 60 mAh·g$^{-1}$ with rapid decay in the first 300 cycles (Fig. 5a). To clarify the actual mechanism, *ex-situ* XRD characterizations of VOP and PA-VOP (16.5 Å phase) cathode before and after 300 cycles are carried out as shown in Fig. 5b. Apparently, the characteristic diffraction peaks of VOP cathode completely disappear after 300 cycles, indicating the phase conversion of VOP during the long-term cycling. In contrast, the PA-VOP (16.5 Å phase) cathode well maintained the initial phase without any impurity diffraction peaks. SEM observation also reveals PA-VOP (16.5 Å phase) sample well maintain the initial plate-like morphology compared to the drastic aggregation of VOP cathode after 300 cycles (Fig. 5c). All results demonstrate the greatly enhanced structural reversibility of our PA-VOP sample after PA intercalation.

To further probe the origin of the remarkable difference on long-lifespan, we compare the chemical stability of both VOP and PA-VOP samples in aqueous 2M ZnSO$_4$ electrolyte. As shown in Fig. 5d, it is clear that VOP sample generally disappears in the bottom of solution followed by a typical color change into deep red after 5 days due to the decomposition/dissolution of VOPO$_4$$\cdot$$n$H$_2$O into VO$_x$(s) and PO$_4$$^{3-}$. However, the PA-VOP (16.5 Å phase) sample still floats on the liquid surface with much better stability after 5 days (Fig. 5d). Such a result inspires our consideration on the difference on surface-wettability of these two samples. Accordingly, Fig. 5e compares the water-based contact angle on the surface of pressed powder of VOP and PA-VOP sample. The PA-VOP and VOP sample shows a contact angle of 56.7° and 9.8° respectively, demonstrating the surface is much more hydrophobic after PA intercalation. Rationally, this hydrophobic surface plays a decisive role in the inhibiting the decomposition/dissolution of VOPO$_4$ in aqueous electrolyte, thus leading to much improved long lifespan of the as-constructed
ZIBs. Most recently, Sun et al. reported that the decomposition/dissolution of VOPO$_4$·2H$_2$O can be prevented by PO$_4^{3-}$ addition and high salt concentration of the aqueous electrolyte$^9$. Such a strategy promotes a long-term cycling over 500 cycles with a stable capacity of only 90 mAh·g$^{-1}$ at 2.0 A·g$^{-1}$ but also increases the battery cost. In contrast, our strategy by PA-intercalation requires no addition of any reagent/salt addition, and realizes much superior cycling stability over 2000 cycles with a capacity of ~200 mAh·g$^{-1}$.

**Discussion**

In summary, we developed a controllable phenylamine-intercalated strategy for layered vanadium phosphates with various interlayer spacings (14.8 Å, 15.6 Å and 16.5 Å). The crystal water in the VOPO$_4$·2H$_2$O was extracted from the interlayer space accompanied with the PA intercalation. The specific capacity is quite dependent on $d$-spacing in PA-VOP system followed by an approximate linear tendency, and the interlayer spacing (16.5 Å) results in a maximum capacity of 268.2 mAh·g$^{-1}$ at 0.1 A·g$^{-1}$ and an energy density of 328.5 Wh·Kg$^{-1}$. Both of the experimental data and theoretical calculation identify that the enlarged spacing can boost fast Zinc-ion diffusion with an ultrahigh diffusion coefficient of $\sim 5.7 \times 10^{-8}$ cm$^2$·s$^{-1}$. The intercalation of PA molecules also significantly increases the hydrophobility in the aqueous electrolyte, inhibiting the decomposition/dissolution of VOPO$_4$·2H$_2$O and remarkably improved long-term cycling stability over 2000 cycles at 5.0 A·g$^{-1}$ with a capacity of ~200 mAh·g$^{-1}$. Our study provides a feasible solution on the sluggish Zn$^{2+}$ diffusion kinetics and poor stability of layered VOPO$_4$·2H$_2$O, and also shows a clear understanding on the interlayer chemistry towards aqueous Zinc-ion storage. Similar strategy may be extended to other layered inorganic compounds with superior ion diffusion kinetics and better battery performance.
Fig. 5 Hydrophobicity improvement. a Long-term cycle test of Zn // VOP and Zn // PA-VOP (16.5 Å phase) battery using 2M Zn(CF$_3$SO$_3$)$_2$ aqueous electrolyte. b XRD patterns and c SEM images of the VOP and PA-VOP (16.5 Å phase) cathode before and after 300 cycles, respectively. d Optical photographs of the as-dispersed VOP and PA-VOP (16.5 Å phase) powder in 2M ZnSO$_4$ aqueous electrolyte at different times. e Water-based angle contact experiment of VOP and PA-VOP (16.5 Å phase) powder.
Methods

