The current invention relates to the preparation of an improved cathode active material for non-aqueous lithium electrochemical cell. In particular, the cathode active material comprises ε-phase silver vanadium oxide prepared by using a γ-phase silver vanadium oxide starting material. The reaction of γ-phase SVO with a silver salt produces the novel ε-phase SVO possessing a lower surface area than γ-phase SVO produced from vanadium oxide (V₂O₅) and a similar silver salt as starting materials. Consequently, the low surface area ε-phase SVO material provides an advantage in greater long-term stability in pulse dischargeable cells.
PREPARATION OF EPSILON-PHASE SILVER VANADIUM OXIDE FROM GAMMA-PHASE SVO STARTING MATERIAL

CROSS REFERENCE TO RELATED APPLICATION

[0001] This application is a divisional of application Ser. No. 10/004,995, filed Dec. 5, 2001, which claims priority based on provisional application Ser. No. 60/254,918, filed Dec. 12, 2000.

BACKGROUND OF THE INVENTION

[0002] 1. Field of the Invention

[0003] This invention relates to the conversion of chemical energy to electrical energy. More particularly, this invention relates to the preparation of an improved cathode active material for non-aqueous lithium electrochemical cells, and still more particularly, a cathode active ε-phase silver vanadium oxide (SVO, Ag₃VₓOₓₙ) prepared using a γ-phase silver vanadium oxide (Agₓ₂VₓOₓₙ) starting material. The product cathode active material can be used in an implantable electrochemical cell, for example of the type powering a cardiac defibrillator, where the cell may run under a light load for significant periods interrupted from time to time by high rate pulse discharges.

[0004] The reaction of γ-phase SVO with a source of silver produces ε-phase SVO that possesses a lower surface area than SVO produced from other vanadium-containing starting materials. The relatively low surface area of this new ε-phase SVO material results in greater long-term stability for the cathode active material in comparison to other forms of SVO with higher specific surface areas.

[0005] 2. Prior Art

[0006] The prior art discloses many processes for manufacturing SVO; however, they result in a product with greater surface area than the material prepared by the current invention. Specifically, U.S. Pat. No. 4,391,729 to Liang et al. discloses the preparation of silver vanadium oxide by a thermal decomposition reaction of silver nitrate with vanadium oxide conducted under an air atmosphere. This decomposition reaction is further detailed in the publication: Leisinger, R. A.; Takeuchi, E. S. Chem. Mater. 1993, 5, 738-742, where the syntheses of SVO from silver nitrate and vanadium oxide under an air atmosphere is presented as a function of temperature. In another reference: Leisinger, R. A.; Takeuchi, E. S. Chem. Mater. 1994, 6, 489-495, the synthesis of SVO from different silver precursor materials (silver nitrate, silver nitrite, silver oxide, silver vanadate, and silver carbonate) is described. The product active materials of this latter publication are consistent with the formation of a mixture of SVO phases prepared under argon, which is not solely ε-phase Ag₃VₓOₓₙ.


Thus, Fleury and Kohlmuller teach that the heat treatment of starting materials under a non-oxidizing atmosphere (such as argon) results in the formation of SVO with a reduced silver content.

[0008] In U.S. Pat. No. 5,955,218 to Crespi et al., the process of heat-treating SVO prepared by a thermal decomposition reaction to improve the electrochemical performance of the material is disclosed. In this patent, thermal decomposition SVO prepared according to U.S. Pat. Nos. 4,310,609 and 4,391,729, both to Liang et al., under an air atmosphere at a somewhat lower temperature of 360° C. is described. However, the 218 patent to Crespi et al. demonstrates that adding a second heat treatment step increases the crystallinity of the resulting active material. The present invention is concerned with the product active material's surface area, and not necessarily its crystallinity.

