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Puiki Leung, Lin Zeng, Tianshou Zhao, Lei Wei *et al.*
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In-plane gradient design of flow fields enables enhanced convections for redox flow batteries

Lyuming Pan,^{†a} Jianyu Xie,^{†a} Jincong Guo,^{†a} Dongbo Wei,^a Honghao Qi,^a
Haoyao Rao,^{†a} Puiki Leung,^{*b} Lin Zeng,^{†a} Tianshou Zhao^{*a} and Lei Wei^{†a}

In the realm of redox flow batteries, the flow field plays a vital role in influencing the overall performances of the redox flow batteries. Inspired by human behavior, an in-plane gradient flow field design featuring a gradient decrease in channel width from the inlet to the outlet is proposed in this work. A three-dimensional multi-physical simulation model was utilized to investigate the transport behaviors and overall battery performance associated with novel flow field configurations. It was indicated that the novel in-plane gradient design can enhance the under-rib convections of the electrolyte in the downstream regions near the outlet, leading to improved uniformity of the active species' distribution over porous electrodes. Consequently, this enhancement substantially reduces concentration polarization losses of redox flow batteries. The maximum power density and rated current density of the proposed design are 553.2 mW cm⁻² and 270.1 mA cm⁻², which are 74.5 mW cm⁻² and 8.3 mA cm⁻² higher than conventional design. These results substantiate the benefits of employing the proposed flow field for achieving high-performance battery designs. Meanwhile, due to its straightforward, efficient, and easily scalable design mechanism, this novel flow field shows great promise for engineering applications of redox flow batteries.

1. Introduction

Future sustainability and carbon neutrality hinge on the advancement of renewable energy technologies.^{1,2} Solar and wind power are among the most popular choices. Yet, their intermittent and unpredictable nature necessitates the urgent development of large-scale energy storage systems to assimilate renewable energy sources.^{3,4} Electrochemical energy storage systems, such as vanadium redox flow batteries (VRFBs), are among the most promising technologies to address this issue due to their inherent safety, recoverable crossover, and scalability. The VRFB consists of the power unit and electrolyte reservoirs,^{5,6} with the former dictating system power and the latter determining capacity. Consequently, the power and capacity of the system are decoupled, and like the pumped storage and compressed air systems, making it well-suited for long-term energy storage in the era of carbon neutrality.

The power unit of VRFBs comprises essential components, namely the flow fields, electrodes and a membrane. The electrolytes are recirculated from external reservoirs and into the power unit by a circulation pump. The active species

dissolved in the aqueous electrolytes are delivered to the porous electrodes through the flow fields, initiating electrochemical redox reactions. Conventionally, VRFBs adopt a flow-through structure, wherein the electrolyte is directly pumped through the electrodes, leading to significant pressure drop and pumping losses.^{7,8} Typically, more than 3 mm thick electrodes were used to minimize pumping losses, but this also causes higher internal ohmic resistance, thereby leading to suboptimal overall performances (lower than 150 mA cm⁻² at an energy efficiency of 80%).

Since then, researchers have devised flow-by designs in which flow fields are incorporated alongside the porous electrodes. Conventional designs include parallel flow fields (PFFs),⁹ serpentine flow fields (SFFs)^{10,11} and interdigitated flow fields (IFFs).¹² In these flow-by setups, the electrolyte flows through the channels and is transported into the electrodes. Adopting thinner electrodes in channel-based cells reduces internal resistance and enhances cell functionality.¹³ Different mass transfer properties are associated with various flow pattern designs. PFF is the design with the lowest pressure drop of inlet and outlet since the electrolyte flows parallel between various flow channels,¹⁴ but it also results in poor mass transfer. Therefore, SFFs and IFFs have gained greater attention in recent research. While the inlet and outlet of the IFF are not connected, the electrolyte must be pumped through the porous electrodes. IFFs, however, experience high-pressure drops when used with densely compressed electrodes. Conversely, SFFs have interconnected flow channels

^a Shenzhen Key Laboratory of Advanced Energy Storage, Department of Mechanical and Energy Engineering, Southern University of Science and Technology, Shenzhen 518055, China. E-mail: zengl3@sustech.edu.cn, zhaots@sustech.edu.cn, weil@sustech.edu.cn

^b Key Laboratory of Low-Grade Energy Utilization Technologies and Systems, MOE, Chongqing University, Chongqing 400030, China. E-mail: p.leung@cqu.edu.cn

[†] These authors contributed equally.



from the inlet to outlet, enabling the electrolyte to flow through both the channel and porous electrode, which minimizes the pressure drop but may lead to uneven distribution. Researchers have focused on modifying porous electrodes for balancing mass transfer and active sites.^{15–18} Li *et al.*¹⁹ proposed a gradient-pore-oriented graphite felt electrode that improves electrochemical activity and enhances mass transport from the nano- to the micro-scale. This gradient-pore-oriented electrode exhibits an energy efficiency as high as 79.74% at 200 mA cm⁻² on VRFBs, suggesting great potential applications in high-performance flow batteries. Although improved VRFB performance by electrode surface modification and microstructures is intuitive, mass transfer still also relies mainly on flow field optimization.

All three flow patterns and the modified ones are directly derived from fuel cell technology and introduced onto VRFBs. For example, Xu *et al.*²⁰ first proposed a convection-enhanced serpentine flow field, which brings about larger pressure differences between adjacent flow channels and thereby enhances in-plane forced flow through the electrode porous layer. Their pioneering design concept has been utilized in VRFBs and presents significantly excellent effects.^{21–23} Typically, optimization of the flow-field parameters, such as channel width, depth, and opening ratio, has not adequately considered the distinct structure of VRFBs and the properties of the active species. Unlike fuel cells, the component adjacent to the flow field is the gas diffusion layer responsible for transporting reactants and products. In VRFBs, on the other hand, the porous electrodes adjacent to the flow field not only facilitate the transport of active species but also serve as active sites for the redox reaction. More importantly, the reactants in fuel cells, such as hydrogen and oxygen, have high diffusion rates, an order of magnitude higher than the reactive ions in flow batteries using liquid electrolytes. While electrochemical reactions in fuel cells mainly occur in the catalytic layer adjacent to the membrane, VRFB reactions primarily take place in the porous electrode beneath the flow channels' ribs of flow channels. Additionally, the reaction in fuel cells involves a gas–liquid phase transition, whereas the VRFBs only involve a single liquid-phase reaction.

In light of these differences in working mode and system characteristics between fuel cells and flow batteries, direct utilization of fuel-cell flow-field designs in flow batteries may give rise to various issues. Hence, it is necessary to optimize the flow field according to the working modes of the flow battery and the requirements of active species.^{24–28} For instance, Xu *et al.*²⁹ explored the design of blocked serpentine flow channels for VRFBs to achieve a more homogeneous distribution of electrolyte flow and enhance active mass transport within the electrodes, thus improving the overall battery performance. Considering the effects of different block heights on reactant distribution and battery voltage, they suggested that the optimized height of 0.14 mm led to the highest net discharge power at 16.73 W. Zheng *et al.*³⁰ proposed a separate serpentine flow field with two flow channels—one located on the porous electrode next to the membrane and the other on a carbon plate. This design features a combined advantage of serpentine and interdigital patterns, guiding the electrolyte along a U-shaped path within a single

channel, passing through porous electrodes rather than channels between them. This design improves the uniformity of the active species' distribution, even with relatively thick electrodes at 5 mm. While these designs demonstrate promising performances, their sophisticated performance improvement mechanisms pose challenges for promotion and scaling-up.

Herein, inspired by the towel-twisting phenomenon, an in-plane gradient flow field (IGFF) is proposed for redox flow batteries. The channel width is designed to decrease gradually from the upstream region at the inlet to the downstream zone at the outlet, deviating from the conventional approaches. This unique gradient change in channel width corresponds to the decrease in reactant concentration. Therefore, the adoption of IGFF enhances the under-rib convection in the near-outlet region, thereby promoting the mass transfer capability of active species throughout the entire porous electrode. Utilizing a 3D multi-physical simulated model, the introduction of IGFF can improve the average reactant concentration, distribution uniformity and current density on the porous electrodes. The maximum power density and rated current density of the proposed design are 553.2 mW cm⁻² and 270.1 mA cm⁻², which are 74.5 mW cm⁻² and 8.3 mA cm⁻² higher than the conventional design. These results clearly demonstrate the advantages of the proposed IGFF in constructing high-performance VRFB. Meanwhile, the straightforward, efficient, and easily scalable design mechanism, opens up possibilities for its application in engineering applications of VRFB stacks.