Materials synthesis. VOPO₄·2H₂O sample is synthesized based on a reported refluxing method with some modifications. Typically, 4.8 g V₂O₅, 26.6 mL H₃PO₄ and 115.4 mL DI water is added to a three-necked flask, respectively, followed by 30 min of magnetic stirring with the addition of 10 mL of HNO₃ to stabilize the oxidation state of vanadium. The mixture is then heated at 110 °C for 16 h. Yellow precipitate is collected from the bottom after cooling down to room temperature, the yellow precipitate is collected by centrifugation, washed with DI water and acetone for 3 times, respectively. The resulting bright yellow VOP sample was dried at 60 °C in the vacuum oven. To introduce phenylamine molecules into the interlayer spacing of VOP, 0.3 g of as-synthesized VOP is mixed with 30 mL of isopropanol in 50 mL Teflon-lined strain-less steel autoclave and followed by the addition of 5 mL phenylamine. The mixture is stirred for 15 min and then heated in an oven at 60 °C for a short time (60, 90 and 120 min) to obtained phenylamine intercalated VOPO₄ (PA-VOP) with varied interlayer spacing (14.8, 15.6 and 16.5 Å), respectively. The black brown sediment in the bottom is then collected and washed with ethanol for 3 times. The PA-VOP powder is finally obtained after vacuum drying at 60 °C for 6h.

Materials characterization. The structure morphology and structure characterization are characterized by scanning electron microscopy (SEM, Phenom Pro X), transmission electron microscopy (TEM, Philips CM 200 FEG Field Emission Microscope) and X-ray diffraction (XRD, Bruker D8-A25 diffractometer using Cu Kα radiation (λ = 1.5406 Å)). The high resolution transmission electron microscopy (HRTEM) and energy dispersive spectrum mapping (EDS-Mapping) is carried along with the TEM measurement to gather the detailed information of phase structure and chemical element distribution of the sample, respectively. Further information of the chemical bonding and microscopic condition is revealed by means of Fourier transform infrared spectroscopy (FTIR, BRUKER TENSOR II) and thermogravimetric analysis (TGA, SDT Q600), respectively. Water-based angle contact experiment (JY-82) is developed based on pressed powder of PA-VOP and VOP samples. Electronic conductivity test is carried out on a four-probe conductivity tester (ST2253y).

Electrochemical measurement. PA-VOP and VOP based cathodes of aqueous ZIBs is severally fabricated by mixing as-prepared samples, acetylene black and polyvinylidene fluorides (PVDF) on the basis of a mass ratio of 7:2:1 with the addition of 1-methyl-2-pyrrolidinone (NMP), then the uniform slurry is coated onto a piece of 304 stainless steel foil and drying at 100 °C for 12 h in a vacuum oven.
The slurry-coated foil is cut into Φ15 mm electrode as cathode, while the zinc foil washed with ethanol and glass fiber membrane is used as the anode and separator, respectively, and 2M Zn(CF$_3$SO$_3$)$_2$ is prepared as the electrolyte. The CR-2032 cell was assembled in air using the beforehand electrodes and other relevant components. LAND battery test system (CT2001A) was employed to evaluate the electrochemical performance of the battery: galvanostatic charge-discharge (GCD), rate capability and long-term cycle performance. Cyclic voltammetry (CV) test at different scan rate is performed on electrochemical workstation (CHI660E). Galvanostatic intermittent titration technique (GITT) is performed under a modified GCD mode, in which an operation period includes two parts: a charge / discharge procedure last for 10 min at 0.05 A·g$^{-1}$ and a followed pause time for 10 min.