[0009] U.S. Pat. No. 5,221,453 to Crespi teaches a method for making an electrochemical cell containing SVO, in which the cathode active material is prepared by a chemical addition reaction of an admixed 2:1 mole ratio of AgVO₄ and VₓOₓ heated in the range of 300° C. to 700° C. for a period of 5 to 24 hours. Crespi does not discuss γ-phase SVO in the context of this invention. Therefore, this process could not manufacture the ε-phase material described by the current invention.

[0010] U.S. Pat. Nos. 6,130,005 and 5,955,218, both to Crespi et al., relate to heat treated silver vanadium oxide materials, for example, γ-phase SVO heat treated to form decomposition-produced SVO (dSVO). The starting material does not appear to be heated for further combination with a source of silver or other metal. Also, U.S. Pat. No. 5,895,733 to Crespi et al. shows a method for synthesizing SVO by using AgO and a vanadium oxide as starting materials. However, the result is not a low surface area ε-phase SVO cathode material, as disclosed in the current invention.

[0011] U.S. Pat. No. 5,545,497 to Takeuchi et al. teaches cathode materials having the general formula of AgₓVₓOₓ. Suitable materials comprise a β-β-phase SVO having in the general formula x=0.35 and y=5.18 and a γ-phase SVO having x=0.74 and y=5.37, or a mixture of the phases thereof. Such SVO materials are produced by the thermal decomposition of a silver salt in the presence of vanadium pentoxide. In addition, U.S. Pat. No. 6,171,729 to Gan et al. shows exemplary alkali metal/solid electrochemical cells in which the cathode may be an SVO of β₁, γ₁ or ε-phase materials. However, none of Gan et al.’s methods are capable of producing a low surface area ε-phase cathode material, as per the current invention.

[0012] Therefore, based on the prior art, there is a need to develop a process for the synthesis of mixed metal oxides, including silver vanadium oxide, having a relatively low surface area. An example is a low surface area SVO prepared using a silver-containing compound and γ-phase SVO as starting materials. The product ε-phase SVO is a cathode active material useful for non-aqueous electrochemical cells having enhanced characteristics, including the high pulse capability necessary for use with cardiac defibrillators.

SUMMARY OF THE INVENTION

[0013] The current invention relates to the preparation of an improved cathode active material for non-aqueous
lithium electrochemical cells, and in particular, a cathode active material that contains e-phase SVO prepared using a γ-phase SVO starting material. The reaction of γ-phase SVO with a source of silver produces e-phase SVO possessing a lower surface area than e-phase SVO produced from other vanadium-containing starting materials. The present synthesis technique is not, however, limited to silver salts since salts of copper, magnesium and manganese can be used to produce relatively low surface area metal oxide active materials as well. The relatively low surface area of the e-phase SVO material provides an advantage in greater long-term stability when used as an active cathode material compared to SVO with a higher specific surface area.

DETAILED DESCRIPTION OF THE PREFERRED EMBODIMENTS

[0014] The current invention discloses that reacting a γ-phase SVO material with a source of silver, or other suitable metal salt, produces pure e-phase SVO (Ag₂V₁/V₂O₁₃). This product material possesses a relatively lower surface areas in comparison to active materials synthesized by a thermal decomposition reaction under an oxidizing atmosphere. Decreased surface area is an unexpected result.

[0015] The thermal reaction of silver nitrate with vanadium oxide under an air atmosphere is a typical example of the preparation of silver vanadium oxide by a decomposition reaction. This reaction is set forth below in Equation 1:

$$\text{AgNO}_3 + 2\text{V}_2\text{O}_5 \rightarrow \text{Ag}_2\text{V}_4\text{O}_{13} + 2\text{NO}_x$$  \hspace{1cm} (1)

[0017] The physical characteristics of SVO material (i.e. particle morphology, surface area, crystallinity, etc.) produced by this reaction are dependent on the temperature and time of reaction. In addition, the reaction environment has a dramatic effect on the product material. The same reaction of silver nitrate with vanadium oxide conducted under an argon atmosphere is depicted below in Equation 2:

$$2\text{AgNO}_3 + 2\text{V}_2\text{O}_5 \rightarrow 2\text{AgVO}_3 + 2\text{Ag}_2\text{V}_3\text{O}_9 + 2\text{NO}_x$$  \hspace{1cm} (2)