2. Results and discussions

2.1. The in-plane gradient design mechanism

Convection and mass transport phenomena are common in nature and human life and play crucial roles in various processes. An example is the water retention phenomenon of a rolled towel of uniform diameter. The presence of numerous pores between the fibers of the towel enables the water to remain in a stable state. This stability is achieved when a towel is rolled into a uniform-diameter tube. It is worth noting that altering objects' parameters (diameter, width, *etc.*) is a practical and effective means of modifying hydraulic pressure and fluid velocity. When it is necessary to extract the held water, humans tend to tighten one end of the towel tube to reduce its diameter, thereby squeezing out water (Fig. 1a). This is a simple example of human behavior to effectively utilize diameter changes for enhancing convections. It can also be observed in other natural objects, such as changes in river width, changes in pipeline diameter, and so on. The human behavior is considered applicable to flow field optimization to enhance the mass transport of active species for VRFB.

The flow channels of conventional SFF designs in VRFB maintain the same width throughout the entire region. It contains multiple long flow channels and U-shaped interconnections to form a connection from the inlet to the outlet, as shown in Fig. 1b. The SFF's repeating structure results in an average hydraulic pressure distribution. Therefore, the concentration of active



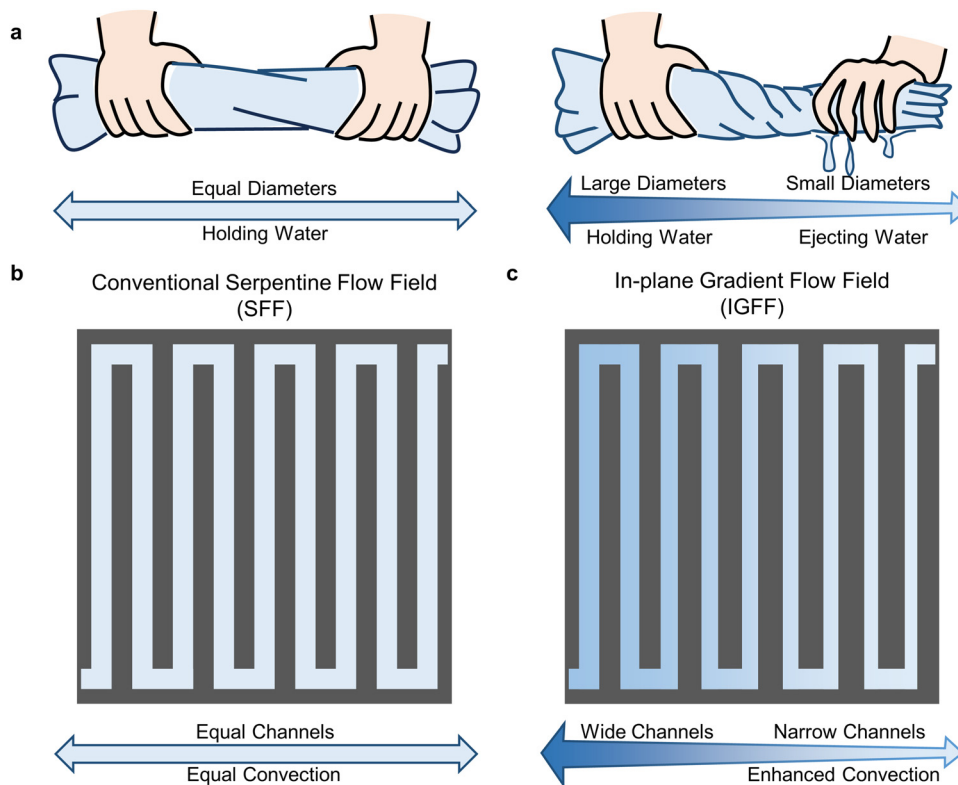


Fig. 1 The (a) in-plane gradient design principles and geometric patterns of (b) SFF and (c) IGFF.

species decreases sequentially from inlet to outlet, accompanied by electrochemical reactions. This non-uniformity may become more significant as the flow rate decreases and the current density increases.^{31,32} To address this issue, it is necessary to increase the hydraulic pressure drop and under-rib convection in the downstream region near the outlet. The human behavior mentioned above inspires us to enhance the electrolyte convection into the porous electrode by adjusting the channel width. As shown in Fig. 1c, the channel width gradually decreases in IGFF, while the flow channels' number, direction, and arrangement remain unchanged compared to SFF. In such cases, the decrease in reactant concentration in the electrolyte can be kept

consistent. The difference between IGFF and SFF can be analyzed and elucidated by comparing hydraulic pressure, fluid velocity, and the electrochemical performance of VRFBs.

2.2. The distributions of pressure, velocity and concentration

In this work, a 3D multi-physical simulation model is established to evaluate the battery performance with different flow fields, which involves the coupling mechanisms of fluid mechanics, electricity and mass transfer. Under the premise of achieving grid independence and model validation, as shown in Fig. 2, the 3D model, which may be used to compare SFF with IGFF, is considered reasonably accurate for VRFBs. Specifically, the

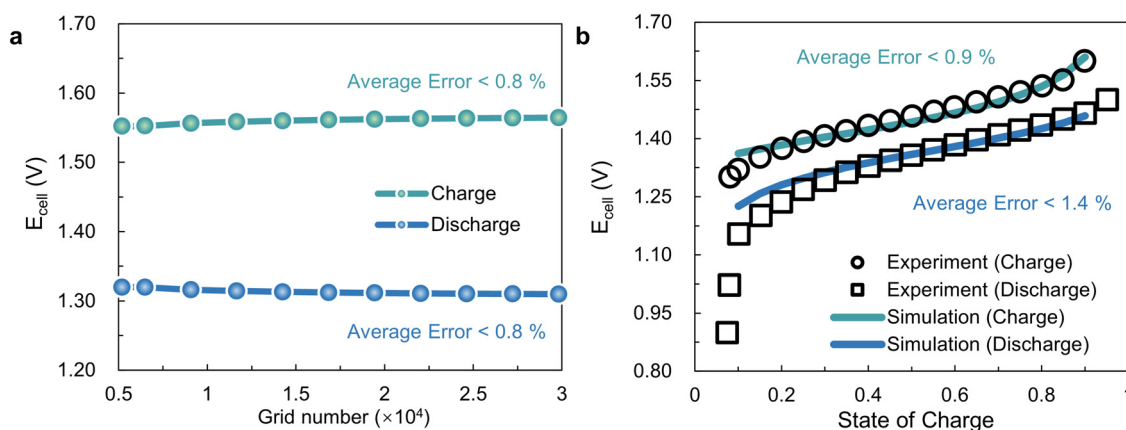


Fig. 2 The (a) grid independence and (b) validation of the 3D multi-physical simulated models.



computed inaccuracy of cell voltages (E_{cell}) is less than 0.8% when the grid number of the 3D models is set to at least 1×10^4 (Fig. 2a). Additionally, the average discrepancies between the actual E_{cell} and the simulated ones for a VRFB applied under a current density of 50 mA cm^{-2} and a specific flow rate of $5 \text{ mL min}^{-1} \text{ cm}^{-2}$ are less than 0.9% and 1.4%, respectively (Fig. 2b). The average validated error for discharging is greater than that for charging, mainly due to significant polarization at a low state of charge regions (SOC < 0.3). This might be due to inaccurate estimation of physical parameters, such as diffusion coefficients or reaction rate coefficients, as reported.³³ Otherwise, the conducted 3D models compare well with experiments for most operating conditions (medium SOC), which are reliable for comparing SFF and IGFF.

Utilizing the aforementioned 3D model as a foundational framework, the intricate interplay of multiple physical domains within VRFB is comprehensively addressed. The fluid dynamics of the electrolyte, spanning both the flow field and porous medium, is rigorously formulated and resolved by invoking the venerable Navier–Stokes equation and Darcy's law, respectively (Fig. 3).³⁴ The innovative in-plane gradient configuration profoundly influences the hydraulic pressure distribution within the flow channel, thereby amplifying the pressure gradient near-outlet region. Specifically, in the case of SFF, the hydraulic pressure differential between adjacent flow channels is maintained at a uniform 55 Pa level (Fig. 3a1). Contrasting this, the IGFF exhibits a remarkable alteration, with the pressure differential escalating to 140 Pa between the final adjacent channels while maintaining the 55 Pa disparity between the initial adjacent channels (Fig. 3a2). The hydraulic pressure distribution across the flow field inherently governs the fluid velocity profile within the porous electrode (Fig. 3b). The fluid velocity

profile demonstrates periodicity and uniformity beneath the ribs of SFF (Fig. 3b1), mirroring the hydraulic pressure distribution. In the context of IGFF, however, a noteworthy evolution becomes evident, as the intensity of under-rib convections gradually increases from the proximal inlet to the adjacent outlet regions (Fig. 3b2). Cross-sectional views (Fig. 3c) offer a lucid visualization of the discernible correlation between hydraulic pressure gradients within the flow channels and the resultant under-rib convection patterns within the porous electrode. The innovative in-plane gradient design, inspired by principles like those observed in natural systems, manifests a tangible enhancement in convective processes, thereby achieving the coveted goal of convective augmentation.