**Fabrication of the flexible all-solid-state battery.** The quasi solid flexible zinc-ion battery was fabricated using aforementioned slurry-coated carbon fiber cloth (CFC) as the cathode (mass loading: 2 mg·cm$^{-2}$), CFC with a layer of zinc electrodeposited coating as the anode, a piece of filter paper as the separator and 2M Zn(ClO$_4$)$_2$ @ polyvinyl alcohol (PVA) as the electrolyte. The electrolyte was prepared by adding 3 g PVA to 30 mL 2M Zn(ClO$_4$)$_2$ aqueous solution little by little with continuous stirring, followed by oil bath treatment at 80 °C for 2h and gathering the thick sample when it came to the room temperature. The electrochemical deposition of zinc layer on the CFC was executed on CHI660E using potentiostatic model at -0.7 V (vs. Zn$^{2+}$/Zn) for 2000 s. The ultimate flexible all-solid-state battery was assembled and sealed with polyimide (PI) film to form a stable sandwich structure.

**Electrochemical calculation method.** The calculation method for capacity contribution of the cathodes is based on the following equation:

\[ I_p = C_1 v + C_2 v^\frac{1}{2} \]  

(1)

where $I_p$ (A·g$^{-1}$) is the peak current density at different scan rate, $v$ (mV·s$^{-1}$) is the specific scan rate, $C_1$ and $C_2$ are the corresponding constant factors of the capacity contribution of surface pseudocapacitive effect and battery-type effect, respectively.

With a deformation of the above equation, the specific contribution rate of different internal mechanisms can be solved according to the following equation:

\[ \frac{I_p}{v^\frac{1}{2}} = C_1 v^\frac{1}{2} + C_2 \]  

(2)

The specific energy density (Wh·kg$^{-1}$) and average specific power density (W·kg$^{-1}$) of the batteries is calculated in terms of the following equations:

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Where $E_s$ is the calculated specific energy density (Wh·kg$^{-1}$), $P_s$ is the average specific power density (W·kg$^{-1}$); $C_s$ (mAh·g$^{-1}$) is the specific capacity of the battery, $V_0$ and $V_1$ is the voltage lower limit and voltage upper limit of discharge procedure, respectively, and $t$ is the discharge time (h). All the parameters calculated are based on the mass loading of the active materials (VOP or PA-VOP).

Diffusion coefficient ($D_{Zn^{2+}}$) of Zinc-ion can be experimentally calculated by GITT method in terms of the following equation:

$$D = \frac{4}{\pi \tau} L^2 \left( \frac{\Delta E_s}{\Delta E_t} \right)^2$$  \hspace{1cm} (5)

Where $D$ is the diffusion coefficient of Zinc-ion, $\tau$ is the relaxation time of current pulse, $L$ is diffusion length which is approximate to the thickness of coated slurry, $\Delta E_s$ and $\Delta E_t$ is the voltage change produced by current pulse and the galvanostatic charge / discharge, respectively.

**Simulation details.** The modeling in this study was performed in the framework of the DFT as implemented in the Vienna Ab initio Simulation Package (VASP). The functional of Perdew-Burke-Ernzerhof based on generalized gradient approximation (GGA) was applied to describe the exchange-correlation energy. In addition, the zero damping DFT-D3 dispersion correction method of Grimme was accounted for VdW interaction in the system. A vacuum space of 15 Å was adopted. The plane-wave cutoff energy was set to be 480 eV, and the k-mesh was determined to be $7 \times 7 \times 1$ according to the convergence test, which makes the energy accuracy within $1.0 \times 10^{-3}$ eV atom$^{-1}$. Finally, a double-layered VOPO$_4$ model was constructed, with the corresponding interlayer spacing obtained from our previous XRD and HRTEM analysis, thereby the diffusion of zinc ion between VOPO$_4$ layers was simulated using the climbing-image nudged elastic band method.

**Data Availability**

The authors declare that all the relevant data are available within the paper and its Supplementary Information file or from the corresponding author on reasonable request.
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Author contributions
L. F. H. supervised the project. L. F. H., Z. Y. W. designed the sample preparation and executed the electrochemical screening experiments. C. J. L. performed DFT calculations and simulations. F. Y. participated in the battery performance measurement and analysis. Y. L. and
R. L. P. conducted material characterizations on SEM, TEM, XRD, XAS, Raman characterizations. Z. M. S. discussed the results and interpreted the data. All authors contributed to the writing of the paper.

Competing interests

The authors declare no competing interests.