



US008771856B2

(12) **United States Patent**  
**Li et al.**

(10) **Patent No.:** **US 8,771,856 B2**  
(45) **Date of Patent:** **Jul. 8, 2014**

(54) **FE-V REDOX FLOW BATTERIES**

(75) Inventors: **Liyu Li**, Richland, WA (US); **Soowhan Kim**, Richland, WA (US); **Zhenguo Yang**, Richland, WA (US); **Wei Wang**, Richland, WA (US); **Jianlu Zhang**, Richland, WA (US); **Baowei Chen**, Richland, WA (US); **Zimin Nie**, Richland, WA (US); **Guanguang Xia**, Pasco, WA (US)

4,814,241 A	3/1989	Nagashima et al.
6,562,514 B1 *	5/2003	Kazacos et al. .... 429/204
7,320,844 B2	1/2008	Skyllas-Kazacos
2003/0143456 A1	7/2003	Kazacos et al.
2005/0244707 A1	11/2005	Skyllas-Kazacos et al.
2006/0183016 A1 *	8/2006	Kazacos et al. .... 429/105
2008/0292964 A1	11/2008	Kazacos et al.
2010/0230632 A1 *	9/2010	Adamson et al. .... 252/182.1
2011/0045332 A1 *	2/2011	Horne et al. .... 429/105
2011/0086247 A1 *	4/2011	Keshavarz et al. .... 429/9
2012/0040216 A1 *	2/2012	Parakulam .... 429/61

(73) Assignee: **Battelle Memorial Institute**, Richland, WA (US)

(\* ) Notice: Subject to any disclaimer, the term of this patent is extended or adjusted under 35 U.S.C. 154(b) by 785 days.

**FOREIGN PATENT DOCUMENTS**

CN	1770503	*	5/2006
JP	04019966 S		5/1990
WO	9512219 A1		5/1995
WO	9635239 A1		11/1996

(21) Appl. No.: **12/892,698**

(22) Filed: **Sep. 28, 2010**

(65) **Prior Publication Data**

US 2012/0077067 A1 Mar. 29, 2012

(51) **Int. Cl.**

**H01M 8/18** (2006.01)

**H01M 8/20** (2006.01)

(52) **U.S. Cl.**

CPC ..... **H01M 8/184** (2013.01); **H01M 8/188** (2013.01); **H01M 8/20** (2013.01)

USPC ..... **429/105**; 429/204

(58) **Field of Classification Search**

CPC ..... H01M 8/20; H01M 8/188; H01M 8/184

USPC ..... 429/105, 204

See application file for complete search history.

(56) **References Cited**

**U.S. PATENT DOCUMENTS**

3,996,064 A	12/1976	Thaller
4,786,567 A	11/1988	Skyllas-Kazacos et al.

**STN Search.\***

Skyllas-Kazacos, Maria, "Novel vanadium chloride/polyhalide redox flow battery", Journal of Power Sources, 2003, 299-302 pps., vol. 124, Elsevier, Sydney, Australia.

\* cited by examiner

*Primary Examiner* — Milton I Cano

*Assistant Examiner* — Stephen Yanchuk

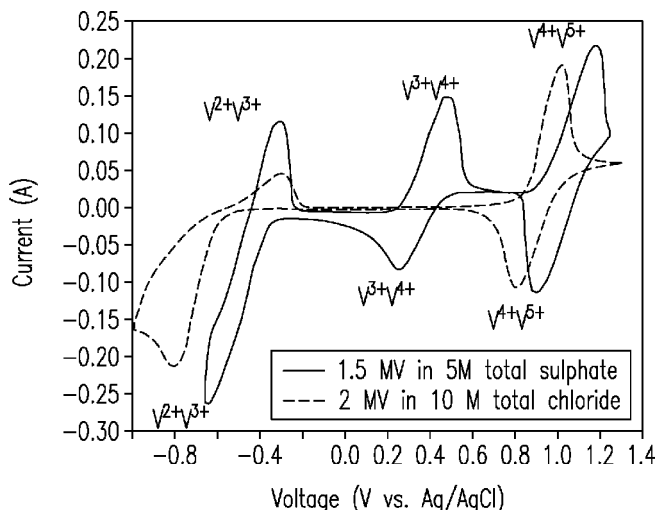
(74) *Attorney, Agent, or Firm* — Klarquist Sparkman, LLP

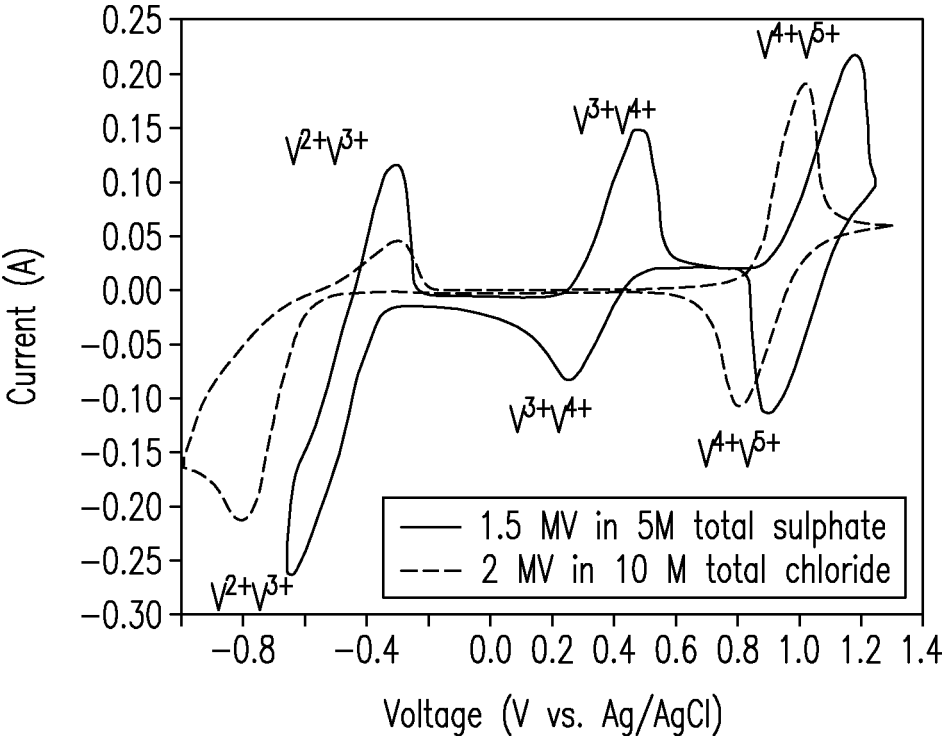
**OTHER PUBLICATIONS**

(57) **ABSTRACT**

A redox flow battery having a supporting solution that includes Cl<sup>-</sup> anions is characterized by an anolyte having V<sup>2+</sup> and V<sup>3+</sup> in the supporting solution, a catholyte having Fe<sup>2+</sup> and Fe<sup>3+</sup> in the supporting solution, and a membrane separating the anolyte and the catholyte. The anolyte and catholyte can have V cations and Fe cations, respectively, or the anolyte and catholyte can each contain both V and Fe cations in a mixture. Furthermore, the supporting solution can contain a mixture of SO<sub>4</sub><sup>2-</sup> and Cl<sup>-</sup> anions.