[0018] Thus, the synthesis of SVO under an inert atmosphere results in the formation of a mixture of silver vanadate (Ag₂VO₃) and γ-phase SVO (Ag₂V₁/V₂O₁₃). This is described in the above-referenced publication by Leising, R. A.; Takeuchi, E. S. Chem. Mater. 1994, 6, 489-495. As reported by Leising et al., a mixture of material phases is less suitable than a single e-phase SVO (Ag₂V₁/V₂O₁₃) as a cathode active material for lithium electrochemical cells. For this reason, argon is typically not preferred for synthesis of SVO cathode active material. A more benign preparation technique for e-phase SVO from vanadium oxide and silver carbonate (Ag₂CO₃) according to Equation 3 below results in the release of CO₂ gas, which is a nontoxic byproduct. However, the specific surface area of the product SVO is also higher than the surface area of SVO prepared from silver nitrate. This is shown below in Table 1.

$$\text{Ag}_2\text{CO}_3 + 2\text{V}_2\text{O}_5 \rightarrow \text{Ag}_2\text{V}_3\text{O}_9 + \text{CO}_2$$  \hspace{1cm} (3)

[0019] Thus, a synthesis technique for SVO using vanadium oxide and either silver oxide or silver carbonate, or other preferred metal salts, while eliminating the formation of toxic NOₓ byproduct, results in an SVO material with a higher specific surface area than SVO produced from vanadium oxide and silver nitrate.

TABLE 1. Specific Surface Area of e-Phase SVO

<table>
<thead>
<tr>
<th>Starting Materials</th>
<th>Synthesis Temperature</th>
<th>BET Surface Area</th>
</tr>
</thead>
<tbody>
<tr>
<td>V₂O₅ + AgNO₃</td>
<td>500° C.</td>
<td>0.42 m²/g</td>
</tr>
<tr>
<td>V₂O₅ + 0.5 AgNO₃</td>
<td>500° C.</td>
<td>0.64 m²/g</td>
</tr>
<tr>
<td>V₂O₅ + 0.5 AgCO₃</td>
<td>500° C.</td>
<td>0.81 m²/g</td>
</tr>
<tr>
<td>Ag₆.5/V₃O₉ + 0.15 AgNO₃</td>
<td>500° C.</td>
<td>0.54 m²/g</td>
</tr>
<tr>
<td>Ag₆.5/V₃O₉ + 0.15 AgCO₃</td>
<td>500° C.</td>
<td>0.44 m²/g</td>
</tr>
</tbody>
</table>

[0020] The present invention is an alternate preparation synthesis that does not produce noxious by-products, such as NOₓ and, additionally, results in an active material with a desirable relatively low surface area. Benefits attributed to the present synthesis process for the formation of a cathode active material are illustrated in the following examples:

EXAMPLE 1

[0021] In contrast to the prior art syntheses described above, SVO of the present invention is prepared using γ-phase SVO (Ag₂V₁/V₂O₁₃) as a starting material instead of V₂O₅. In particular, a 12.90-gram sample of Ag₁.₂/V₂O₃ was combined with a 1.09-gram sample of Ag₂O, and heated to 500° C. under a flowing air atmosphere for about 16 hours. The sample was then cooled, mixed and reheated under a flowing air atmosphere at about 500° C. for about 24 hours. At this point, the material was cooled and analyzed by x-ray powder diffraction and BET surface area measurements. The x-ray powder diffraction data confirmed the formation of e-phase SVO (Ag₂V₁/V₂O₁₃). The material displayed a BET surface area of 0.54 m²/g.