Fluid velocity exerts a profound influence on the spatial distribution of active species, particularly the under-rib convections (Fig. 4). The operational dynamics of vanadium electrolytes within the battery are characterized by a cyclic inflow and outflow, dynamically responding to alterations in the SOC. These dynamic responses manifest as observable fluctuations in the concentrations of reactants at the inlet and outlet of the flow field (Fig. 4a). As an illustrative example, consider the concentrations of VO_2^+ ($c\text{VO}_2^+$, as the reactants at the positive side for the discharge process) at the inlet and outlet are 0.85 M and 0.72 M, respectively. Remarkably, whether adopting SFF (Fig. 4a1) or IGFF (Fig. 4a2), these inflow and outflow concentrations are equivalent. However, the spatial distribution of $c\text{VO}_2^+$ within the porous electrode presents a striking contrast (Fig. 4b). In SFF-based batteries, $c\text{VO}_2^+$ exhibits a progressive decline from the inlet to the outlet (Fig. 4b1). This decline corresponds to the gradual consumption of active species driven by electrochemical reactions. In a noteworthy departure, IGFF-based batteries maintain the $c\text{VO}_2^+$ concentration near-outlet region at the same level

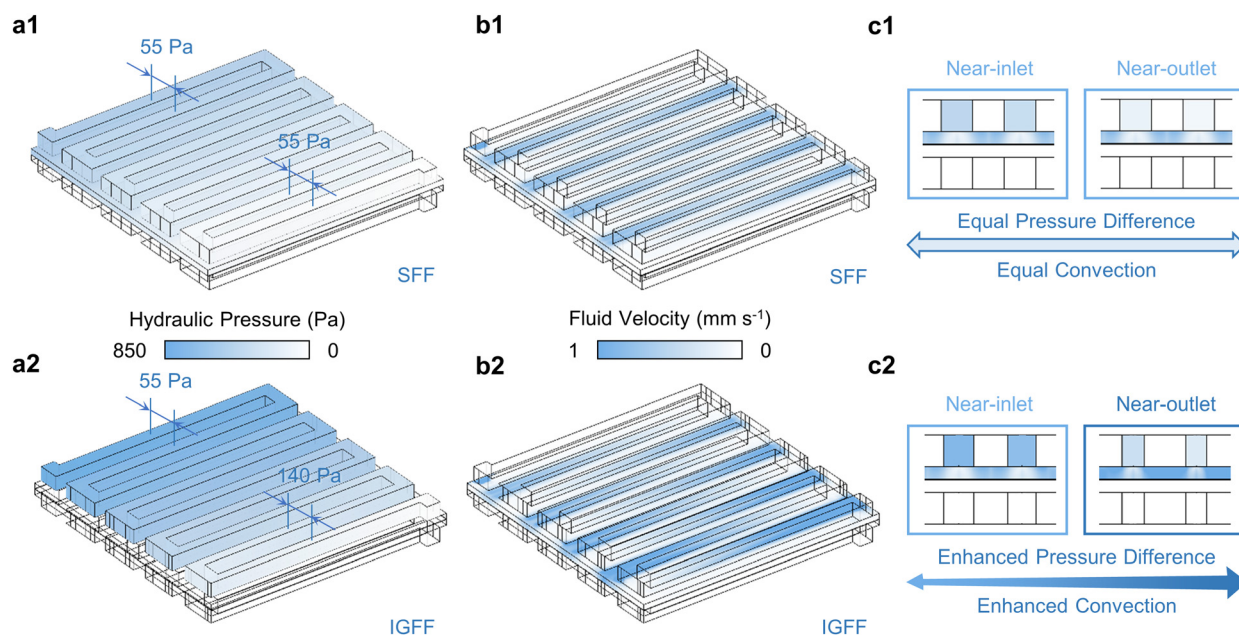


Fig. 3 The (a) hydraulic pressure within flow channels, (b) fluid velocities within porous electrodes and (c) in the cross-view for SFF and IGFF-based batteries.



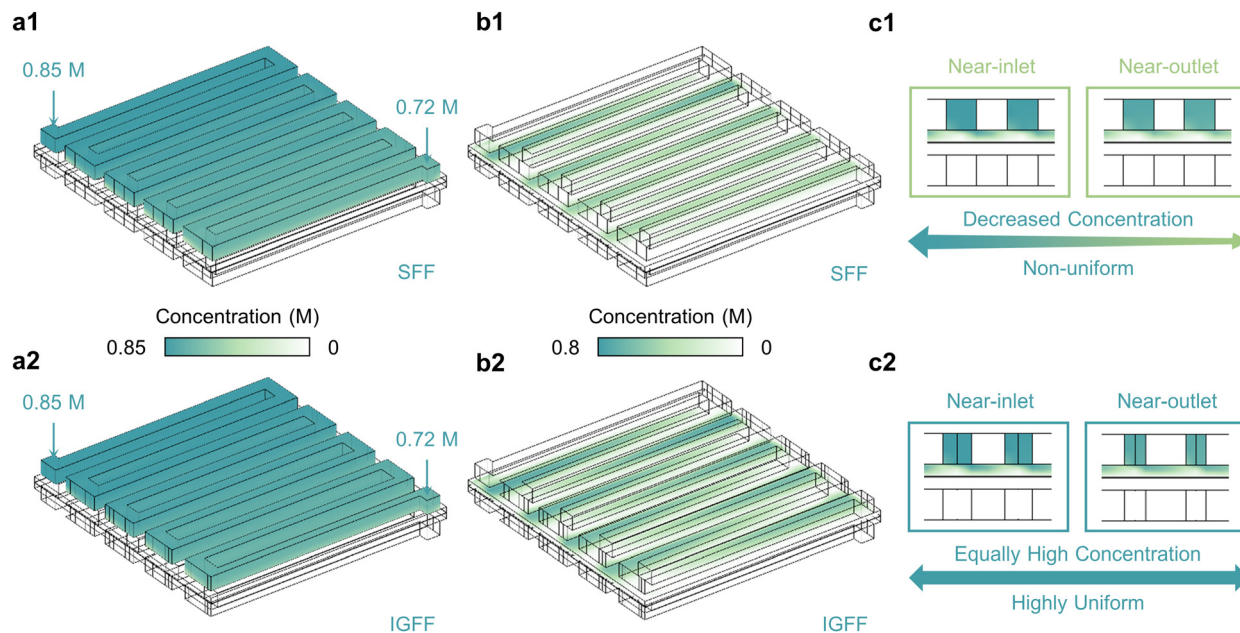


Fig. 4 The reactant concentrations (a) within flow channels, (b) within porous electrodes and (c) in the cross-view for SFF and IGFF-based batteries.

as near-inlet (Fig. 4b2). This intriguing phenomenon, substantiated by cross-sectional perspectives (Fig. 4c), underscores the pivotal role played by the innovative in-plane gradient design, inspired by human behavior, in fostering enhanced under-rib convections. The ramifications of these findings on the electrochemical performance of batteries are profound. The achievement of more uniform and elevated concentrations of active ingredients has the potential to significantly enhance battery performance. Notably, the concentration elevation near the outlet region can be attributed to the augmented under-rib convection patterns stemming from the innovative in-plane gradient design—a concept informed by the principles observed in human life.

The active species' concentration within the porous electrode exhibits intricate interplay with cross-surface distance, current density, and electrolyte flow rate (Fig. 5a–c). During the discharge process, V^{2+} (Fig. 5a1) and VO_2^+ (Fig. 5a2) function as active species for the negative and positive electrodes, respectively. In the context of the through-plane direction, it becomes evident that the average concentration within the porous electrode diminishes as the relative distance from the membrane increases. This behavior can be attributed to the underlying convection patterns beneath the ribs, where fluid velocity tends to be comparatively subdued. In addition, applying higher current densities accelerates the consumption of reactants, resulting in a commensurate reduction in the concentrations of V^{2+} (cV^{2+} , as the reactants are at the negative side for the discharge process), as demonstrated in Fig. 5b. Augmenting the flow rate of the electrolyte offers a direct avenue to bolster the supply of reactants, consequently elevating cV^{2+} levels within the porous electrode (Fig. 5c). A comprehensive assessment necessitates the concurrent consideration of both current density and electrolyte flow rate (Fig. 5d and e). For instance, focusing on the negative electrode during discharge, two key metrics, namely the average

concentration of V^{2+} (cV^{2+}_{average}) and the uniformity factor of cV^{2+} distribution ($UF_{cV^{2+}}$), serve as quantitative benchmarks for discerning between SFF and IGFF. With increasing current density, both cV^{2+}_{average} and $UF_{cV^{2+}}$ exhibit a diminishing trend, but the performance differential between SFF and IGFF steadily amplifies. Notably, the cV^{2+}_{average} consistently registers at lower levels for SFF compared to IGFF; however, this discrepancy dwindles as the electrolyte flow rate is heightened (Fig. 5d). Analogously, $UF_{cV^{2+}}$ follows a parallel trajectory of variation (Fig. 5e). These findings collectively signify that at elevated current densities and reduced flow rates, IGFF exerts a more pronounced positive influence on performance when compared to SFF, underscoring the intricate dynamics of these multi-factorial interactions.