**18 Claims, 11 Drawing Sheets**





*Fig. 1*

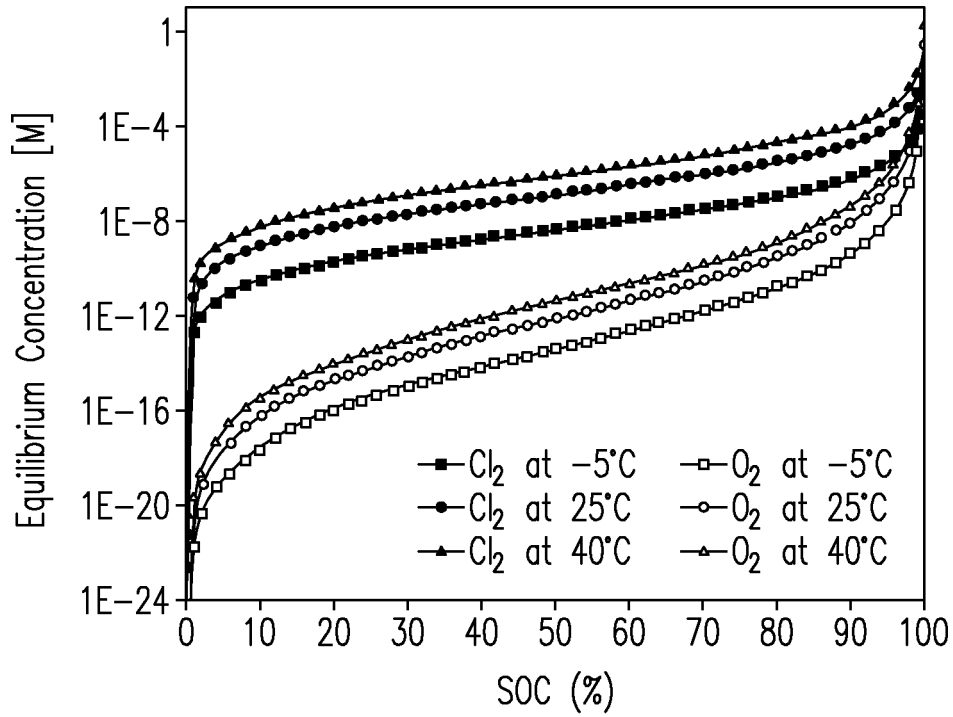


Fig. 2a

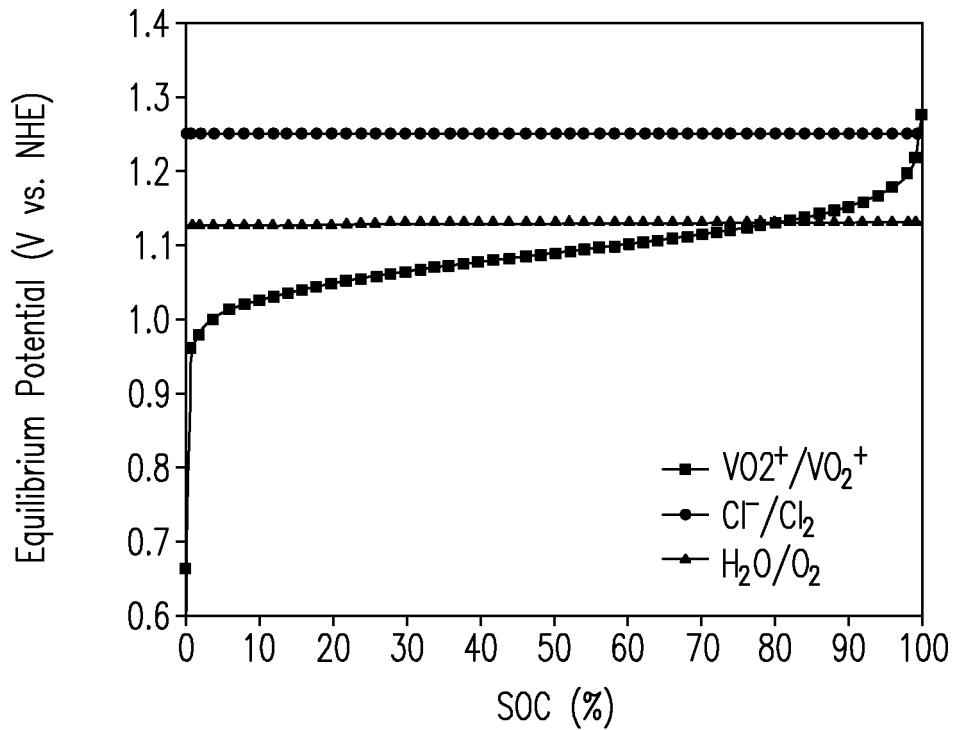
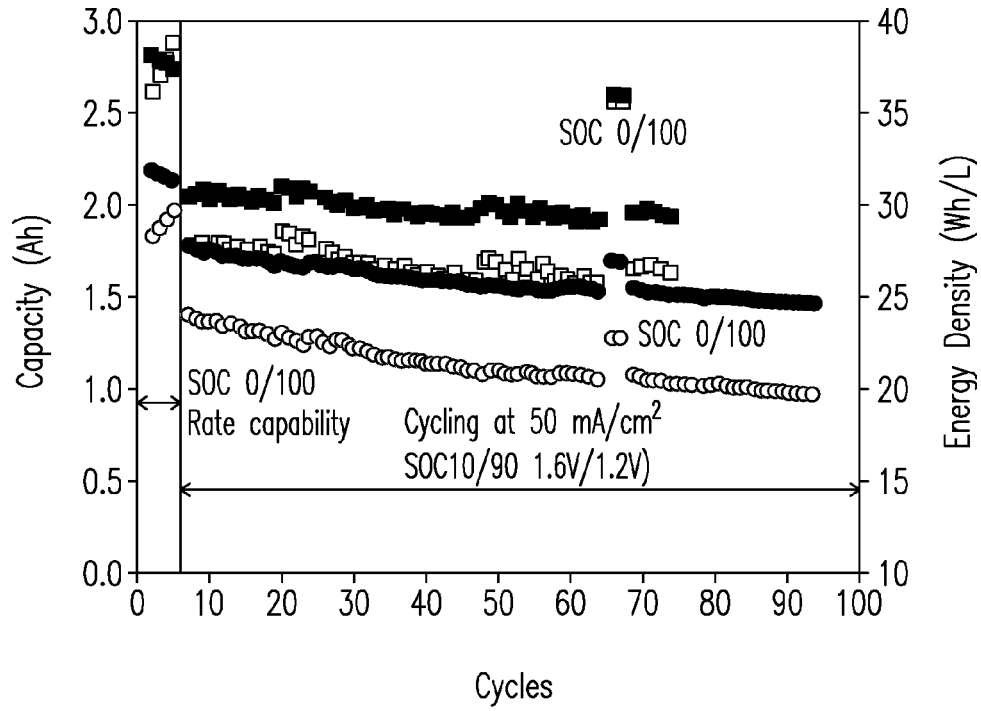
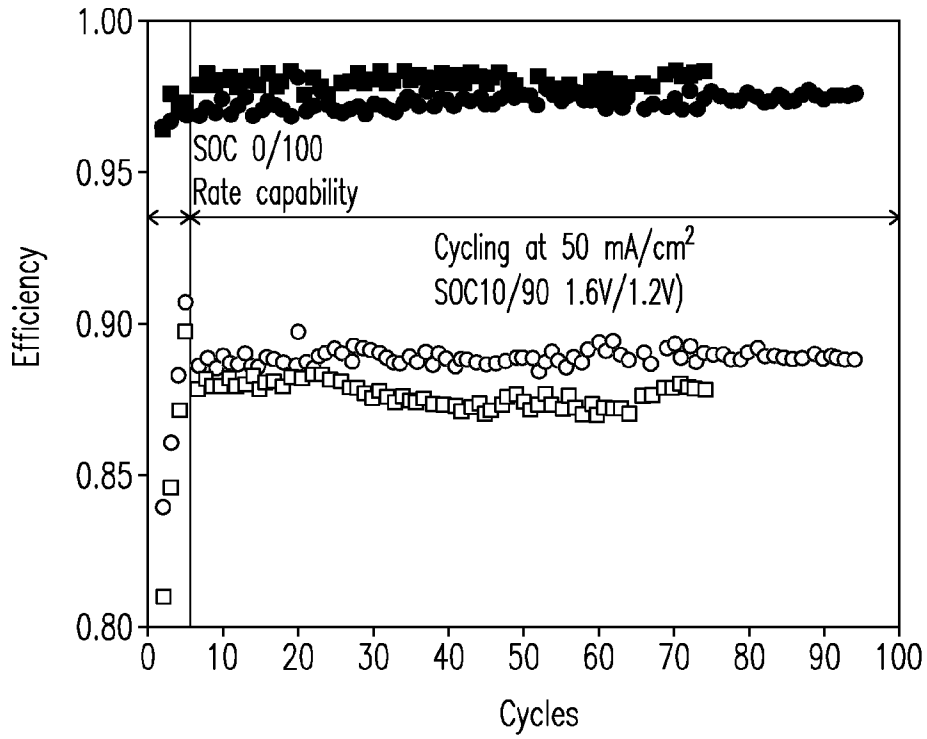


Fig. 2b



- Capacity: 2.3 MV in 10 M total chloride
- Capacity: 1.7 MV in 5 M total sulphate
- Energy Density: 2.3 MV in 10 M total chloride
- Energy Density: 1.7 MV in 5 M total sulphate

*Fig. 3a*



- Capacity: 2.3 MV in 10 M total chloride
- Capacity: 1.7 MV ub 5 M total sulphate
- Energy Density: 2.3 MV in 10 M total chloride
- Energy Density: 1.7 MV in 5 M total sulphate

Fig. 3b

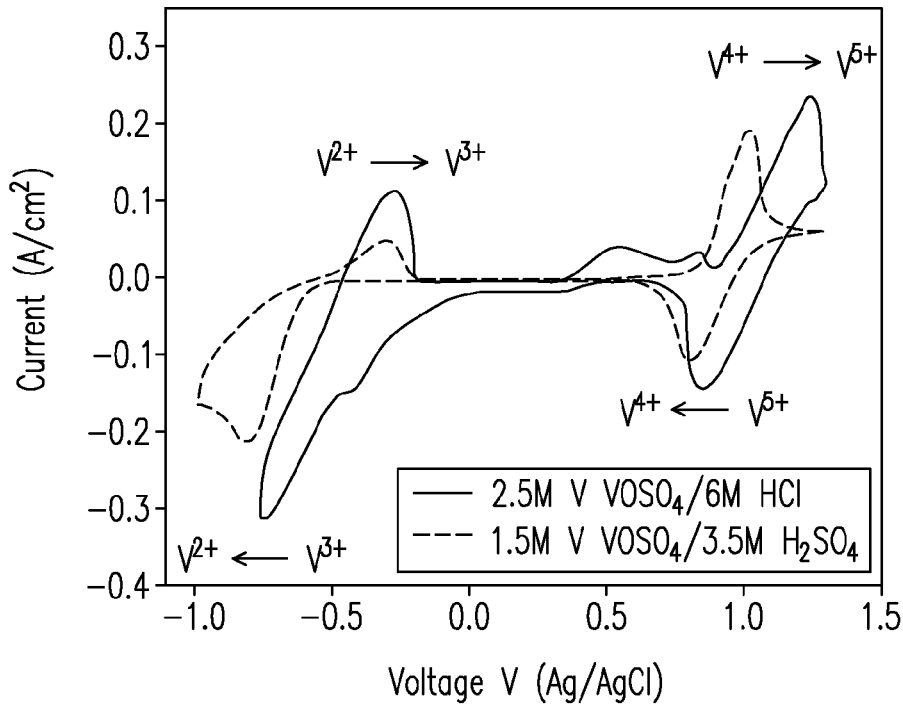


Fig. 4

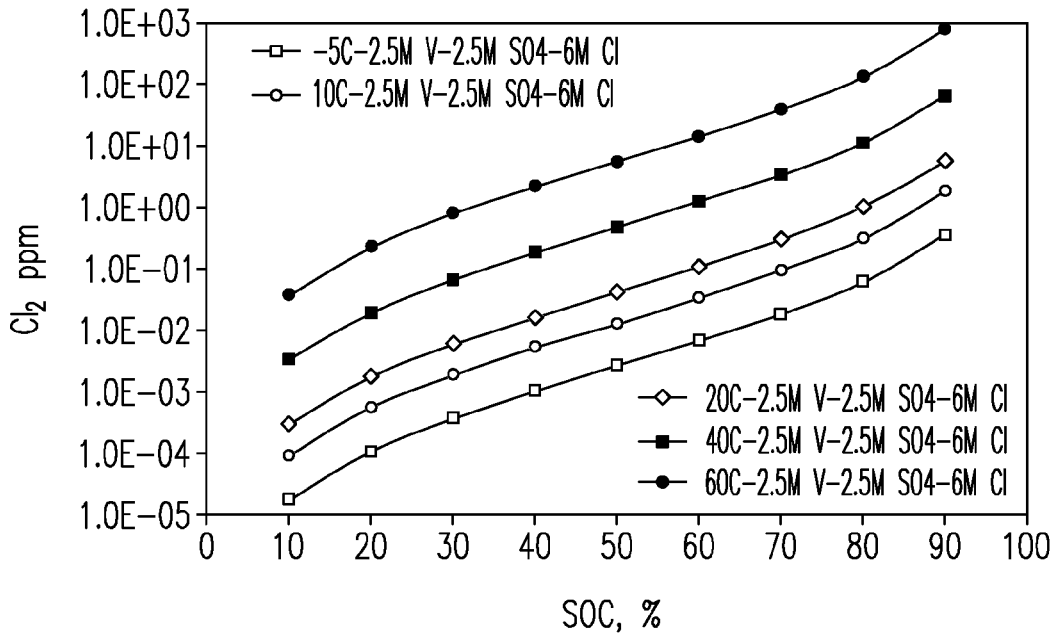
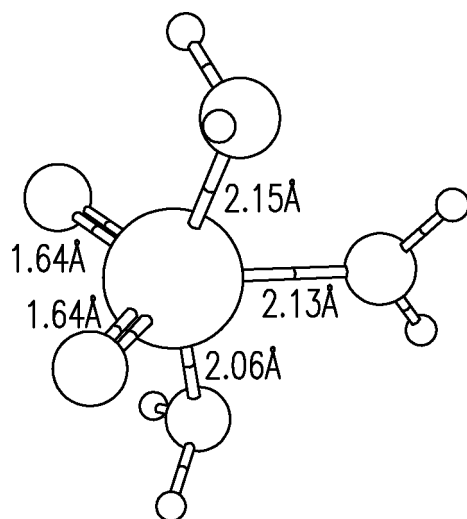
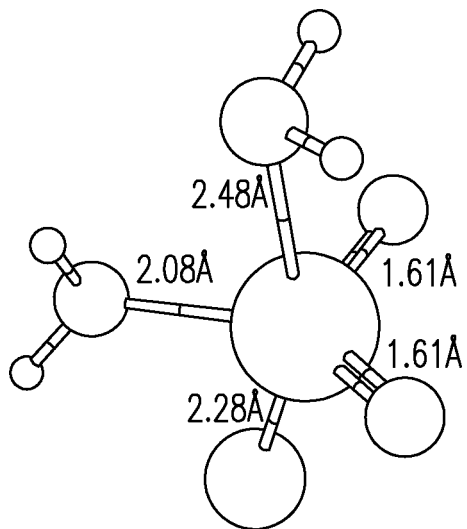