COMPARATIVE EXAMPLE 1

[0022] As a comparison, SVO was prepared by a prior art combination reaction. In particular, a 9.00-gram sample of V₂O₅ was combined with a 5.73-gram sample of Ag₂O, and heated to about 500° C. under a flowing air atmosphere for about 16 hours. The sample was then cooled, mixed and reheated under a flowing air atmosphere at about 500° C. for about 24 hours. At this point the material was cooled and analyzed by x-ray powder diffraction and BET surface area measurements. The material displayed a BET surface area of 0.64 m²/g, which is significantly higher than the specific surface area of the material prepared in Example 1.

EXAMPLE 2

[0023] e-phase SVO according to the present invention was also prepared using a γ-phase SVO starting material in combination with silver carbonate. In particular, a 5.00-gram sample of Ag₁.₂/V₂O₉ was combined with a 0.50-gram sample of Ag₂CO₃, and heated to about 500° C. under a flowing air atmosphere for about 16 hours. The sample was then cooled, mixed and reheated under a flowing air atmosphere at about 500° C. for about 24 hours. At this point, the material was cooled and analyzed by x-ray powder diffraction and BET surface area measurements. The x-ray powder diffraction data confirmed the formation of e-phase SVO (Ag₂V₁/V₂O₁₃), while the material displayed a BET surface area of 0.44 m²/g.

COMPARATIVE EXAMPLE 2

[0024] As a comparison to Example 2, SVO was prepared using V₂O₅ and Ag₂CO₃. In particular, a 15.00-gram sample
of V₂O₅ was combined with an 11.37-g sample of Ag₂O, and heated to about 450 °C. under a flowing air atmosphere for about 16 hours. The sample was then cooled, mixed and reheated under a flowing air atmosphere at about 500 °C. for about 24 hours. At this point the material was cooled and analyzed by x-ray powder diffraction and BET surface area measurements. The material displayed a BET surface area of 0.81 m²/g, which is nearly twice the specific surface area of the material prepared in Example 2.

EXAMPLE 3

[0025] Copper silver vanadium oxide or CSVO (Cs0.2Ag0.8V₂O₅) was prepared according to the present invention using γ-phase SVO as a starting material in combination with copper(II) oxide. In particular, a 1.80-gram sample of Ag₁₂.V₄O₈.₅ was combined with a 0.10-gram sample of CuO, and heated to about 450 °C. under a flowing air atmosphere for about 16 hours. The sample was then cooled, mixed and reheated under a flowing air atmosphere at about 500 °C. for about 24 hours. At this point, the material was cooled and analyzed by BET surface area measurements. The material displayed a BET surface area of 0.31 m²/g.

COMPARATIVE EXAMPLE 3

[0026] As a comparison to the product of Example 3, CSVO was prepared via the prior art decomposition method using V₂O₅, Cu(NO₃)₂ and AgNO₃. In particular, a 1.36 gram sample of V₂O₅ was combined with a 0.99 gram sample of AgNO₃ and a 0.34 gram sample of Cu(NO₃)₂·2.5H₂O, and heated to about 400 °C. under a flowing air atmosphere for about 16 hours. The sample was then cooled, mixed and reheated under a flowing air atmosphere at about 500 °C. for about 44 hours. At this point, the product material was cooled and analyzed by BET surface area measurement. The material displayed a BET surface area of 0.45 m²/g, which is significantly higher than the specific surface area of the CSVO material prepared in Example 3. Thus, in addition to the toxic implications of released NO₂ gas, the preparation of CSVO by the prior art method provides a material with a higher specific surface area than the new preparation technique.

[0027] The above detailed description and examples are intended for the purpose of illustrating the invention, and are not to be construed as limiting. For example, starting materials other than silver oxide and silver carbonate are reacted with γ-phase silver vanadium oxide to form e-phase silver vanadium compounds. The list includes: silver lactate (Ag₂C₆H₅O₇·H₂O, Tₘ 120 °C), silver triflate (AgCF₃SO₃, Tₘ 286 °C), silver pentafluorobispropionate (AgC₅F₅O₂, Tₘ 242 °C), silver laurate (AgC₁₂H₂₅O₂, Tₘ 212 °C), silver myristate (AgC₁₄H₂₅O₂, Tₘ 211 °C), silver palmitate (AgC₁₄H₂₇O₂, Tₘ 209 °C), silver stearate (AgC₁₇H₃₃O₂, Tₘ 205 °C), silver vanadate (AgV₂O₇·rH₂O), copper carbonate (Cu₂CO₃), copper oxide (CuO, Tₘ 1,446 °C), copper carbonat (Cu₂CO₃), manganese carbonate (MnCO₃), manganese oxide (MnO, Tₘ 1,650 °C), magnesium carbonate (MgCO₃, Tₘ 350 °C), magnesium oxide (MgO, Tₘ 2,826 °C), and combinations and mixtures thereof.