2.3. The performance comparison between SFF and IGFF

The in-plane gradient design, in addition to fostering under-rib convections and uniform distributions, bestows a remarkable enhancement in the electrochemical performance of VRFB. Evident within the polarization curves presented in Fig. 6, these curves faithfully depict the correlation between E_{cell} and the current densities, effectively encapsulating the polarization characteristics intrinsic to the battery. In VRFB, subjecting the system to current densities equal to or surpassing 150 mA cm^{-2} precipitates a discernible upswing in concentration overpotential, thereby inducing a steep decline in E_{cell} . To illustrate, when scrutinizing a scenario wherein the electrolyte flow rate stands at 1 mL min^{-1} and the current density is maintained at 300 mA cm^{-2} (Fig. 6a), the E_{cell} for SFF-based VRFB hovers at approximately 1.00 V. This outcome concurs with the previously expounded disparities in mass transfer associated with SFF. By stark contrast, under analogous conditions, the E_{cell} of IGFF-based VRFB escalates to 1.13 V (Fig. 6b), constituting a noteworthy increment of approximately 13% relative to SFF-based VRFB. This higher discharge



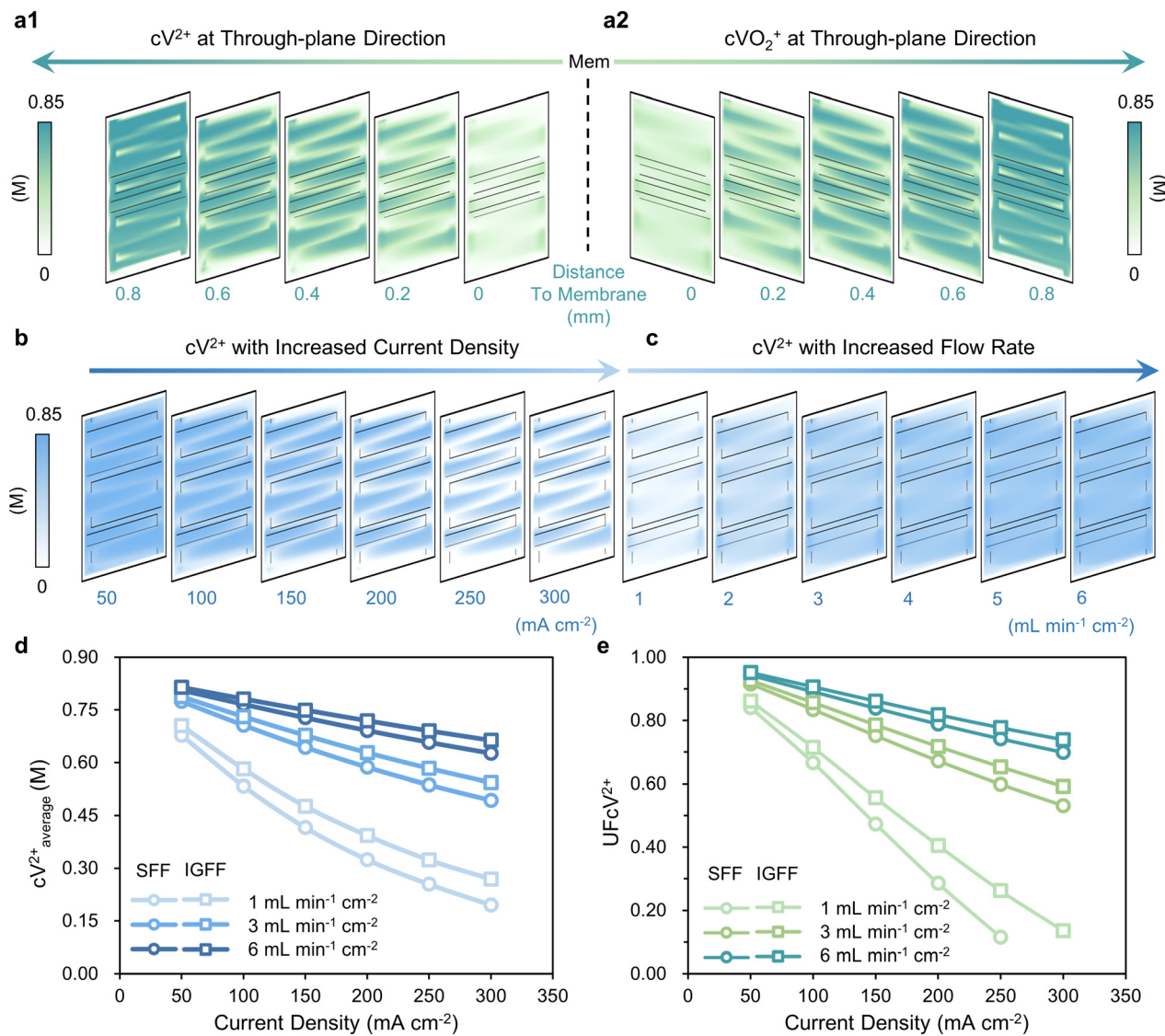


Fig. 5 The distributions of (a1) cV^{2+} for the negative side and (a2) cVO_2^+ for the positive side at different distances to the membrane, with (b) increased current densities and (c) flow rates. The (d) $cV^{2+}_{average}$ and (e) corresponding $UF_{cV^{2+}}$ for SFF and IGFF with different operating conditions.

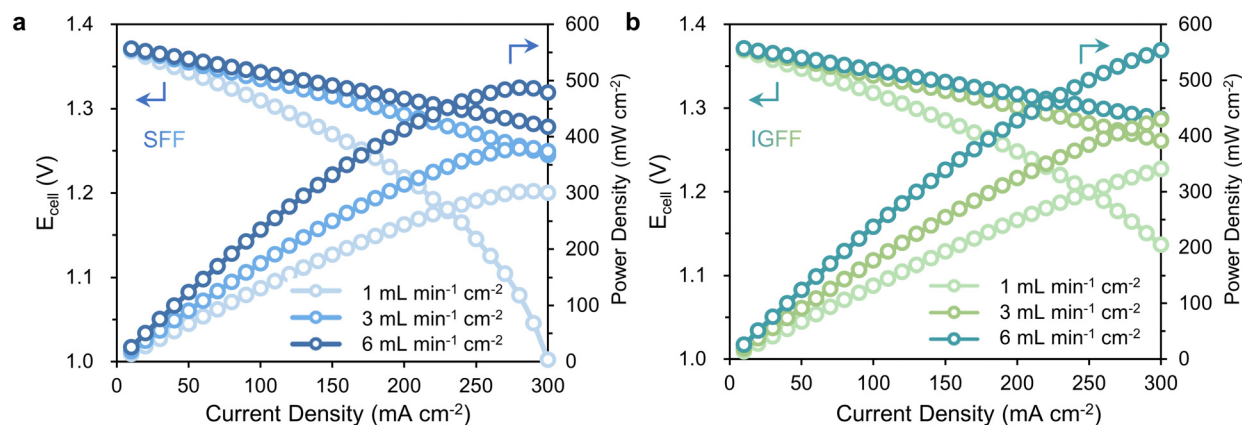


Fig. 6 The polarization curves of (a) SFF-based VRFBs and (b) IGFF-based VRFBs.



voltage coupled with diminished polarization collectively translates into an augmented power density. The maximum power densities (W_{\max}) of IGFF-based batteries are 341.0, 429.8 and 553.2 mW cm^{-2} , respectively, across a spectrum of electrolyte flow rates comprising 1, 3 and 6 $\text{mL min}^{-1} \text{cm}^{-2}$. Notably, these values substantially outstrip their SFF-based counterparts, which record power densities of 300.7, 374.4 and 478.7 mW cm^{-2} , conclusively underscoring the discernible advantages conferred by IGFF in terms of electrochemical performance and power output across various operational conditions.