Fig. 5



*Fig. 6a*



*Fig. 6b*

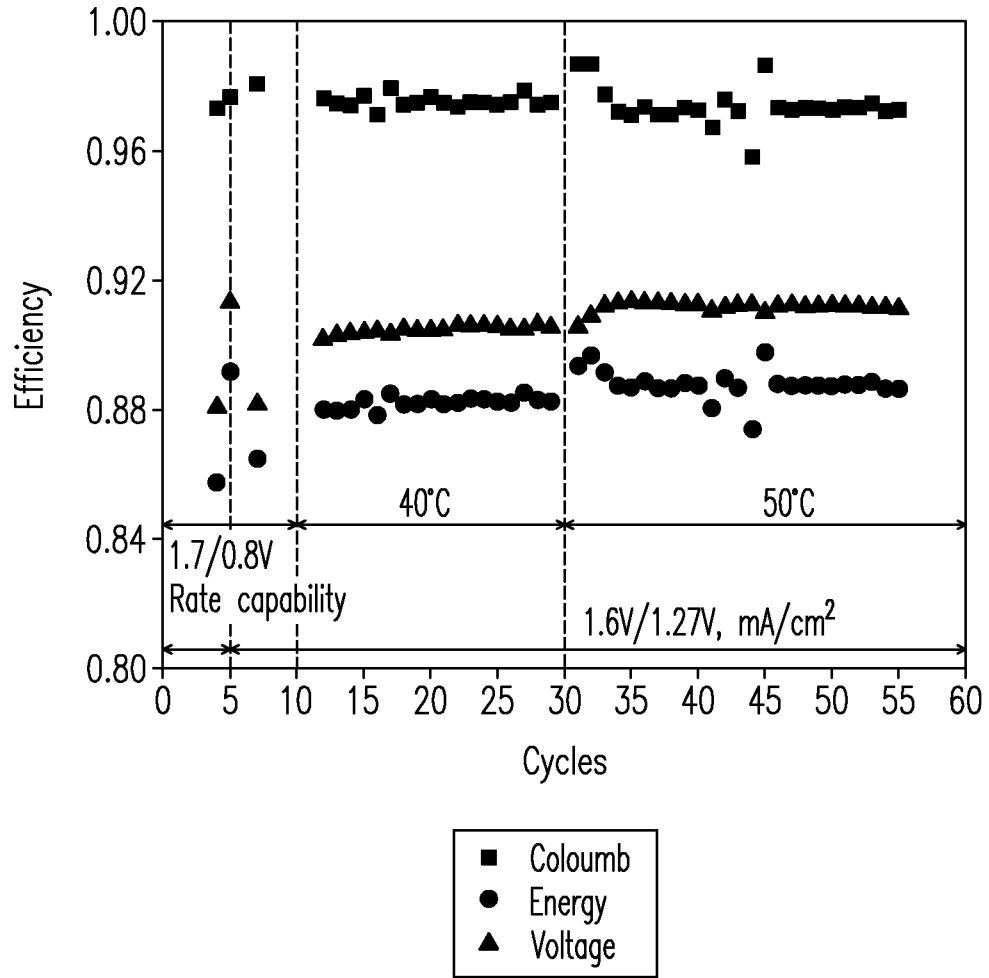
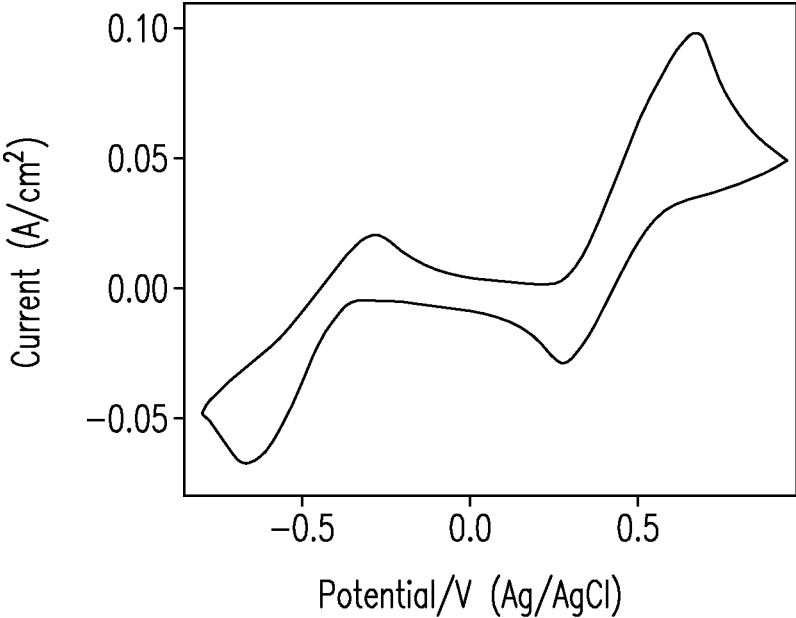
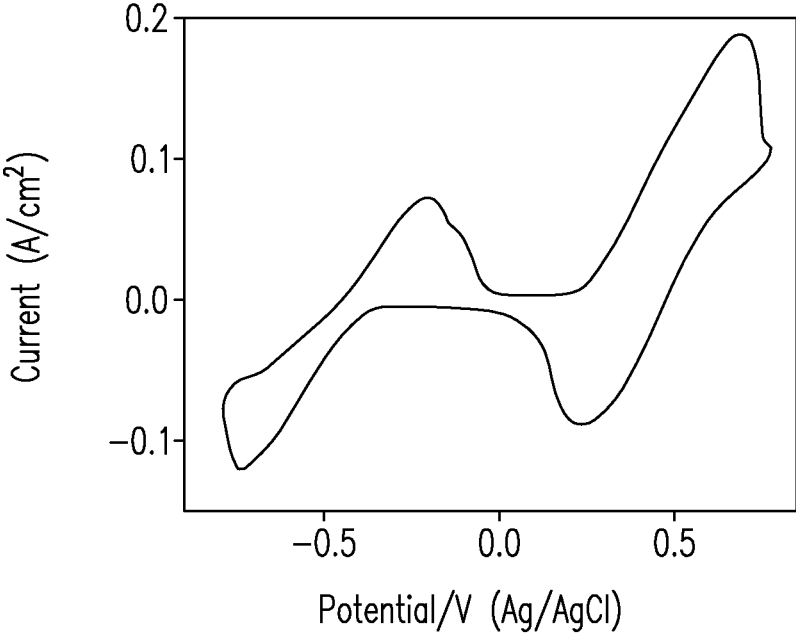