[0028] While the starting materials are described as being heated to a preferred temperature of about 500 °C., it is contemplated by the scope of the present invention that suitable heating temperatures range from about 300 °C. to about 550 °C., depending on the specific starting materials. Also, heating times for both the first and second heating steps range from about 5 hours to about 30 hours. Longer heating times are required for lower heating temperatures. Further, while the present invention has been described in the examples as requiring two heating events with an ambient mixing in between, that is not necessarily imperative. Some synthesis protocols according to the present invention may require one heating step with periodic mixing, or multiple heating events with periodic ambient mixing.

[0029] The product mixed metal oxides according to the present invention includes: e-phase SVO (Ag₉V₂O₁₉), CSVO (Cs₀.₂Ag₀.₈V₂O₅), MnSVO (Mn₁₋₅Ag₁₋₅V₂O₅), and MgSVO (Mg₁₋₅Ag₁₋₅V₂O₅). The use of the above mixed metal oxides as a cathode active material provides an electrochemical cell that possesses sufficient energy density and discharge capacity required meeting the vigorous requirements of implantable medical devices. These types of cells comprise an anode of a metal selected from Groups IA, IIA and IIB of the Periodic Table of the Elements. Such anode active materials include lithium, sodium, potassium, etc., and their alloys and intermetallic compounds including, for example, Li—Mg, Li—Si, Li—Al, Li—B and Li—Si—B alloys and intermetallic compounds. The preferred anode comprises lithium. An alternate anode comprises a lithium alloy such as a lithium-aluminum alloy. The greater the amounts of aluminum present by weight in the alloy, however, the lower the energy density of the cell.

[0030] The form of the anode may vary, but preferably the anode is a thin metal sheet or foil of the anode metal, pressed or rolled on a metallic anode current collector, i.e., preferably comprising titanium, titanium alloy or nickel, to form an anode component. Copper, tungsten and tantalum are also suitable materials for the anode current collector. In the exemplary cell of the present invention, the anode component has an extended tab or lead of the same material as the anode current collector, i.e., preferably nickel or titanium, integrally formed therewith such as by welding and contacted by a weld to a cell case of conductive metal in a case-negative electrical configuration. Alternatively, the anode may be formed in some other geometry, such as a bobbin shape, cylinder or pellet to allow an alternate low surface cell design.

[0031] Before the previously described e-phase active materials are fabrication into an anode electrode for incorporation into an electrochemical cell, they are preferably mixed with a binder material, such as a powdered fluoropolymer, more preferably powdered polytetrafluoro-ethylene or powdered polyvinylidenefluoride, present at about 1 to about 5 weight percent of the cathode mixture. Further, up to about 10 weight percent of a conductive diluent is preferably added to the cathode mixture to improve conductivity. Suitable materials for this purpose include acetylene black, carbon black and/or graphite or a metallic powder such as of nickel, aluminum, titanium and stainless steel. The preferred cathode active mixture thus includes a powdered fluoropolymer binder present at about 3 weight percent, a conductive diluent present at about 3 weight percent and about 94 weight percent of the cathode active material. For example, depending on the application of the electrochemical cell, the range of cathode compositions is...
from about 99% to about 80%, by weight, ε-phase silver vanadium oxide mixed with carbon graphite and PTFE.