The observed enhancements in W_{\max} exhibit a discernible correlation with distinct operational conditions. Notably, under relatively low flow rates, IGFF exhibits a significantly more substantial performance improvement compared to SFF (Fig. 7a). This trend also manifests in the relationship between alterations in voltage efficiency (VE) and flow rate (Fig. 7b), an aspect we have previously elucidated in our previous report.²¹ Herein, the concept of the critical flow rate assumes paramount importance. This critical flow rate is rigorously defined as the specific operational condition at which the incremental gain in VE fails to exceed 1% per $\text{mL min}^{-1} \text{cm}^{-2}$. It is imperative to recognize that VE exhibits a noteworthy improvement below the critical flow rate, marking these domains as “sensitive regions”. In stark contrast, higher flow rate regimes are designated as “insensitive regions”, where the amelioration in VE becomes constrained. Astonishingly, the critical flow rate for IGFF-based VRFBs registers at 3.0 $\text{mL min}^{-1} \text{cm}^{-2}$, a value notably lower than the corresponding 3.5 $\text{mL min}^{-1} \text{cm}^{-2}$ observed for SFF. It is essential to highlight that the applied electrolyte flow rate

must exceed this critical threshold to unlock a high VE. Nonetheless, it is imperative to acknowledge that the electrolyte flow rate is not boundless, its upper limits tethered to the pressure drop between the flow field's inlet and outlet (ΔP). This intricate relationship is vividly illustrated in Fig. 7c, wherein ΔP demonstrates an almost linear ascent in tandem with escalating electrolyte flow rates. Significantly, IGFF exhibits a more pronounced amplitude in ΔP compared to SFF. Furthermore, it is imperative to note that in large-size cells exceeding 100 cm^2 ,^{27,35–38} the range of ΔP typically spans from 90–380 Pa cm^{-2} (equating to 1.15–4.92 kPa in this work). This range is primarily delimited by considerations encompassing pipeline design, sealing methodologies, and the intricate art of system integration. Therefore, the prospect of a lower critical flow rate is undeniably gratifying, as it imparts a heightened degree of operational flexibility, ensuring that VRFBs can be judiciously maintained within reasonable and manageable parameters.

The holistic evaluation of pump-based voltage efficiency (VE_{pump}), thoughtfully amalgamating VE and ΔP into a singular metric, furnishes a lucid basis for direct comparisons (Fig. 7d–f). Given the intricate interplay between VE_{pump} and current density, we introduce two pivotal parameters that merit in-depth discussion: the “reversal point” and the “rated current density”. The concept of the “reversal point” serves to delineate a threshold of current density. Conventional designs hold sway below this inflection point, while novel designs predominate above it. Concomitantly, the “rated current density” denotes the maximum permissible current density at which VE_{pump} steadfastly maintains a lofty threshold of 80% or greater. For

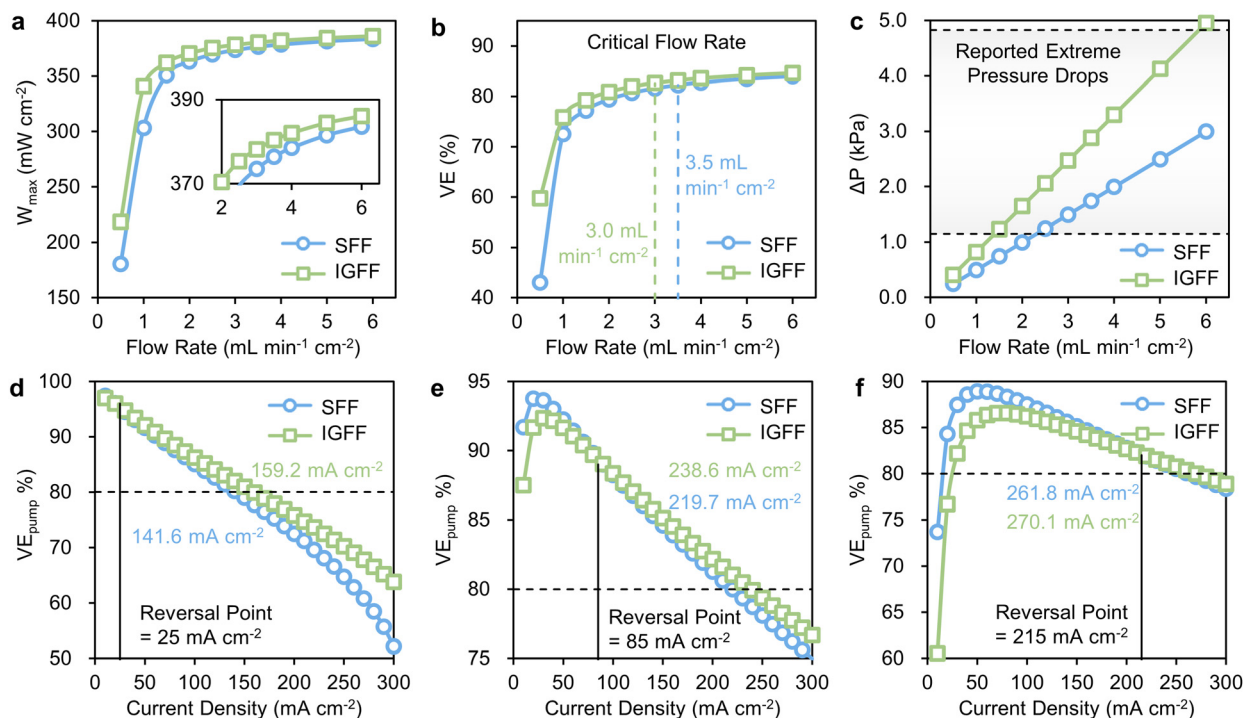


Fig. 7 The relationships of (a) W_{\max} , (b) VE and (c) ΔP with flow rates for SFF and IGFF-based VRFBs. The relationships of VE_{pump} with current densities for SFF and IGFF-based VRFBs under flow rates at (d) 1, (e) 3 and (f) 6 $\text{mL min}^{-1} \text{cm}^{-2}$.



illustrative purposes, it is considered the scenario of a modest flow rate of $1 \text{ mL min}^{-1} \text{ cm}^{-2}$. Here, the “reversal point” stands at 25 mA cm^{-2} , signifying that IGFF-based VRFBs exhibit superior VE_{pump} values relative to their SFF counterparts when subjected to current densities surpassing this threshold (Fig. 7d). Notably, the rated current densities for SFF-based and IGFF-based VRFBs stand at 141.6 mA cm^{-2} and 159.2 mA cm^{-2} , respectively. As the transition to a moderate electrolyte flow rate of $3 \text{ mL min}^{-1} \text{ cm}^{-2}$ (Fig. 7e), both the reversal point and rated current density exhibit upward mobility. Specifically, the VE_{pump} of IGFF-based VRFB eclipses that of SFF-based VRFBs under the current densities exceeding 85 mA cm^{-2} . Most notably, the rated current density for IGFF-based VRFBs soars to an impressive 238.6 mA cm^{-2} , constituting an 8.6% augmentation relative to the SFF configuration's 219.7 mA cm^{-2} . With the ascent to a higher electrolyte flow rate of $6 \text{ mL min}^{-1} \text{ cm}^{-2}$, the expanse between the reversal point (215 mA cm^{-2}) and the rated current density (270.1 mA cm^{-2}) for IGFF diminishes. This conveys that permissible current densities converge within a narrow window of approximately 55 mA cm^{-2} to ensure the sustenance of high VE_{pump} values, as shown in Fig. 7f. While an increased electrolyte flow rate intuitively augments mass transfer, it simultaneously introduces multifaceted challenges encompassing sealing, pressure differentials, and intricate operational considerations. Nonetheless, the rate current density is still 8.3 mA cm^{-2} higher than that of SFF. These findings cogently underscore both the efficacy and indispensability of the innovative in-plane gradient design.

2.4. Basic approaches for flow field optimization evaluation

The in-plane gradient flow field design presented herein stands as a testament to remarkable advancements in enhancing the electrochemical performance of VRFBs. Moreover, this study introduces a novel lexicon of concepts, including “critical flow

rate”, “reversal point”, and “rated current density” which collectively furnish a comprehensive framework for the multi-faceted evaluation of various contributing factors. Notably, the augmentation of under-rib convections, as realized through IGFF, represents a universal panacea applicable not only to the present study but also to previous investigations in flow field optimization.^{38–40} The intricate mechanism underpinning performance enhancement *via* flow field design warrants concise elucidation and synthesis. Consequently, we proffer a generalized evaluation criterion distilled from the tenets of enhanced convection-based optimization, thereby encapsulating the quintessence of flow field refinement (Fig. 8a). This holistic framework encompasses several sequential steps, promulgating a structured approach to propel advancements in flow field optimization:

(1) **Flow rate evaluation.** Elucidates the intricate interplay between VE and flow rate, thereby establishing the lower threshold of the insensitivity region, denoted as the “critical flow rate”. Subsequently, gauge “extreme flow rate”, contingent on the upper limit of pressure drops, a factor crucial for delineating pipeline design, the integrity of sealing mechanisms, and seamless system integration. The imperative lies in minimizing this “critical flow rate” to ensure a sufficiently expansive operational range for VRFBs (Fig. 8b).