Fig. 7





*Fig. 8a*



*Fig. 8b*

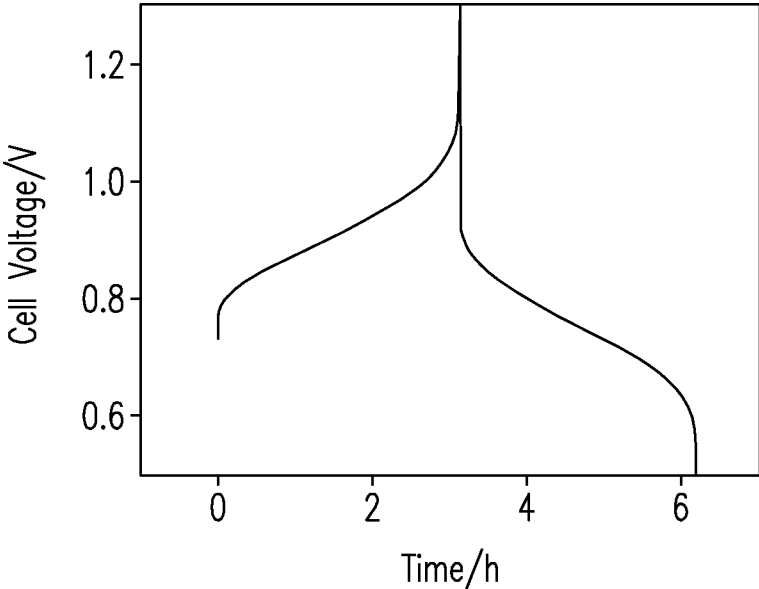


Fig. 9a

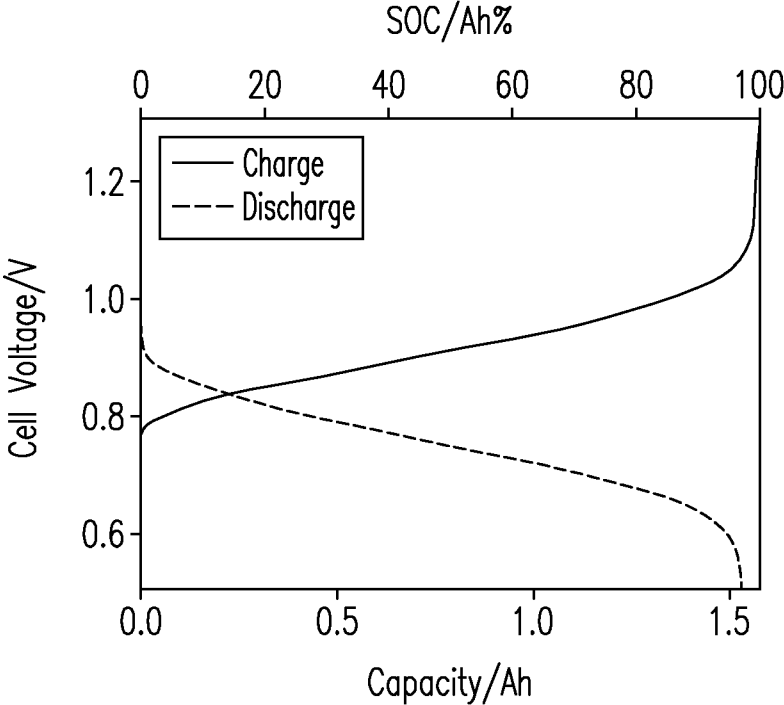


Fig. 9b

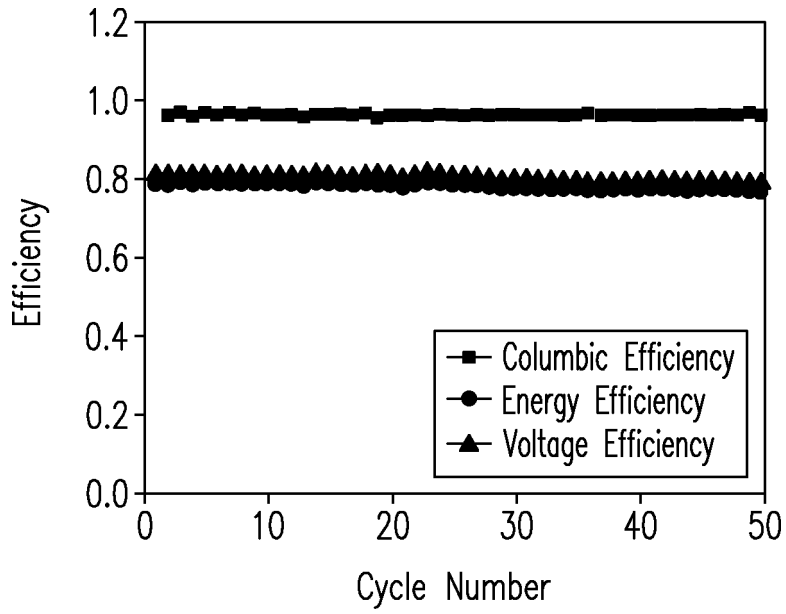


Fig. 9c

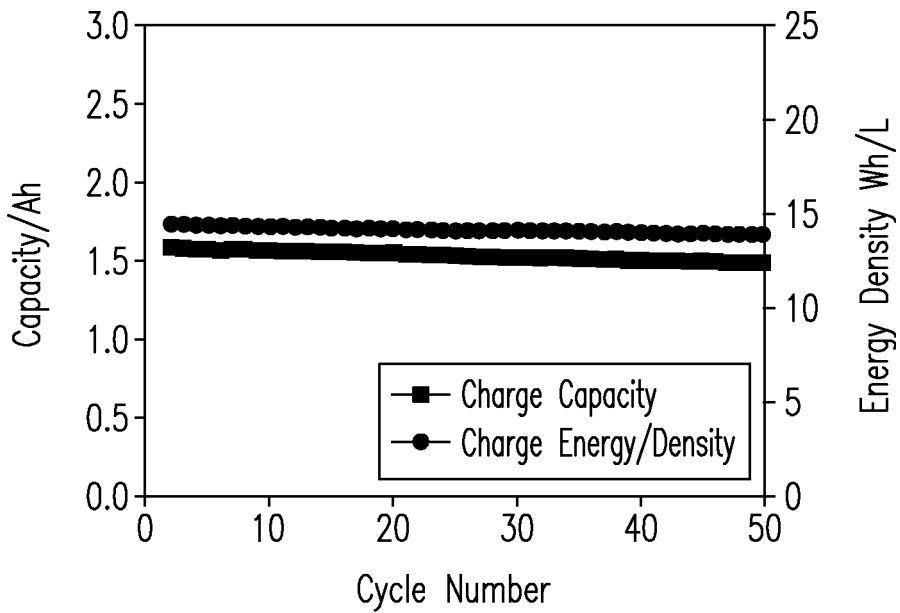


Fig. 9d

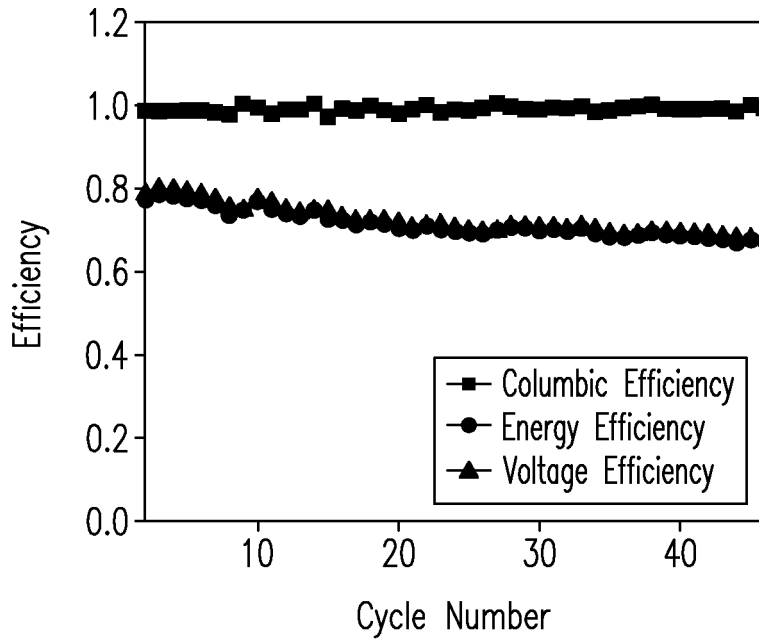


Fig. 10a

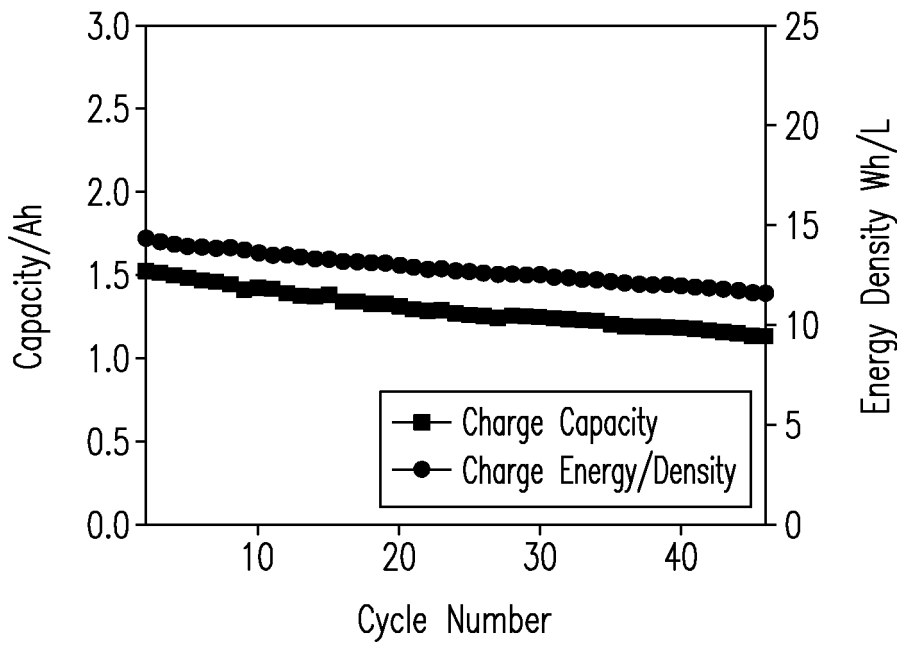


Fig. 10b

1

**FE-V REDOX FLOW BATTERIES**STATEMENT REGARDING FEDERALLY  
SPONSORED RESEARCH OR DEVELOPMENT

This invention was made with Government support under Contract DE-AC0576RL01830 awarded by the U.S. Department of Energy. The Government has certain rights in the invention.

## BACKGROUND

A redox flow battery (RFB) stores electrical energy in reduced and oxidized species dissolved in two separate electrolyte solutions. The anolyte and the catholyte circulate through a cell electrode separated by a porous membrane. Redox flow batteries are advantageous for energy storage because they are capable of tolerating fluctuating power supplies, repetitive charge/discharge cycles at maximum rates, overcharging, overdischarging, and because cycling can be initiated at any state of charge.

However, among the many redox couples upon which redox flow batteries are based, a number of disadvantages exist. For example, many systems utilize redox species that are unstable, are highly oxidative, are difficult to reduce or oxidize, precipitate out of solution, and/or generate volatile gases. In many ways, the existing approaches to addressing these disadvantages have been ad hoc and can include the imposition of restrictive operating conditions, the use of expensive membranes, the inclusion of catalysts on the electrodes, and/or the addition of external heat management devices. These approaches can significantly increase the complexity and the cost of the total system. Therefore, a need for improved redox flow battery systems exists.

## SUMMARY

The present invention includes redox flow battery systems having a supporting solution that comprises  $\text{Cl}^-$  anions. In one embodiment, a vanadium-based redox flow battery system is characterized by an anolyte comprising  $\text{V}^{2+}$  and  $\text{V}^{3+}$  in a supporting solution and a catholyte comprising  $\text{V}^{4+}$  and  $\text{V}^{5+}$  in a supporting solution. The supporting solution can comprise  $\text{Cl}^-$  ions or a mixture of  $\text{SO}_4^{2-}$  and  $\text{Cl}^-$  ions. The use of  $\text{Cl}^-$  ions can improve the energy density and the stability of an all-vanadium battery compared to the traditional use of  $\text{SO}_4^{2-}$  ions.

Supporting solutions comprising both  $\text{SO}_4^{2-}$  and  $\text{Cl}^-$  ions can further improve the performance and characteristics by all-vanadium batteries by increasing the solubility of the vanadium cations as described in greater detail below. In particular embodiments, the concentration ratio of  $\text{Cl}^-$  to  $\text{SO}_4^{2-}$  can be between 1:100 and 100:1. In other embodiments, the ratio can be between 1:10 and 10:1. In still other embodiments, the ratio can be between 1:3 and 3:1.

For all-vanadium batteries, the  $\text{Cl}^-$  in the supporting solution can improve stability of the vanadium cations. For example, in traditional flow redox batteries,  $\text{V}^{5+}$  can tend to form  $\text{V}_2\text{O}_5$  at temperatures above  $40^\circ\text{C}$ . However, the presence of  $\text{Cl}^-$  ions in the supporting solution can result in the formation of  $\text{VO}_2\text{Cl}(\text{H}_2\text{O})_2$ , a stable, neutral species. Accordingly, embodiments of the present invention can operate at cell temperatures greater than  $40^\circ\text{C}$ . Preferably, the cell temperature during operation is between  $-35^\circ\text{C}$ . and  $60^\circ\text{C}$ . Furthermore, the embodiments of the present invention can operate without thermal management devices actively regulating the cell temperature. In conventional all-vanadium flow

2

redox batteries, thermal management devices are required to maintain the battery below the cell temperature at which the V cations come out of solution.

Further still, vanadium cation concentrations in batteries of the present invention can exceed those of traditional  $\text{SO}_4^{2-}$ -based batteries. In some embodiments, the vanadium cation concentration is greater than 0.5M. In others, the vanadium cation concentration is greater than 1.7M. In still others, the vanadium cation concentration is greater than 2.5M.

In a preferred embodiment, the state of charge condition is greater than 0% and less than 100% during operation. In other words, the batteries are preferably not operated to full charge or discharge states.

In another embodiment of the present invention, a redox flow battery having a supporting solution comprising  $\text{Cl}^-$  ions comprises an anolyte comprising  $\text{V}^{2+}$  and  $\text{V}^{3+}$  in the supporting solution, a catholyte comprising  $\text{Fe}^{2+}$  and  $\text{Fe}^{3+}$  in the supporting solution, and a membrane separating the anolyte and the catholyte. The anolyte and catholyte can comprise V cations and Fe cations, respectively, or the anolyte and catholyte can each contain both V and Fe cations in a mixture. In some instances, the concentrations of the Fe cations and/or the V cations can be greater than 0.5M.

Relative to some highly oxidative redox couples, the Fe and V couple is less aggressive. Accordingly, expensive oxidation-resistant membranes such as sulfonated tetrafluoroethylene based fluoropolymer-copolymers are not necessary. On a cost basis, other less expensive options can be preferable. Accordingly, some embodiments of the Fe/V battery system comprise hydrocarbon-based membranes or microporous separators. One example of a hydrocarbon membrane includes, but is not limited to a sulfonated poly(phenylsulfone) membrane. Other ion exchange membranes can be suitable.

In another embodiment, the Fe/V battery system comprises electrodes, which do not contain a redox catalyst, in contact with the anolyte and the catholyte. Redox catalysts are sometimes necessary for species that are difficult to reduce and/or oxidize and can include metals or metal oxides. Redox catalysts are preferably absent from the electrodes used in embodiments of the present invention.

Some embodiments of the Fe/V battery system operate at cell temperatures below  $60^\circ\text{C}$ . In other embodiments, the system operates at cell temperatures between  $-20^\circ\text{C}$ . and  $50^\circ\text{C}$ . In preferred embodiments, the system does not include a heat management device actively regulating the cell temperature. In particular, no heat management device is utilized to heat the Fe/V battery system.

In the Fe/V battery systems, supporting solutions comprising both  $\text{SO}_4^{2-}$  and  $\text{Cl}^-$  ions can further improve the performance and characteristics by increasing the solubility of the cations as described in greater detail below. In particular embodiments, the concentration ratio of  $\text{Cl}^-$  to  $\text{SO}_4^{2-}$  can be between 1:100 and 100:1. In other embodiments, the ratio can be between 1:10 and 10:1. In still other embodiments, the ratio can be between 1:3 and 3:1. Instances in which the supporting solution comprises both  $\text{Cl}^-$  to  $\text{SO}_4^{2-}$  and the anolyte and catholyte both comprise V and Fe cations, the concentration of  $\text{V}^{2+}$ ,  $\text{V}^{3+}$ ,  $\text{Fe}^{2+}$ , and  $\text{Fe}^{3+}$  are greater than 1.5M. Instances in which the anolyte comprises V cations and the catholyte comprises Fe cations, the concentrations of  $\text{V}^{2+}$  and  $\text{V}^{3+}$  are greater than 2M in the anolyte and concentrations of  $\text{Fe}^{2+}$  and  $\text{Fe}^{3+}$  are greater than 2M in the catholyte.

In yet another embodiment, a redox flow battery system comprises a supporting solution that comprises a mixture of  $\text{SO}_4^{2-}$  and  $\text{Cl}^-$ . As described elsewhere herein, a supporting solution having mixed  $\text{SO}_4^{2-}$  and  $\text{Cl}^-$  can provide increased

energy density and improved stability and solubility of one or more of the ionic species in the catholyte and/or anolyte, such as  $\text{Fe}^{2+}$ ,  $\text{Fe}^{3+}$ ,  $\text{Cr}^{2+}$ ,  $\text{Cr}^{3+}$ , and others. In particular embodiments, the concentration ratio of  $\text{Cl}^-$  to  $\text{SO}_4^{2-}$  can be between 1:100 and 100:1. In other embodiments, the ratio can be between 1:10 and 10:1. In still other embodiments, the ratio can be between 1:3 and 3:1. In still other embodiments, other halogen ions can be mixed with  $\text{SO}_4^{2-}$ , including but not limited to,  $\text{F}^-$ .

The purpose of the foregoing abstract is to enable the United States Patent and Trademark Office and the public generally, especially the scientists, engineers, and practitioners in the art who are not familiar with patent or legal terms or phraseology, to determine quickly from a cursory inspection the nature and essence of the technical disclosure of the application. The abstract is neither intended to define the invention of the application, which is measured by the claims, nor is it intended to be limiting as to the scope of the invention in any way.

Various advantages and novel features of the present invention are described herein and will become further readily apparent to those skilled in this art from the following detailed description. In the preceding and following descriptions, the various embodiments, including the preferred embodiments, have been shown and described. Included herein is a description of the best mode contemplated for carrying out the invention. As will be realized, the invention is capable of modification in various respects without departing from the invention. Accordingly, the drawings and description of the preferred embodiments set forth hereafter are to be regarded as illustrative in nature, and not as restrictive.

#### DESCRIPTION OF DRAWINGS

Embodiments of the invention are described below with reference to the following accompanying drawings.

FIG. 1 is a graph of current versus voltage comparing all-vanadium RFBs using chloride-containing and sulfate-containing supporting solutions.

FIG. 2 compares thermodynamic equilibrium concentrations (a) and equilibrium potentials (b) of chlorine and oxygen gases in vanadium chloride RFB systems.

FIG. 3 compares cyclic performances of vanadium chloride RFB systems and vanadium sulfate RFB systems.

FIG. 4 compares cyclic voltammetry curves of a vanadium-chloride-sulfate solution and a vanadium sulfate solution.