(2) **Current density evaluation.** Thoroughly elucidate the intricate relationship between VE_{pump} and current density, leading to the identification of critical parameters such as the “reversal point” and the “rated current density” (defined as the maximum current density where VE_{pump} exceeds 80%) for both conventional and novel designs. Notably, the reversal point should invariably fall below the rated current density. These analyses delineate three distinct regions based on the parameters as mentioned above (Fig. 8c). The first region,

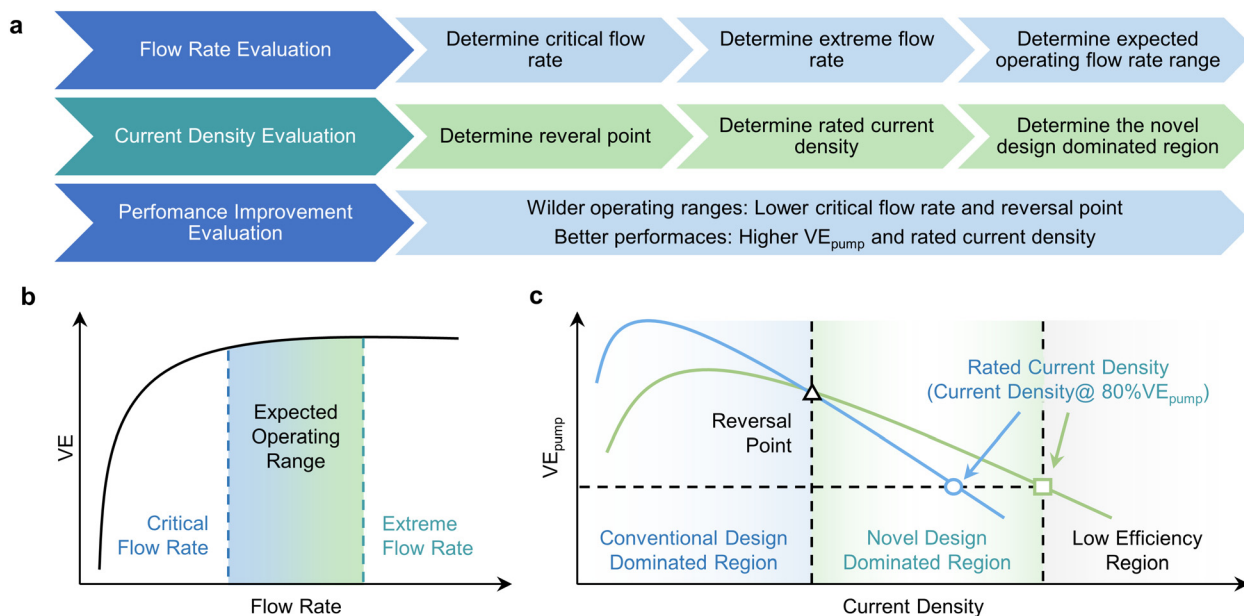


Fig. 8 (a) The basic approaches for evaluation. (b) The overall trend of VE and the determination of operating flow rates. (c) The overall trend of VE_{pump} and the determination of operating current densities.



characterized by current densities below the reversal point, represents the realm of conventional designs where power output remains relatively modest. The intermediate region, spanning between the reversal point and the rated current density, is governed by novel design principles, featuring superior VE_{pump} values. It's crucial to note that higher current density regions within regions of lower efficiency are inherently unsuitable for practical applications.

(3) **Performance improvement evaluation.** The quantitative assessment of electrochemical performance enhancement within the defined parameter range is pivotal. This evaluation hinges on a comprehensive understanding of flow rates and current densities. Notably, pursuing higher VE_{pump} values and elevated rated current densities is both a desirable and anticipated outcome.

3. Conclusions

In this work, a novel design, the so-called in-plane gradient flow field, is proposed for VRFBs. The enhanced under-rib convection in the low-concentration region results in a more uniform distribution of the active species within the porous electrode. Compared with the conventional SFF, the adoption of the newly designed flow field can significantly improve the battery performance. Based on a 3D multi-physical simulated model, introducing an IGFF can improve the average reactant's concentration and distribution uniformity, as well as the VE_{pump} of the battery. The adoption of the proposed IGFF reaches an insensitive area of VE under a flow rate of over $3 \text{ mL min}^{-1} \text{ cm}^{-2}$, and the VE_{pump} surpasses conventional SFF at a current density of over 85 mA cm^{-2} . The W_{max} and maximum rated current density of IGFF-based VRFB are 553.2 mW cm^{-2} and 270.1 mA cm^{-2} , respectively, which are 74.5 mW cm^{-2} and 8.3 mA cm^{-2} higher than SFF-based VRFB. In addition, the basic criteria for evaluating flow field optimization have been clarified. The performance improvement of the flow field by strengthening the convection under the ribs is related to the specific flow rate and current density, and it is necessary to achieve lower critical flow rate and turning points to broaden the application range of the novel design. The obtained results provide compelling evidence that the proposed IGFF is advantageous for the development of high-performance VRFBs. Furthermore, owing to its straightforward, efficient, and easily scalable design inspired by human behavior, the IGFF design holds great potential for application in engineering VRFBs and large-scale energy storage systems.

4. Experimental section

Geometry details

The 3D model consists of two flow fields, two porous electrodes and a membrane with an actual area of $3.6 \times 3.6 \text{ cm}^2$. The geometric structure of the SFF and IGFF is similar. Both are composed of 1 inlet, 1 outlet, 9 long flow channels and 8 short flow channels. Besides, the depth of the channel is 2 mm. The only difference between the SFF and IGFF is that from the

beginning of the first long flow channel near the inlet, the width of each subsequent long flow channel will be decreased by 0.1 mm, the width of the first long flow channel is 2 mm, and the width of the last long channel which is near the outlet is 1.2 mm. The width of every long flow channel of SFF is 2 mm. The two flow fields are analyzed and compared by numerical simulation, described in detail in the methods section.

Physical assumptions

The physical assumptions for the model are as follows:

- (1) The electrolyte is regarded as an incompressible fluid, and its flow is considered as laminar flow.
- (2) The electrodes are isotropic and uniform.
- (3) The whole model is regarded as an isothermal model.
- (4) Ignore all side reactions, including hydrogen evolution reactions.
- (5) The influence of gravity is not taken into account.
- (6) The electrolyte shall be regarded as a dilute solution.

Governing equations

The mass conservation of both the flow channels and porous media is expressed as:

$$\rho \nabla \cdot \vec{u} = 0 \quad (1)$$

where ρ is the density of the electrolyte and \vec{u} is the velocity of the electrolyte.

The momentum conservation of electrolytes in flow channels and porous media can be expressed by the Navier–Stokes equation and Brinkman equation, respectively:⁴¹

$$\rho(\vec{u} \cdot \nabla) \vec{u} = -\nabla p + \nabla \cdot [\mu \nabla \vec{u} + (\nabla \vec{u})^T] + \vec{f} \quad (2)$$

$$\frac{\rho}{\varepsilon} (\vec{u} \cdot \nabla) \frac{\vec{u}}{\varepsilon} = -\nabla p + \nabla \cdot \left[\frac{\mu}{\varepsilon} \nabla \vec{u} + (\nabla \vec{u})^T \right] - \left(\frac{\mu}{k} + \beta_F |\vec{u}| \right) \vec{u} + \vec{f} \quad (3)$$

where ε is the porosity of the porous media, p is the pressure, μ is the dynamic viscosity of the electrolyte, and k is the permeability of the porous media. β_F is the Forchheimer drag coefficient, which is ignored in the simulation. Since gravity is also not taken into account in the model, the volume force acting on the fluid is zero.

The permeability of the porous media is calculated by the Carman–Kozeny equation:⁴²

$$k = \frac{d_f^2 \varepsilon^3}{16 k_{\text{ck}} (1 - \varepsilon)^2} \quad (4)$$

where d_f is the diameter of the fiber and k_{ck} is the Carman–Kozeny constant, which is a dimensionless number that depends on the shape and orientation of the fiber material.

The Nernst–Planck equation describes the transfer process of dilute species in porous media. The equation includes diffusion item, migration item and convection item:⁴³

$$\vec{N}_i = -D_i^{\text{eff}} \nabla c_i - \frac{z_i c_i D_i^{\text{eff}} F}{RT} \nabla \varphi_i + \vec{u} c_i \quad (5)$$



The conservation of each species can be expressed as:⁴³

$$\nabla \cdot \vec{N}_i = S_i \quad (6)$$

where the subscript i represents a certain species, and \vec{N}_i represents the flux of the species, S_i is the source term of the species, and is used to represent the generation rate due to the electrochemical reactions. c_i and z_i represent the concentration and charge of a species i , respectively. R is the ideal gas constant, T is the temperature, and F is the Faraday's constant. φ_i is the potential of a species in the liquid phase.