FIG. 5 is a graph of equilibrium concentrations of chlorine in the positive side of a vanadium-chloride-sulfate cell at various states of charge.

FIG. 6 is a diagram depicting structures of  $\text{VO}_2^-$  in sulfuric acid (a) and in hydrochloric acid (b).

FIG. 7 is a graph of cyclic coulombic efficiency, voltage efficiency, and energy efficiency for a vanadium-chloride-sulfate RFB system.

FIG. 8 are cyclic voltammetry curves in a Fe/V and Cl-containing solution using two different electrodes.

FIG. 9 contains graphs demonstrating the electrochemical performance of an Fe/V redox flow cell using a Cl-containing supporting solution.

FIG. 10 shows cyclic Coulombic efficiency, voltage efficiency, and energy efficiency (a) as well as cell charge capacity and charge energy density change (b) for a Fe/V cell employing S-Radel as membrane

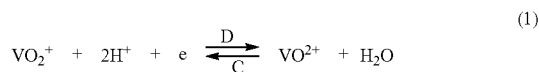
#### DETAILED DESCRIPTION

The following description includes the preferred best mode as well as other embodiments of the present invention. It will

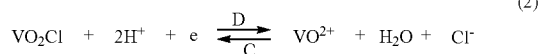
be clear from this description of the invention that the invention is not limited to these illustrated embodiments but that the invention also includes a variety of modifications and embodiments thereto. Therefore the present description should be seen as illustrative and not limiting. While the invention is susceptible of various modifications and alternative constructions, it should be understood, that there is no intention to limit the invention to the specific form disclosed, but, on the contrary, the invention is to cover all modifications, alternative constructions, and equivalents falling within the spirit and scope of the invention as defined in the claims.

FIGS. 1-10 show a variety of embodiments and aspects of the present invention. Referring first to FIG. 1, current versus voltage data is plotted for vanadium ion redox couples using either chloride or sulfate supporting solutions. Three redox couples were observed in the chloride system, indicating that two redox couples ( $\text{VO}^{2+}/\text{VO}_2^+$  for the positive and  $\text{V}^{2+}/\text{V}^{3+}$  for the negative) can be employed for a redox flow battery. Electrochemical reversibility of the  $\text{V}^{4+}/\text{V}^{5+}$  couple was similar to that of a sulfate system, but that of the  $\text{V}^{2+}/\text{V}^{3+}$  was significantly improved in the chloride system. For example, the peak potential difference is 0.351 V in the sulfate system and 0.509 V in the chloride system.

According to quantum chemistry calculations, the  $\text{V}^{5+}$  species in the chloride solution forms  $\text{VO}_2\text{Cl}(\text{H}_2\text{O})_2$ , which is a more stable neutral species than  $[\text{VO}_2(\text{H}_2\text{O})_3]^+$ , the species commonly formed in the sulfate solution. However,  $\text{V}^{2+}$ ,  $\text{V}^{3+}$  and  $\text{V}^{4+}$  in the chloride solution have a similar structure to that in the sulfate solution. Based on the above, the half cell reaction shown in Eq. (2) for the positive pole describes well the electrochemistry. The standard potential of this half cell reaction is expected to be slightly higher than that of the conventional sulfate system resulting from a different  $\text{V}^{5+}$  species. By forming this new structure, the thermal stability of the  $\text{V}^{5+}$  in the chloride solution was significantly improved.



$$E^\circ = 1.0 \text{ V vs. NHE}$$



$$E^\circ = 1.0 \text{ V} + \alpha \text{ V vs. NHE}$$



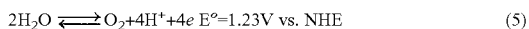
$$E^\circ = -0.25 \text{ V vs. NHE}$$

In the chloride system, oxygen and chlorine gas evolution during charging can reduce coulombic efficiency. Referring to FIG. 2(a), equilibrium concentrations of chlorine or oxygen estimated from thermodynamic equilibrium for Eq. (1) and (4), and Eq. (1) and (5), respectively, are shown as a function at the state of charge (SOC) at various temperatures. It should be noted that hypochlorite can be negligible because the equilibrium constant of Eq. (6) is  $6.35\text{E-}13$  at  $25^\circ \text{C}$ . The actual concentrations of the chlorine should be lower than the values depicted in FIG. 2(a) due to complex formation. Within a typical operation window of redox flow batteries (i.e., SOC of 20-80%), the chlorine concentration is negligible even at  $40^\circ \text{C}$ . However, gas evolution may be significant at SOC values approaching 100%.

Chlorine has much higher solubility in water than oxygen; Henry's constant of chlorine and oxygen in water at  $25^\circ \text{C}$ . is

## 5

0.062 mol/L-atm and 0.0013 mol/L-atm, respectively. Assuming partial pressure of oxygen and chlorine is 0.1 bar, the equilibrium potential of Eq. (4) and (5) was calculated for 2.3 M V in 10 M total chloride system, and is shown in FIG. 2(b) as a function of SOC. Based on the data, VO<sub>2</sub><sup>+</sup> is thermodynamically stable from oxygen evolution below an 80% SOC, and from chlorine evolution below a 98% SOC. To maintain saturation of chlorine in the electrolyte solution, the flow battery is preferably operated in a closed system. A closed system is also advantageous to prevent rapid oxidation of V<sup>2+</sup> and V<sup>3+</sup> by air and to minimize electrolyte loss.



In addition to thermodynamic equilibrium, electrode overpotential can contribute to gas evolution. The equilibrium potential of reaction (4) is higher than that of reaction (5), but oxygen evolution can be negligible compared to chlorine evolution because of a higher overpotential on the electrode. For example, the chlorine evolution overpotential on a graphite porous electrode was 0.12 V at 25° C. at charge current of 22 mA/cm<sup>2</sup> for a Zn/Cl<sub>2</sub> battery (see N. Watanabe, T. Touhara, New Mat. New Processes, 1 (1981) 62). This overpotential was higher than that of the oxidation reaction in Eq. (2) above. Therefore, the chlorine evolution reaction can be negligible except for an SOC of ~100%. Because the electrode overpotential of chlorine evolution decreases with increasing temperature, charging is preferably controlled below SOC of 90–95% to prevent chlorine evolution, especially at elevated temperature.

The thermal stabilities of different vanadium ion species in either sulfate or chloride supporting solutions are summarized in Table 1. In the sulfate system, with more than 1.7 M vanadium, V<sup>2+</sup> and V<sup>4+</sup> experienced precipitation at low temperatures (–5° C. and 25° C.), and V<sup>5+</sup> suffered from precipitation at 40° C. In the chloride system, thermal stability was significantly improved. V<sup>2+</sup>, V<sup>4+</sup> and V<sup>3+</sup> were stable for more than 10 days in the temperature ranges of –5 and 50° C. for 2.3 M vanadium. According to nuclear magnetic resonance data (not shown), V<sup>5+</sup> in the sulfate solution exists as a form of [VO<sub>2</sub>(H<sub>2</sub>O)<sub>3</sub>]<sup>+</sup>. With increasing temperature, this complex decomposed into VO(OH)<sub>3</sub> and H<sub>3</sub>O<sup>+</sup>, and then VO(OH)<sub>3</sub> is converted into a precipitate of V<sub>2</sub>O<sub>5</sub>·3H<sub>2</sub>O. As mentioned elsewhere herein, V<sup>5+</sup> is believed to exist as a stable neutral form of VO<sub>2</sub>Cl(H<sub>2</sub>O)<sub>2</sub> in the chloride solution. Regardless, the supporting solutions comprising Cl<sup>–</sup> can enable better stability at higher temperature.

TABLE 1

Comparison of thermal stability of V <sup>m+</sup> for chloride and sulfate systems.						
V <sup>m+</sup> species	V <sup>m+</sup> [M]	H <sup>+</sup> [M]	SO <sub>4</sub> <sup>2–</sup> [M]	Cl <sup>–</sup> [M]	T (° C.)	Time for precipitation
V <sup>2+</sup>	2	6	5	0	–5	419 hr
	2	6	5	0	25	Stable (>20 d)
	2	6	5	0	40	Stable (>20 d)
V <sup>3+</sup>	2	4	5	0	–5	Stable (>20 d)
	2	4	5	0	25	Stable (>20 d)
	2	4	5	0	40	Stable (>20 d)
V <sup>4+</sup> (VO <sup>2+</sup> )	2	6	5	0	–5	18 hr
	2	6	5	0	25	95 hr
V <sup>5+</sup> (VO <sub>2</sub> <sup>+</sup> )	2	8	5	0	–5	Stable (>20 d)
	2	8	5	0	25	Stable (>20 d)
	2	8	5	0	40	95 hr

## 6

TABLE 1-continued

Comparison of thermal stability of V <sup>m+</sup> for chloride and sulfate systems.						
V <sup>m+</sup> species	V <sup>m+</sup> [M]	H <sup>+</sup> [M]	SO <sub>4</sub> <sup>2–</sup> [M]	Cl <sup>–</sup> [M]	T (° C.)	Time for precipitation
V <sup>2+</sup>	1.8	8.4	5	0	40	358 hr
	2.3	5.4	0	10	–5	Stable (>20 d)
	2.3	5.4	0	10	25	Stable (>20 d)
V <sup>3+</sup>	2.3	5.4	0	10	40	Stable (>20 d)
	1.5	3.0	0	7.5	–5	Stable (>20 d)
	1.8	3.0	0	8.4	–5	124 hr
V <sup>4+</sup> (VO <sup>2+</sup> )	2.3	3.1	0	10	–5	96 hr
	2.3	3.1	0	10	25	Stable (>20 d)
	2.3	3.1	0	10	40	Stable (>20 d)
V <sup>5+</sup> (VO <sub>2</sub> <sup>+</sup> )	2.3	5.4	0	10	–5	Stable (>20 d)
	2.3	5.4	0	10	25	Stable (>20 d)
	2.3	5.4	0	10	40	Stable (>20 d)
V <sup>5+</sup> (VO <sub>2</sub> <sup>+</sup> )	2.3	7.7	0	10	–5	Stable (>20 d)
	2.3	7.7	0	10	25	Stable (>20 d)
	2.3	7.7	0	10	40	Stable (>20 d)
	2.3	7.7	0	10	50	Stable (>10 d)

When operation of an all Cl<sup>–</sup> system occurs at, or below, freezing temperatures (i.e., 0° C.), the tank containing the electrolyte is preferably insulated to maintain waste heat from the flow battery, which can be approximately 20% of total energy. Operation above the freezing temperature, energy density can be improved by approximately 35% owing to higher vanadium concentration compared to the sulfate system. Stabilization of the V<sup>3+</sup> species at the lower temperature can be achieved by using a supporting solution containing both SO<sub>4</sub><sup>2–</sup> and Cl<sup>–</sup>, as is described in greater detail elsewhere herein.

Typical energy efficiency of vanadium redox flow batteries is about 80%; indicating 20% of the energy is released as waste heat during each cycle. Assuming an adiabatic system, the electrolyte temperature can increase by about 4° C. per cycle. The thermal stability of electrolytes at higher temperatures can be a major concern, especially on hot days. For conventional all vanadium sulfate systems, active thermal management devices such as heat exchangers are commonly employed to maintain the cell temperature below 40° C. and to prevent precipitation of V<sup>5+</sup>. An active thermal management system is not preferable and is a significant parasitic energy loss. Embodiments of the present invention based on vanadium and Cl-containing supporting solution can be operated at a wide range of temperatures between 0 to 50° C. without an active thermal management system, improving significant system efficiency and also reducing cost.

Flow cell performance for different chloride and sulfate systems were evaluated under the identical test conditions. The results at different discharging current densities were summarized in Table 2. Energy density of the chloride system was ~38 Wh/L, 30% higher than that of the sulfate system, resulting from the higher solubility of vanadium in the chloride solution. This higher energy density can reduce the system cost by reducing tank size and footprint. Columbic efficiency of the chloride system was 94–97% under operation of SOC between 0 and 100% (not inclusive), comparable to that of the sulfate system, which was 95–97%.

TABLE 2

Comparison of discharging rate capability for VSRFB (1.7M V in 5M total sulphate) and VCRFB (2.3M V in 10M total chloride).										
CD (mA/cm <sup>2</sup> )	Capacity		Energy density*		Efficiency					
	(Ah)		(Wh/L)		Coulomb		Energy		Voltage	
	Cl <sup>-</sup>	SO <sub>4</sub> <sup>2-</sup>	Cl <sup>-</sup>	SO <sub>4</sub> <sup>2-</sup>	Cl <sup>-</sup>	SO <sub>4</sub> <sup>2-</sup>	Cl <sup>-</sup>	SO <sub>4</sub> <sup>2-</sup>	Cl <sup>-</sup>	SO <sub>4</sub> <sup>2-</sup>
100	2.75	2.14	35.5	27.9	0.94	0.95	0.80	0.83	0.85	0.87
75	2.75	2.14	36.6	28.4	0.96	0.96	0.84	0.85	0.87	0.89
50	2.75	2.14	37.8	29.1	0.97	0.96	0.87	0.88	0.90	0.91
25	2.74	2.13	38.7	29.7	0.97	0.97	0.90	0.91	0.92	0.94

\*Note that energy density was calculated only by electrolyte volume.

Cyclic performance of both systems at ambient temperature was also evaluated by cycling between 1.6V and 1.2V, which are shown in FIG. 3. The capacities of both systems slightly decreased with cycles because of different transport rate of vanadium species across the membrane. This capacity loss can be recovered by remixing and rebalancing anolyte and catholyte because a single element of V is used for both solutions. Energy and coulombic efficiencies for the chloride system was stable with cycles and comparable to those of sulfate system. It can be concluded that the novel all vanadium chloride flow battery can be stably operated in a comparable energy efficiency to the sulfate system, while delivering energy density of ~38 Wh/L, 30% higher than the sulfate system. Chlorine evolution or V<sup>5+</sup> electrolyte stability in the chloride solution was not an issue under closed operation conditions.

Electrolyte for the all vanadium chloride systems described above was prepared by dissolving V<sub>2</sub>O<sub>3</sub> in concentrated HCl (38%). The electrolyte for the all vanadium sulfate system was fabricated by dissolving VOSO<sub>4</sub>·3.8 H<sub>2</sub>O in sulfuric acid (95.8%).

Cyclic voltammetry (CV) tests for the chloride system were conducted with identical graphite felts (φ=5 mm mm) used in flow cell testing to identify redox couples and electrochemical reversibility using Solartron 1287 potentiostat. The scan rate was 0.5 mV/s.

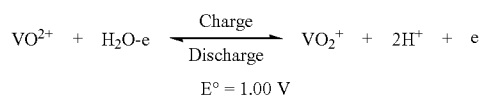
Cell performance of two different systems was measured using a flow cell system under identical test conditions. The apparent area of the graphite felt was 10 cm<sup>2</sup> (2 cm×5 cm), in contact with NAFION 117 membrane, a sulfonated tetrafluoroethylene based fluoropolymer-copolymer. Other proton-exchange membranes can be suitable. 2.3 M vanadium in 10 M total chloride solution and 1.7 M V in 5 M total sulphate solution were used for performance comparison. Each electrolyte volume and flow rate was 50 mL and 20 mL/min, respectively. The effect of different discharging current densities was evaluated in the first 5 cycles with the same charging current of 50 mA/cm<sup>2</sup>. The flow cell was charged to 1.7 V and then discharged to 0.8 V. After that, the flow cell was cycled between 1.6 V and 1.2 V at 50 mA/cm<sup>2</sup>.

The electrolyte stability tests were carried out in polypropylene tubes at -5, 25, 40, and 50° C., using about 5 ml solution for each sample. During the stability tests, the samples were kept static without any agitation, and were monitored daily by naked eye for the formation of precipitation.

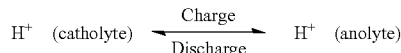
Referring to Table 3, which summarizes the stability of V<sup>2+</sup>, V<sup>3+</sup>, V<sup>4+</sup>, and V<sup>5+</sup> in sulfuric acid solutions, conventional sulfuric acid-only vanadium redox flow batteries (VRFB) can typically only be operated at cell temperatures

between 10° C. and 40° C. with vanadium concentration in the electrolytes less than 1.7 M (with an energy density <25 Wh/L). The electrochemical reactions of an all vanadium sulfate redox flow battery are represented by the following equations.

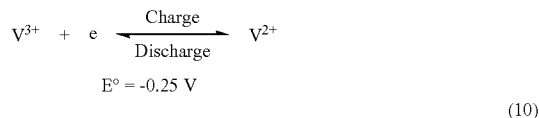
Catholyte: (7)



Membrane: (8)



Anolyte: (9)



Overall: (10)

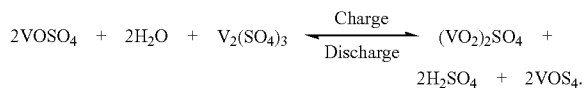
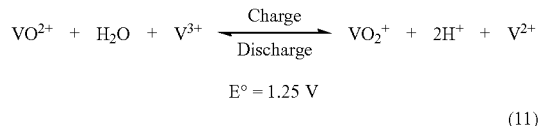


TABLE 3

Stability of V <sup>m+</sup> cations in H <sub>2</sub> SO <sub>4</sub> solution					
V <sup>m+</sup> specie	V <sup>m+</sup> , M	H <sup>+</sup> , M	SO <sub>4</sub> <sup>2-</sup> , M	T, ° C.	Time for precipitation
V <sup>2+</sup>	2	6	5	-5	Stable (>10 d)
	2	6	5	25	Stable (>10 d)
	2	6	5	40	Stable (>10 d)
V <sup>3+</sup>	2	4	5	-5	Stable (>10 d)
	2	4	5	25	Stable (>10 d)
	2	4	5	40	Stable (>10 d)
V <sup>4+</sup>	2	6	5	-5	18 hr

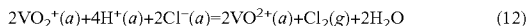


TABLE 3-continued

Stability of V <sup>n+</sup> cations in H <sub>2</sub> SO <sub>4</sub> solution					
V <sup>n+</sup> specie	V <sup>n+</sup> , M	H <sup>+</sup> , M	SO <sub>4</sub> <sup>2-</sup> , M	T, ° C.	Time for precipitation
(VO <sup>2+</sup> )	2	6	5	25	95 hr
	2	6	5	40	Stable (>10 d)
V <sup>5+</sup>	2	8	5	-5	Stable (>10 d)
(VO <sup>2+</sup> )	2	8	5	25	Stable (>10 d)
	2	8	5	40	95 hr

As mentioned earlier, since the standard potential of reaction  $2\text{Cl}^- - 2\text{e} = \text{Cl}_2(\text{g})$  ( $E^\circ = 1.36 \text{ V}$ ) is much higher than that of Reaction (7), the supporting solution in a VRFB system can comprise  $\text{Cl}^-$  either as a  $\text{SO}_4^{2-}$  and  $\text{Cl}^-$  mixture or comprising  $\text{Cl}^-$  as the only anion. Moreover, as is described elsewhere herein, the use of mixed  $\text{SO}_4^{2-}$  and  $\text{Cl}^-$  in the supporting solution is not limited to vanadium-based redox flow batteries. Chloride and sulfate ions in the supporting solution can help stabilize relatively higher concentrations of other cations as well.

FIG. 4 shows the cyclic voltammetry curve of a solution containing 2.5 M  $\text{VOSO}_4$  and 6 M  $\text{HCl}$ . This curve is similar to that of a solution containing 1.5 M  $\text{VOSO}_4$  and 3.5 M  $\text{H}_2\text{SO}_4$ . Referring to FIG. 5, the equilibrium concentrations of  $\text{Cl}_2$  gas in a vanadium sulfate-chloride catholyte solution (containing 2.5 M vanadium, 2.5 M sulfate, and 6 M chloride) under different state-of-charge (SOC) conditions were calculated according to Reaction 12. Under normal flow battery operation conditions (i.e.  $T < 40^\circ \text{C}$ . and  $\text{SOC} < 80\%$ ), the equilibrium concentration of  $\text{Cl}_2$  gas is less than 10 ppm. Due to its high solubility in water (0.57 g  $\text{Cl}_2$  per 100 g water at  $30^\circ \text{C}$ .), most of the  $\text{Cl}_2$  gas generated should be dissolved in the catholyte solutions. At high temperatures, SOC values higher than 80% are preferably avoided to minimize the  $\text{Cl}_2$  gas evolution. Nevertheless, a closed system can be used to minimize continuous  $\text{Cl}_2$  gas generation and to prevent  $\text{Cl}_2$  gas emission to the environment. Such closed systems are commonly required for the conventional vanadium sulfate flow battery system to prevent oxidation of  $\text{V}^{2+}$  and  $\text{V}^{3+}$  by  $\text{O}_2$  in air, and to prevent water loss from electrolyte solutions.



The stability of different V<sup>n+</sup> cations in Cl<sup>-</sup>-containing solutions was evaluated at a temperature range of  $-5^\circ \text{C}$ . to  $40^\circ \text{C}$ . The results are given in Table 4. More than 2.3 M  $\text{VOCl}_2$  and  $\text{VO}_2\text{Cl}$  were stabilized in  $\sim 6 \text{ M}$   $\text{HCl}$  solution over a temperature range of  $-5^\circ \text{C}$ . to  $40^\circ \text{C}$ ., which is much higher than those in the sulfuric acid solution ( $\sim 1.5 \text{ M}$  vanadium) over the same temperature range. The  $\text{Cl}^-$  anions appear to stabilize  $\text{VO}_2^+$  and  $\text{VO}^{2+}$  cations in the solution. Similar to that in the  $\text{H}_2\text{SO}_4$  solution, more than 2.3 M  $\text{V}^{2+}$  was also stabilized in  $\sim 6 \text{ M}$   $\text{HCl}$  solution at  $-5^\circ \text{C}$ . to  $40^\circ \text{C}$ . However, compared to that in the  $\text{H}_2\text{SO}_4$  solution, the stability of  $\text{V}^{3+}$  in  $\text{HCl}$  solution was decreased. At  $-5^\circ \text{C}$ ., only about 1.5 M  $\text{V}^{3+}$  could be stabilized in 3 M  $\text{HCl}$ , whereas more than 2 M  $\text{V}^{3+}$  was stabilized in 2 M  $\text{H}_2\text{SO}_4$  (see Table 4).

TABLE 4

Stability of V <sup>n+</sup> cations in HCl solution					
V <sup>n+</sup> specie	V <sup>n+</sup> , M	H <sup>+</sup> , M	Cl <sup>-</sup> , M	T, ° C.	Time for precipitation
V <sup>2+</sup>	2.3	5.4	10	-5	Stable (>10 d)
	2.3	5.4	10	25	Stable (>10 d)
	2.3	5.4	10	40	Stable (>10 d)

TABLE 4-continued

Stability of V <sup>n+</sup> cations in HCl solution					
V <sup>n+</sup> specie	V <sup>n+</sup> , M	H <sup>+</sup> , M	Cl <sup>-</sup> , M	T, ° C.	Time for precipitation
V <sup>3+</sup>	1.5	3.0	7.5	-5	Stable (>10 d)
	1.8	3.0	8.4	-5	124 hr
	2.3	3.1	10	-5	96 hr
V <sup>4+</sup>	2.3	3.1	10	25	Stable (>10 d)
	2.3	3.1	10	40	Stable (>10 d)
	2.3	5.4	10	-5	Stable (>10 d)
(VO <sup>2+</sup> )	2.3	5.4	10	25	Stable (>10 d)
	2.3	5.4	10	40	Stable (>10 d)
	2.3	5.4	10	40	Stable (>10 d)
V <sup>5+</sup>	2.3	7.7	10	-5	Stable (>10 d)
	2.3	7.7	10	25	Stable (>10 d)
	2.3	7.7	10	40	Stable (>10 d)

Based on the stability test results above,  $\text{Cl}^-$  anions can help stabilizing  $\text{VO}^{2+}$  and  $\text{VO}_2^+$  cations, and  $\text{SO}_4^{2-}$  anions can help stabilize  $\text{V}^{3+}$  cations. Both  $\text{Cl}^-$  and  $\text{SO}_4^{2-}$  anions can stabilize  $\text{V}^{2+}$  cations. Accordingly, a sulfuric acid and hydrochloric acid mixture can stabilize high concentrations of all four vanadium cations. Table 5 gives the stability of different V<sup>n+</sup> cations in two mixed  $\text{SO}_4^{2-}$  and  $\text{Cl}^-$  solutions at  $-5^\circ \text{C}$ . to  $40^\circ \text{C}$ . Without optimization, about 2.5 M of all four V<sup>n+</sup> cations were effectively stabilized in the 2.5 M  $\text{SO}_4^{2-}$ -6 M  $\text{Cl}^-$  mixed acid solution. At a higher vanadium concentration (3M),  $\text{V}^{2+}$ ,  $\text{VO}^{2+}$ , and  $\text{VO}_2^+$  were also stabilized in the 3 M  $\text{SO}_4^{2-}$ -6 M  $\text{Cl}^-$  mixed acid solution at  $-5^\circ \text{C}$ . to  $40^\circ \text{C}$ . However,  $\text{V}^{3+}$  was only stable for about 8 days at  $-5^\circ \text{C}$ . Precipitation of  $\text{VOCl}$  was observed. Due to the large amount of heat generation during the operation of a VRFB system, it is not difficult to keep the cell temperature of the electrolytes higher than  $-5^\circ \text{C}$ . even when the ambient temperature is  $-5^\circ \text{C}$ . or lower. Also, since a VRFB system is always operated under 80 to 90% state-of-charge and state-of-discharge conditions, the highest concentration of  $\text{V}^{3+}$  in a 3 M all vanadium flow battery system is 2.7 M (mixing with 0.3 M  $\text{V}^{2+}$ , at the end of 90% discharge) or 2.4 M (mixing with 0.6 M  $\text{V}^{2+}$ , at the end of 80% discharge). Therefore, in one embodiment, by using a sulfuric acid and hydrochloric acid mixture as the supporting solution, the VRFB system uses a supporting solution with a total vanadium concentration higher than 3 M.