The effective diffusion coefficient D_i^{eff} can be obtained by Bruggeman's modification:⁴⁴

$$D_i^{\text{eff}} = \varepsilon^{\frac{3}{2}} D_i \quad (7)$$

The charge conservation on the positive and negative sides is solved as the following two equations, respectively:

$$\nabla \cdot \vec{i}_s = -\nabla \cdot \vec{i}_l = i_{\text{pos}} \quad (8)$$

$$\nabla \cdot \vec{i}_s = -\nabla \cdot \vec{i}_l = i_{\text{neg}} \quad (9)$$

where \vec{i}_s and \vec{i}_l are the current densities of the solid phase and the liquid phase, respectively, and they can be calculated as:

$$\vec{i}_s = -\sigma_s^{\text{eff}} \nabla \varphi_s \quad (10)$$

$$\vec{i}_l = F \sum_i z_i \vec{N}_i \quad (11)$$

where σ_s^{eff} is the effective conductivity of the porous electrodes, which can be calculated as:

$$\sigma_s^{\text{eff}} = (1 - \varepsilon)^{\frac{3}{2}} \sigma_s \quad (12)$$

where σ_s is the electronic conductivity of the porous electrodes. The source term S_i is related to the generation rate of the electrochemical reaction, and each source term can be calculated as:

$$S_{V^{2+}} = \frac{i_{\text{neg}}}{F} \quad (13)$$

$$S_{V^{3+}} = -\frac{i_{\text{neg}}}{F} \quad (14)$$

$$S_{V^{4+}} = \frac{i_{\text{pos}}}{F} \quad (15)$$

$$S_{V^{5+}} = -\frac{i_{\text{pos}}}{F} \quad (16)$$

The results can also be obtained using the Butler-Volmer equations:⁴³

$$i_{\text{pos}} = aFk_{0,\text{pos}} c_{V^{4+}}^{z_{p,c}} c_{V^{5+}}^{z_{p,a}} \left[\frac{c_{V^{5+}}^s}{c_{V^{5+}}} \exp\left(-\frac{\alpha_c F \eta_{\text{pos}}}{RT}\right) - \frac{c_{V^{4+}}^s}{c_{V^{4+}}} \exp\left(-\frac{\alpha_a F \eta_{\text{pos}}}{RT}\right) \right] \quad (17)$$

$$i_{\text{neg}} = aFk_{0,\text{neg}} c_{V^{2+}}^{z_{n,c}} c_{V^{3+}}^{z_{n,a}} \left[\frac{c_{V^{3+}}^s}{c_{V^{3+}}} \exp\left(-\frac{\alpha_c F \eta_{\text{neg}}}{RT}\right) - \frac{c_{V^{2+}}^s}{c_{V^{2+}}} \exp\left(-\frac{\alpha_a F \eta_{\text{neg}}}{RT}\right) \right] \quad (18)$$

where $k_{0,\text{pos}}$ and $k_{0,\text{neg}}$ are the reaction rate coefficients on the positive and the negative sides, respectively. α_a and α_c is the charge transfer coefficient on the anode and cathode sides, respectively.

The concentrations of vanadium ions at the liquid-solid interfaces can be calculated *via* bulk concentration and the corresponding parameters. On the positive side, the surface concentrations of V^{4+} and V^{5+} are derived as:⁴⁵

$$c_{V^{4+}}^s = \frac{B_2 c_{V^{5+}} + (1 + B_2) c_{V^{4+}}}{1 + A_2 + B_2} \quad (19)$$

$$c_{V^{5+}}^s = \frac{A_2 c_{V^{4+}} + (1 + A_2) c_{V^{5+}}}{1 + A_2 + B_2} \quad (20)$$

where

$$A_2 = \frac{k_{0,\text{pos}}}{k_m} c_{V^{4+}}^{z_c-1} c_{V^{5+}}^{z_a} \exp(\alpha_a F \eta_{\text{pos}}) \quad (21)$$

$$B_2 = \frac{k_{0,\text{pos}}}{k_m} c_{V^{4+}}^{z_c} c_{V^{5+}}^{z_a-1} \exp(-\alpha_c F \eta_{\text{pos}}) \quad (22)$$

For the negative side, the surface concentrations of V^{2+} and V^{3+} are derived as:

$$c_{V^{2+}}^s = \frac{B_1 c_{V^{3+}} + (1 + B_1) c_{V^{2+}}}{1 + A_1 + B_1} \quad (23)$$

$$c_{V^{3+}}^s = \frac{A_1 c_{V^{2+}} + (1 + A_1) c_{V^{3+}}}{1 + A_1 + B_1} \quad (24)$$

where

$$A_1 = \frac{k_{0,\text{neg}}}{k_m} c_{V^{2+}}^{z_c-1} c_{V^{3+}}^{z_a} \exp(\alpha_a F \eta_{\text{neg}}) \quad (25)$$

$$B_1 = \frac{k_{0,\text{neg}}}{k_m} c_{V^{2+}}^{z_c} c_{V^{3+}}^{z_a-1} \exp(-\alpha_c F \eta_{\text{neg}}) \quad (26)$$

The mass transfer coefficient in the above equation can be roughly calculated as:⁴⁶

$$k_m = 1.6 \times 10^{-4} \bar{u}^{0.4} \quad (27)$$

The overpotential on the positive and negative sides can be defined as the following two equations, respectively:⁴⁷

$$\eta_{\text{pos}} = \varphi_{\text{pos,s}} - \varphi_{\text{pos,l}} - E_{\text{eq,pos}} \quad (28)$$

$$\eta_{\text{neg}} = \varphi_{\text{neg,s}} - \varphi_{\text{neg,l}} - E_{\text{eq,neg}} \quad (29)$$

The equilibrium potential on the positive and negative sides can be calculated by the Nernst equation, respectively:⁴⁵

$$E_{\text{eq,pos}} = E_{\text{pos}}^0 + \frac{RT}{F} \ln\left(\frac{c_{V^{5+}} c_{H^+}}{c_{V^{4+}}}\right) \quad (30)$$



$$E_{\text{eq,neg}} = E_{\text{neg}}^0 + \frac{RT}{F} \ln \left(\frac{c_{\text{V}^{3+}}}{c_{\text{V}^{2+}}} \right) \quad (31)$$

where E_{pos}^0 and E_{neg}^0 are the equilibrium potentials under the standard conditions, then the expressions of the battery's discharging voltage and charging voltage can be calculated as the following two equations, respectively:

$$E_{\text{cell,discharge}} = E_{\text{eq,pos}} - E_{\text{eq,neg}} - \eta_{\text{neg}} + \eta_{\text{pos}} - IR_{\text{cell}} \quad (32)$$

$$E_{\text{cell,charge}} = E_{\text{eq,pos}} - E_{\text{eq,neg}} - \eta_{\text{pos}} + \eta_{\text{neg}} + IR_{\text{cell}} \quad (33)$$

where I is the current density and R_{cell} is the cell electrical resistance.

Post processes

Average reactant concentrations, uniformity factor, pressure drop, VE and VE_{pump} are key indexes commonly used to evaluate the quality of a flow field design.

The VE of the battery can be calculated as follows:^{48,49}

$$\text{VE} = \frac{E_{\text{cell,discharge}}}{E_{\text{cell,charge}}} \quad (34)$$

where $E_{\text{cell,charge}}$ and $E_{\text{cell,discharge}}$ are the cell voltages during the charge and discharge processes, respectively.

In addition, the pressure drop refers to the difference between the inlet and outlet pressure. The pumping loss per active area is usually expressed by:

$$W_{\text{pump}} = \frac{2q\Delta P}{\eta} \quad (35)$$

where η represents the pump's efficiency, which equals 0.6, and q represents the flow rate per active area.

Considering the influence of pumping loss, the VE_{pump} of the battery can be calculated as the following:^{48,49}

$$\text{VE}_{\text{pump}} = \frac{IE_{\text{cell,discharge}} - W_{\text{pump}}}{IE_{\text{cell,charge}} + W_{\text{pump}}} \quad (36)$$

The cell voltages are calculated under an SOC of 50% and initial vanadium concentration of 1.7 M.

The average reactant concentration is calculated at the negative electrodes with the concentration of V^{2+} ions, which can be defined as:

$$c_{\text{V}^{2+}}^{\text{average}} = \frac{1}{V} \iiint c_{\text{V}^{2+}} dV \quad (37)$$

The uniformity factor is used to evaluate the distribution of current density and reactant concentration under a flow field design. For example, the uniformity factor of the V^{2+} ion concentration is defined as:^{23,50}

$$\text{UF}_{\text{cV}^{2+}} = 1 - \frac{1}{c_{\text{V}^{2+}}^{\text{average}}} \sqrt{\frac{1}{V} \iiint (c_{\text{V}^{2+}} - c_{\text{V}^{2+}}^{\text{average}})^2 dV} \quad (38)$$

where V is the volume of the electrode, $c_{\text{V}^{2+}}$ is the concentration

of the V^{2+} ions, and $c_{\text{V}^{2+}}^{\text{average}}$ is the average concentration of the V^{2+} ions.

Author contributions

Lyuming Pan: conceptualization, methodology, simulations, writing – original draft. Jianyu Xie: methodology, simulations, data processes. Jincang Guo: conceptualization, methodology, writing – original draft. Dongbo Wei: simulations, data processes. Honghao Qi: methodology, writing – original draft. Haoyao Rao: methodology, data processes. Puiki Leung: writing – original draft, writing – review & editing. Lin Zeng: writing – original draft, writing – review & editing. Tianshou Zhao: resources, project administration, funding acquisition, supervision. Lei Wei: conceptualization, funding acquisition, project administration, writing – review & editing.

Conflicts of interest

The authors declare no conflict of interest.

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References

- 1 L. Zhang and G. Yu, *Angew. Chem., Int. Ed.*, 2021, **60**, 15028–15035.
- 2 J. Che, F. Wang, C. Song, R. Wang and Y. Li, *Chem. Eng. J.*, 2023, **463**, 142435.
- 3 E. Sánchez-Díez, E. Ventosa, M. Guarnieri, A. Trovò, C. Flox, R. Marcilla, F. Soavi, P. Mazur, E. Aranzabe and R. Ferret, *J. Power Sources*, 2021, **481**, 228804.
- 4 P. Leung, X. Li, C. Ponce de León, L. Berlouis, C. T. J. Low and F. C. Walsh, *RSC Adv.*, 2012, **2**, 10125.
- 5 G. L. Soloveichik, *Chem. Rev.*, 2015, **115**, 11533–11558.
- 6 X. Ke, J. M. Prah, J. I. D. Alexander, J. S. Wainright, T. A. Zawodzinski and R. F. Savinell, *Chem. Soc. Rev.*, 2018, **47**, 8721–8743.
- 7 K. Zhang, J. Xiong, C. Yan and A. Tang, *Appl. Energy*, 2020, **272**, 115093.
- 8 K. Zhang, C. Yan and A. Tang, *J. Power Sources*, 2020, **479**, 228816.
- 9 K. M. Lisboa, J. Marschewski, N. Ebejer, P. Ruch, R. M. Cotta, B. Michel and D. Poulikakos, *J. Power Sources*, 2017, **359**, 322–331.



- 10 D. S. Aaron, Q. Liu, Z. Tang, G. M. Grim, A. B. Papandrew, A. Turhan, T. A. Zawodzinski and M. M. Mench, *J. Power Sources*, 2012, **206**, 450–453.
- 11 Q. H. Liu, G. M. Grim, A. B. Papandrew, A. Turhan, T. A. Zawodzinski and M. M. Mench, *J. Electrochem. Soc.*, 2012, **159**, A1246–A1252.
- 12 J. Sun, M. Zheng, Z. Yang and Z. Yu, *Energy*, 2019, **173**, 637–646.
- 13 R. M. Darling and M. L. Perry, *J. Electrochem. Soc.*, 2014, **161**, A1381–A1387.
- 14 Z. Huang, A. Mu, L. Wu, H. Wang and Y. Zhang, *Int. J. Hydrogen Energy*, 2021, **46**, 31952–31962.
- 15 R. Wang, M. Hao, C. He, Z. Tu, F. Chong and Y. Li, *Appl. Catal., B*, 2023, **332**, 122773.
- 16 R. Wang, C. He, M. Hao, F. Chong, Z. Zhao, X. Wu, Z. Tu, T. Fan and Y. Li, *ACS Sustainable Chem. Eng.*, 2023, **11**, 5953–5962.
- 17 X. Zhou, X. Zhang, L. Mo, X. Zhou and Q. Wu, *Small*, 2020, **16**, e1907333.
- 18 Y. Lv, Z. Pan, X. Zhou, N. Xiong, F. Wang, Y. Li and Q. Wu, *Chem. – Asian J.*, 2022, **17**, e202200825.
- 19 R. Wang, Y. Li and Y.-L. He, *J. Mater. Chem. A*, 2019, **7**, 10962–10970.
- 20 C. Xu and T. S. Zhao, *Electrochem. Commun.*, 2007, **9**, 497–503.
- 21 L. Wei, Z. X. Guo, J. Sun, X. Z. Fan, M. C. Wu, J. B. Xu and T. S. Zhao, *Int. J. Heat Mass Transfer*, 2021, **179**, 121747.
- 22 Z. Guo, J. Sun, X. Fan and T. Zhao, *J. Power Sources*, 2023, **583**, 233540.
- 23 S. B. Wan, H. R. Jiang, Z. X. Guo, C. X. He, X. W. Liang, N. Djilali and T. Zhao, *Energy Environ. Sci.*, 2022, **15**, 2874–2888.
- 24 J. Lee, J. Kim and H. Park, *Int. J. Hydrogen Energy*, 2019, **44**, 29483–29492.
- 25 J. Marschewski, L. Brenner, N. Ebejer, P. Ruch, B. Michel and D. Poulidakos, *Energy Environ. Sci.*, 2017, **10**, 780–787.
- 26 X. Ke, J. I. D. Alexander, J. M. Prahl and R. F. Savinell, *J. Power Sources*, 2014, **270**, 646–657.
- 27 E. Knudsen, P. Albertus, K. T. Cho, A. Z. Weber and A. Kojic, *J. Power Sources*, 2015, **299**, 617–628.
- 28 S. Kumar and S. Jayanti, *J. Power Sources*, 2017, **360**, 548–558.
- 29 M.-Y. Lu, Y.-H. Jiao, X.-Y. Tang, W.-W. Yang, M. Ye and Q. Xu, *J. Energy Storage*, 2021, **35**, 102284.
- 30 J. Sun, M. Zheng, Y. Luo and Z. Yu, *J. Power Sources*, 2019, **428**, 136–145.
- 31 Q. Xu and T. S. Zhao, *Prog. Energy Combust. Sci.*, 2015, **49**, 40–58.
- 32 J. Sun, H. R. Jiang, B. W. Zhang, C. Y. H. Chao and T. S. Zhao, *Appl. Energy*, 2020, **259**, 114198.
- 33 X. Wu, R. Wang and Y. Li, *Electrochim. Acta*, 2022, **433**, 141267.
- 34 O. C. Esan, X. Shi, Z. Pan, X. Huo, L. An and T. S. Zhao, *Adv. Energy Mater.*, 2020, **10**, 2000758.
- 35 Y. Zeng, F. Li, F. Lu, X. Zhou, Y. Yuan, X. Cao and B. Xiang, *Appl. Energy*, 2019, **238**, 435–441.
- 36 D. Reed, E. Thomsen, B. Li, W. Wang, Z. Nie, B. Koeppel and V. Sprenkle, *J. Power Sources*, 2016, **306**, 24–31.
- 37 S. Kim, E. Thomsen, G. Xia, Z. Nie, J. Bao, K. Recknagle, W. Wang, V. Viswanathan, Q. Luo, X. Wei, A. Crawford, G. Coffey, G. Maupin and V. Sprenkle, *J. Power Sources*, 2013, **237**, 300–309.
- 38 Z. Guo, J. Ren, J. Sun, B. Liu, X. Fan and T. Zhao, *J. Power Sources*, 2023, **564**, 232757.
- 39 L. Pan, J. Sun, H. Qi, M. Han, Q. Dai, J. Xu, S. Yao, Q. Li, L. Wei and T. Zhao, *Proc. Natl. Acad. Sci. U. S. A.*, 2023, **120**, e2305572120.
- 40 L. Pan, J. Sun, H. Qi, M. Han, L. Chen, J. Xu, L. Wei and T. Zhao, *J. Power Sources*, 2023, **570**, 233012.
- 41 J. Sun, H. R. Jiang, B. W. Zhang, C. Y. H. Chao and T. S. Zhao, *Appl. Energy*, 2020, **259**, 11.
- 42 S. Zhu, R. H. Pelton and K. Collver, *Chem. Eng. Sci.*, 1995, **50**, 3557–3572.
- 43 J. Newman and N. P. Balsara, *Electrochemical systems*, John Wiley & Sons, 2021.
- 44 A. A. Shah, M. J. Watt-Smith and F. C. Walsh, *Electrochim. Acta*, 2008, **53**, 8087–8100.
- 45 A. Shah, R. Tangirala, R. Singh, R. G. A. Wills and F. C. Walsh, *J. Electrochem. Soc.*, 2011, **158**, A671–A677.
- 46 D. Schmal, J. Vanerkel and P. J. Vanduin, *J. Appl. Electrochem.*, 1986, **16**, 422–430.
- 47 D. J. You, H. M. Zhang and J. Chen, *Electrochim. Acta*, 2009, **54**, 6827–6836.
- 48 Y. K. Zeng, F. H. Li, F. Lu, X. L. Zhou, Y. P. Yuan, X. L. Cao and B. Xiang, *Appl. Energy*, 2019, **238**, 435–441.
- 49 Y. K. Zeng, T. S. Zhao, X. L. Zhou, L. Zeng and L. Wei, *Appl. Energy*, 2016, **182**, 204–209.
- 50 Q. Ye, T. S. Zhao and C. Xu, *Electrochim. Acta*, 2006, **51**, 5420–5429.