TABLE 5

Stability of V <sup>n+</sup> in the $\text{SO}_4^{2-}$ + $\text{Cl}^-$ solutions							
V <sup>n+</sup> specie	V <sup>n+</sup> [M]	H <sup>+</sup> [M]	SO <sub>4</sub> <sup>2-</sup> [M]	Cl <sup>-</sup> [M]	T (° C.)	Time for precipitation	
V <sup>2+</sup>	3	6	3	6	-5	Stable (>10 d)	
	2.5	6	2.5	6	-5	Stable (>10 d)	
	2.5	6	2.5	6	25	Stable (>10 d)	
	2.5	6	2.5	6	40	Stable (>10 d)	
	3	6	3	6	40	Stable (>10 d)	
V <sup>3+</sup>	3	3	3	6	-5	192 hr (8 d)	
	2.5	3.5	2.5	6	-5	Stable (>10 d)	
	2.5	3.5	2.5	6	25	Stable (>10 d)	
	2.5	3.5	2.5	6	40	Stable (>10 d)	
	3	3	3	6	40	Stable (>10 d)	
V <sup>4+</sup>	3	6	3	6	-5	Stable (>10 d)	
	(VO <sup>2+</sup> )	2.5	6	2.5	6	-5	Stable (>10 d)
	2.5	6	2.5	6	25	Stable (>10 d)	
	2.5	6	2.5	6	40	Stable (>10 d)	
	3	6	3	6	40	Stable (>10 d)	
V <sup>5+</sup>	3	9	3	6	-5	Stable (>10 d)	
	(VO <sup>2+</sup> )	2.5	8.5	2.5	6	-5	Stable (>10 d)
	2.5	8.5	2.5	6	25	Stable (>10 d)	
	2.5	8.5	2.5	6	40	Stable (>10 d)	
	3	9	3	6	40	Stable (>10 d)	
2.7 V <sup>3+</sup> +	7.7	3	6	50	Stable (>10 d)		

TABLE 5-continued

Stability of V <sup>n+</sup> in the SO <sub>4</sub> <sup>2-</sup> + Cl <sup>-</sup> solutions						
V <sup>n+</sup> specie	V <sup>n+</sup> [M]	H <sup>+</sup> [M]	SO <sub>4</sub> <sup>2-</sup> [M]	Cl <sup>-</sup> [M]	T (° C.)	Time for precipitation
0.3 V <sup>4+</sup>						
2.7 V <sup>5+</sup> + 0.3 V <sup>4+</sup>		7.7	3	6	60	Stable (>10 d)

At temperatures higher than 40° C., in traditional all-vanadium sulfate RFBs the stability of V<sup>5+</sup> might decrease due to the formation of V<sub>2</sub>O<sub>5</sub>. However, as shown in Table 5, embodiments of the present invention using mixed SO<sub>4</sub><sup>2-</sup>-Cl<sup>-</sup> solutions exhibit excellent stability with a mixture of 2.7 M V<sup>5+</sup> and 0.3 M V<sup>4+</sup> (corresponding to 90% of state-of-charge of a 3 M VRFB system) at temperatures as high as 60° C., indicating that Cl<sup>-</sup> anions can effectively stabilize the VO<sub>2</sub><sup>+</sup> cations. As described elsewhere herein, quantum chemistry calculations show that, in Cl<sup>-</sup>-containing solutions, a stable neutral species can form having the formula VO<sub>2</sub>Cl(H<sub>2</sub>O)<sub>2</sub>. Referring to FIG. 6, a diagram depicts the molecular structure of [VO<sub>2</sub>(H<sub>2</sub>O)<sub>3</sub>]<sup>+</sup> and of VO<sub>2</sub>Cl(H<sub>2</sub>O)<sub>2</sub>. In this structure, one Cl<sup>-</sup> anion, two O<sup>2-</sup> anions, and two H<sub>2</sub>O molecules complex with one V<sup>5+</sup> in the first coordination shell. Without Cl<sup>-</sup> anions in the solution, two O<sup>2-</sup> anions, and three H<sub>2</sub>O molecules complex with V<sup>5+</sup> in the first coordination shell and a positively-charged specie with [VO<sub>2</sub>(H<sub>2</sub>O)<sub>3</sub>]<sup>+</sup> formula forms. Quantum chemistry calculations also indicate that, at elevated temperatures, this positively charged species is prone to convert to V<sub>2</sub>O<sub>5</sub>-3H<sub>2</sub>O by de-protonation (Reaction 13) and condensation (Reaction 14). The structural differences appear to account for the much improved stability of VO<sub>2</sub><sup>+</sup> cations in Cl<sup>-</sup>-containing solutions. Due to the forma-

lost as waste heat during each cycle. For an adiabatic system, this heat can raise the temperature of the whole system by about 5° C. Due to the large amount of waste heat generation, stability of electrolytes at high temperature range is a major concern, especially during hot days. The embodiments of the present invention encompassing all-vanadium RFBs utilizing mixed SO<sub>4</sub><sup>2-</sup>-Cl<sup>-</sup> supporting solutions system can not only improve the energy density, but can also expand the operation temperature window from 10-40° C. to -5-60° C. During the cold winter days, limited insulation can easily keep the temperature of the system above -5° C. Accordingly, in preferred embodiments, no active heat management is needed

Several small VRFB cells were used to evaluate the performances of two vanadium sulfate-chloride mixed systems (with 2.5 M and 3.0 M vanadium). For comparison, the performance of a vanadium sulfate system (with 1.6 M vanadium) was also measured. The results are summarized in Table 6. The sulfate-chloride mixed systems show much higher energy density than the sulfate system. Even with higher vanadium concentration, the all vanadium sulfate-chloride mixed systems still showed similar energy efficiency to that of the vanadium sulfate system. FIG. 7 provides the cyclic coulombic efficiency, voltage efficiency, and energy efficiency of the 2.5 M all vanadium sulfate-chloride mixed acid system at different ambient temperatures. Stable performance was observed with this new system. During a course of 20 days of operation, the gas-phase pressures of the anolyte and catholyte containers remained constant, indicating no significant gas evolution occurred in the whole system. The viscosity and density of a solution containing 2.5 M VOSO<sub>4</sub> and 6 M HCl at 30° C. is 6.1 cP and 1.40 g/ml respectively, slightly lower than the 6.4 cP and 1.45 g/ml for a solution containing 2.0 M VOSO<sub>4</sub> and 3.0 M H<sub>2</sub>SO<sub>4</sub>.

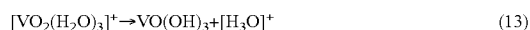
TABLE 6

Performance of all vanadium redox flow cells using the mixed SO <sub>4</sub> <sup>2-</sup> -Cl <sup>-</sup> supporting solutions												
Current of	Energy Density Wh/L			Coulombic Efficiency η <sub>c</sub>			Energy Efficiency η <sub>E</sub>			Voltage Efficiency η <sub>V</sub>		
	2.5VS 6Cl	3VS 6Cl	1.6V 4.5S	2.5VS 6Cl	3VS 6Cl	1.6V 4.5S	2.5VS 6Cl	3VS 6Cl	1.6V 4.5S	2.5VS 6Cl	3VS 6Cl	1.6V 4.5S
100	36.2	39.5	22.3	0.95	0.95	0.94	0.81	0.76	0.83	0.85	0.80	0.88
75	37.5	40.8	22.4	0.96	0.96	0.94	0.84	0.81	0.85	0.88	0.84	0.90
50	38.5	41.8	22.6	0.96	0.97	0.94	0.87	0.85	0.87	0.91	0.88	0.92
25	39.2	43.1	22.6	0.96	0.97	0.94	0.90	0.89	0.88	0.93	0.91	0.94

1. Cell operation conditions: 10 cm<sup>2</sup> flow cell, Charged to 1.7 V by 50 mA/cm<sup>2</sup> current.

2. 2.5VS 6HCl: 2.5M V 2.5M SO<sub>4</sub><sup>2-</sup> 6M Cl<sup>-</sup>; 3VS6HCl: 3M V 3M SO<sub>4</sub><sup>2-</sup> 6M Cl<sup>-</sup>; 1.6V 4.5S: 1.6M V 4.5M SO<sub>4</sub><sup>2-</sup>.

tion of stable VO<sub>2</sub>Cl(H<sub>2</sub>O)<sub>2</sub> structure, the equilibrium concentration of Cl<sub>2</sub> gas in the catholyte solution should be lower than that shown in FIG. 5.



In embodiments comprising mixed SO<sub>4</sub><sup>2-</sup>-Cl<sup>-</sup> solutions, the stability of V<sup>4+</sup> is controlled by the solubility of VOSO<sub>4</sub>, and the stability of V<sup>3+</sup> is controlled by the solubility of VOCl. The improvement of V<sup>4+</sup> stability is due to the decrease of SO<sub>4</sub><sup>2-</sup> concentration in the solution, and the improvement of V<sup>3+</sup> stability is due to the decrease of Cl<sup>-</sup> concentration. V<sup>2+</sup> cation is stable in both Cl<sup>-</sup> and SO<sub>4</sub><sup>2-</sup>-containing solutions.

In traditional all-vanadium sulfate RFBs, energy efficiency is about 80%, which means about 20% of the total energy is

The experiment details related to the all-vanadium RFBs using mixed SO<sub>4</sub><sup>2-</sup>-Cl<sup>-</sup> supporting solutions are as follows. The flow cells consisted of two graphite electrodes, two gold-coated copper current collectors, two PTFE gaskets, and a Nafion® 117 membrane. The active area of the electrode and the membrane was about 10 cm<sup>2</sup>. An Arbin battery tester was used to evaluate the performance of flow cells and to control the charging and discharging of the electrolytes. A Solartron 1287 potentiostat was employed for cyclic voltammetry (CV) experiments. The flow rate was fixed at 20 mL/min, which was controlled by a peristaltic pump. A balanced flow cell contained about 50 mL anolyte and 50 mL catholyte.

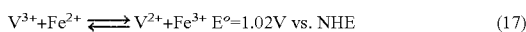
For cell performance evaluation and electrolyte solution preparation, the cell was normally charged at a current density of 50 mA/cm<sup>2</sup> to 1.7 V and discharged to 0.8 V with a

current density of 25 to 100 mA/cm<sup>2</sup>. Cell cycling tests were performed at 90% state-of-charge and state-of-discharge at a fixed charging and discharging current density of 50 mA/cm<sup>2</sup>.

The electrolyte solutions of V<sup>2+</sup>, V<sup>3+</sup>, V<sup>4+</sup> and V<sup>5+</sup> used in this work were prepared electrochemically in flow cells using VOSO<sub>4</sub> (from Alfa Aesar) and VCl<sub>3</sub> as starting chemicals. VCl<sub>3</sub> solutions were prepared by dissolving V<sub>2</sub>O<sub>3</sub> (from Alfa Aesar) in HCl solutions. The electrolyte stability tests were carried out in polypropylene tubes at -5° C., ambient temperature, 40° C., 50° C., and 60° C., using about 5 ml solution for each sample. During the stability tests, the samples were kept static without any agitation, and were monitored daily by naked eye for the formation of precipitation. Solution viscosity was measured using a Ubbelohde calibrated viscometer tube.

Thermodynamic calculations of reaction 2VO<sub>2</sub><sup>+(a)</sup>+4H<sup>+</sup>(a)+2Cl<sup>-</sup>(a)=2VO<sup>2+</sup> were carried out using HSC Chemistry® 6.1 program from Outotec Research Oy. Quantum chemistry calculations were carried out using the Amsterdam Density Functional (ADF) program.

Yet another embodiment of the present invention encompasses a redox flow battery system based on the redox couple of Fe and V. In this system, the anolyte comprises V<sup>2+</sup> and V<sup>3+</sup> in the supporting solution while the catholyte comprises Fe<sup>2+</sup> and Fe<sup>3+</sup> in the supporting solution. The redox reactions and their standard potentials can be described as follows:



The Fe/V system of the present invention can provide significant benefits while circumventing some of the intrinsic issues of conventional systems. For example, certain embodiments of the Fe/V system do not use catalysts and/or high-temperature management systems, which add to the complexity and cost of the system. Moreover the evolution of H<sub>2</sub> gas during charging is minimized, since the working potential of V<sup>2+</sup>/V<sup>3+</sup> (-0.25 V) is considerably higher than that of others, such as Cr<sup>2+</sup>/Cr<sup>3+</sup> (-0.41 V). In the catholyte, the Fe<sup>2+</sup>/Fe<sup>3+</sup> redox couple is electrochemically reversible and can be less oxidative than other common ionic species, such as V<sup>4+</sup>/V<sup>5+</sup>, which can result in higher stability at high temperatures while avoiding expensive, oxidation-resistant membrane materials, such as sulfonated tetrafluoroethylene based fluoropolymer-copolymer.

In one example using mixed Fe and V reactant solutions, an electrolyte for Fe/V redox flow battery tests was prepared by dissolving VCl<sub>3</sub> (99%) and FeCl<sub>2</sub> (98%) in concentrated HCl (38%). Cyclic voltammetry (CV) was carried out in Fe/V+HCl solutions with various concentrations to identify redox couples and electrochemical reversibility using a SOLARTRON 1287 potentiostat (SOLARTRON ANALYTICAL, USA). A Pt wire and Ag/AgCl electrode were used as the counter and reference electrodes, respectively. Glassy carbon electrodes and graphite felt (φ=5.5 mm) sealed onto a metal current collector were used as the working electrodes. The scan rate was 0.5 mV/s. Identical graphite felts without redox catalysts were used in both CV and flow cell testing.

Cell performance was measured under constant current methods using a flow cell system. The apparent area of graphite felt was 10 cm<sup>2</sup> (2 cm×5 cm), in contact with membrane. 1.25 M Fe/V in 2.3 M HCl solution and 1.17 M Fe/V in 2.15 M HCl solution were used with either a sulfonated tetrafluoroethylene based fluoropolymer-copolymer (i.e., NAFION) or a low-cost hydrocarbon membrane such as sulfonated poly

(phenylsulfone) membrane (i.e., S-RADEL), respectively. Each electrolyte volume and flow rate was 50 mL and 20 mL/min. The flow cell was charged to 1.3 V and then discharged to 0.5 V at a current density of 50 mA/cm<sup>2</sup>.

The chemical stability of commercially available membranes was determined by soaking them in 0.15 M Fe<sup>3+</sup> and 7 M total chloride solution at 40° C., and in 0.1 M V<sup>5+</sup> and 5 M total sulfate solution for comparison. During the stability tests, the samples were kept static without any agitation, and were monitored daily by naked eye for changes of color indicating oxidation of the membrane.

FIGS. 8(a) and (b) show CV results of 1.5 M Fe and V in a 1 M hydrochloric acid supporting solution using glassy carbon and graphite felt electrode, respectively. The current density is normalized to the geometrical area of the working electrode. Similar CV spectra were observed on both the glassy carbon and graphite felt working electrode with the graphite felt electrode demonstrating higher over potential due to the low conductivity. Two redox peaks were observed indicating two redox reactions, Fe<sup>3+</sup>/Fe<sup>2+</sup> for the positive and V<sup>2+</sup>/V<sup>3+</sup> for the negative. More importantly, no significant hydrogen evolution current was observed at potentials below the V<sup>3+</sup> reduction peak, indicating that no significant gas evolution occurred at the negative electrode during the charging process when employing a V<sup>2+</sup>/V<sup>3+</sup> redox couple. Oxidation and reduction peaks appear in the forward and reverse scans on the positive side, which revealed a reversible redox couple of Fe<sup>3+</sup>/Fe<sup>2+</sup> with a potential at approximately 0.5 V. Similarly, there is no anodic current observed associated with evolution of the Cl<sub>2</sub> and/or O<sub>2</sub> gas. Thus, the Fe<sup>3+</sup>/Fe<sup>2+</sup> and V<sup>3+</sup>/V<sup>2+</sup> redox couples in chloride supporting solution can be used in the negative and positive half cells according to embodiments of the present invention.

FIG. 3 shows the results of Fe/V flow cell testing with a NAFION 117 membrane. A plot of cell voltage versus time is given in FIG. 3(a). FIG. 3(b) demonstrates the cell voltage profile with respect to the cell capacity and the cell SOC. The SOC is calculated against the maximum charge capacity. Referring to FIG. 3(b), the Fe/V redox flow cell can be charged and discharged to a SOC in the range of 0~100%. A utilization ratio of close to 100% can be achieved. Up to 50 cycles, the Fe/V cell demonstrated stable columbic efficiency of ~97%, energy efficiency of ~78%, and voltage efficiency of ~80% as shown in FIG. 3(c). The Fe/V cell also demonstrated excellent capacity and energy density retention capability as shown in FIG. 3(d) with 0.1% loss per cycle in charge capacity over 50 cycles.

Commercially available, low-cost membranes, including a micro-porous separator, can be used in place of relatively expensive NAFION (i.e., sulfonated tetrafluoroethylene based fluoropolymer-copolymer) membranes. Suitable alternative membranes can include, but are not limited to, hydrocarbon-based commercially available ion-exchange membranes; for example, SELEMION® anion exchange membrane (APS, from Asahi Glass, Japan), SELEMION® cation exchange membrane (CMV, from Asahi Glass, Japan), and sulfonated poly(phenylsulfone) membrane (S-RADEL® (RADEL® from Solvay Advanced Polymers, USA), and micro-porous separators typically used in lithium battery, for example: CELGARD® micro-porous separator (Celgard, USA).

The electrochemical performance of a Fe/V cell employing a S-RADEL membrane was then evaluated using identical test protocols to that of Nafion membrane. The cell performance data is shown in FIGS. 5(a) and (b). Similar Coulombic efficiency, voltage efficiency, and energy efficiency with that of Nafion membrane were obtained.

15

In a preferred embodiment, the energy density of Fe/V RFB systems can be improved by using a supporting solution comprising SO<sub>4</sub><sup>2-</sup>Cl<sup>-</sup> mixed ions to increase the reactant concentration in the anolyte and catholyte. Referring to Table 7, the solubility of Fe<sup>2+</sup> and Fe<sup>3+</sup> ions is higher in H<sub>2</sub>SO<sub>4</sub>—HCl mixed acids than in hydrochloric acid.

TABLE 1

Stability of Fe <sup>n+</sup> cations in the H <sub>2</sub> SO <sub>4</sub> —HCl mixed solutions						
Fe <sup>n+</sup> specie	Fe <sup>n+</sup> , M	H <sup>+</sup> , M	SO <sub>4</sub> <sup>2-</sup> , M	Cl <sup>-</sup> , M	T, ° C.	Time for precipitation
Fe <sup>2+</sup>	2	4	2	4	25	Stable (>6 d)
Fe <sup>3+</sup>	2	6	2	6	25	Stable (>6 d)

While a number of embodiments of the present invention have been shown and described, it will be apparent to those skilled in the art that many changes and modifications may be made without departing from the invention in its broader aspects. The appended claims, therefore, are intended to cover all such changes and modifications as they fall within the true spirit and scope of the invention.

We claim:

1. A redox flow battery system having:  
A membrane separating an anolyte and a catholyte;  
the anolyte comprising V<sup>2+</sup> and V<sup>3+</sup> in a negative supporting solution comprising an initial concentration of at least 0.5M Fe cations and Cl<sup>-</sup> anions;  
the catholyte comprising Fe<sup>2+</sup> and Fe<sup>3+</sup> in a positive supporting solution comprising an initial concentration of at least 0.5M V cations and Cl<sup>-</sup> anions;  
wherein the initial concentration of Fe cations in the negative supporting solution is substantially equal to the initial concentration of V cations in the positive supporting solution.
2. The system of claim 1, wherein the Fe<sup>2+</sup> and Fe<sup>3+</sup> concentrations are greater than 0.5 M.
3. The system of claim 1, wherein the V<sup>2+</sup> and V<sup>3+</sup> concentrations are greater than 0.5M.
4. The system of claim 1, wherein the membrane is a hydrocarbon-based membrane.
5. The system of claim 1, wherein the membrane is a micro-porous separator.

16

6. The system of claim 1, wherein the membrane is not a sulfonated tetrafluoroethylene based fluoropolymer-copolymer.

7. The system of claim 1, further comprising negative and positive electrodes in contact with the anolyte and the catholyte, respectively, wherein the electrodes do not contain a redox catalyst.

8. The system of claim 1, having a cell temperature less than 60° C. during operation.

9. The system of claim 1, having a cell temperature between -20° C. and 50° C. during operation.

10. The system of claim 1, wherein the positive supporting solution, the negative supporting solution, or both further comprises SO<sub>4</sub><sup>2-</sup> anions.

11. The system of claim 10, wherein the Cl<sup>-</sup> to SO<sub>4</sub><sup>2-</sup> concentration ratio is between 1:100 and 100:1.

12. The system of claim 10, wherein the Cl<sup>-</sup> to SO<sub>4</sub><sup>2-</sup> concentration ratio is between 1:10 and 10:1.

13. The system of claim 10, wherein the Cl<sup>-</sup> and SO<sub>4</sub><sup>2-</sup> concentration ratio is between 1:3 and 3:1.

14. The system of claim 12, wherein concentrations of V<sup>2+</sup>, V<sup>3+</sup>, Fe<sup>2+</sup>, and Fe<sup>3+</sup> are greater than 1M in the anolyte and in the catholyte.

15. The system of claim 10, wherein concentrations of V<sup>2+</sup> and V<sup>3+</sup> are greater than 1.5M in the anolyte and concentrations of Fe<sup>2+</sup> and Fe<sup>3+</sup> are greater than 1.5M in the catholyte.

16. The system of claim 1, wherein said cations of V have a concentration greater than 0.5 M in the positive supporting solution and said cations of Fe have a concentration greater than 0.5 M in the negative supporting solution.

17. The system of claim 1, having a catholyte side of the membrane and an anolyte side of the membrane, wherein the catholyte side of the membrane has a total concentration of Fe and V that is substantially equal to that of the anolyte side of the membrane.

18. A redox flow battery system comprising:  
A membrane separating an anolyte and a catholyte  
the anolyte comprising a total concentration of V<sup>2+</sup> and V<sup>3+</sup> cations greater than 0.5 M in a an initial negative supporting solution comprising Cl<sup>-</sup> anions and greater than a 0.5 M concentration of Fe cations;  
the catholyte comprising a total concentration of Fe<sup>2+</sup> and Fe<sup>3+</sup> cations greater than 0.5 M in a an initial positive supporting solution comprising Cl<sup>-</sup> anions and greater than a 0.5 M concentration of V cations.

\* \* \* \* \*