[0032] Cathode components for incorporation into an electrochemical cell according to the present invention may be prepared by rolling, spreading or pressing the cathode active materials onto a suitable current collector selected from the group consisting of stainless steel, titanium, tantalum, platinum, gold, aluminum, cobalt-nickel alloys, nickel-containing alloys, highly alloyed ferritic stainless steel containing molybdenum and chromium, and nickel-, chromium- and molybdenum-containing alloys. The preferred current collector material is titanium and, most preferably, the titanium cathode current collector has a thin layer of graphite/carbon material, iridium, iridium oxide or platinum applied thereto. Cathodes prepared as described above may be in the form of one or more plates or operatively associated with at least one or more plates of anode material, or in the form of a strip wound with a corresponding strip of anode material in a structure similar to a “jellyroll”. In order to prevent internal short circuit conditions, the cathode is separated from the Group IA, IIA or IIB anode by a suitable separator material. The separator is the electrically insulative material, and the separator material also is chemically unreactive with the anode and cathode active materials and both chemically unreactive with and insoluble in the electrolyte. In addition, the separator material has a degree of porosity sufficient to allow flow there through of the electrolyte during the electrochemical reaction of the cell. Illustrative separator materials include fabrics woven from fluoropolymeric fibers including polyvinylidene fluoride, polyethylene tetrafluoroethylene, and polyethylene chlorotrifluoroethylene used either alone or laminated with a fluoropolymeric microporous film, nonwoven glass, polypropylene, polyethylene, glass fiber materials, ceramics, a polytetrafluoroethylene membrane commercially available under the designation TFELEX (Chempast Inc.), a polypropylene membrane commercially available under the designation CELGARD (Celanese Plastics Company, Inc.) and a membrane commercially available under the designation DEXIGLAS (C. H. Dexter, Div., Dexter Corp.).

[0034] The electrochemical cell of the present invention further includes a nonaqueous, ionically conductive electrolyte that serves as a medium for migration of ions between the anode and the cathode electrodes during the electrochemical reactions of the cell. The electrochemical reaction at the electrodes involves conversion of ions in atomic or molecular forms that migrate from the anode to the cathode. Thus, nonaqueous electrolytes suitable for the present invention are substantially inert to the anode and cathode materials, and they exhibit those physical properties necessary for ionic transport, namely, low viscosity, low surface tension and wettability.

[0035] A suitable electrolyte has an inorganic, ionically conductive salt dissolved in a nonaqueous solvent, and more preferably, the electrolyte includes an ionicizable alkali metal salt dissolved in a mixture of aprotic organic solvents comprising a low viscosity solvent and a high permittivity solvent. The inorganic, ionically conductive salt serves as the vehicle for migration of the anode ions to intercalate or react with the cathode active material. Preferably, the ion forming alkali metal salt is similar to the alkali metal comprising the anode.

[0036] In the case of an anode comprising lithium, the alkali metal salt of the electrolyte is lithium based salt. Known lithium salts that are useful as a vehicle for transport of alkali metal ions from the anode to the cathode include LiPF₆, LiBF₄, LiAsF₆, LiSbF₆, LiClO₄, LiO₂, LiAlCl₄, LiGaCl₄, LiCl₂SO₄CF₃, Li₃N(SO₂CF₂)₂, LiSCN, Li₂O₂, LiC₆F₅SO₃, LiO₂CF₃, LiSO₂F, LiH(C₆H₅)₄, LiCF₃SO₃, and mixtures thereof.

[0037] Low viscosity solvents useful with the present invention include esters, linear and cyclic ethers and dialkyl carbonates such as tetrahydrofuran (THF), methyl acetate (MA), diglyme, triglyme, tetraglyme, dimethyl carbonate (DMC), 1,2-dimethoxyethane (DME), 1,2-dioxyethane (DEE), 1-ethoxy-2-methoxyethane (EHE), ethyl methyl carbonate, methyl propyl carbonate, ethyl propyl carbonate, diethyl carbonate, dipropyl carbonate, and mixtures thereof. Suitable high permittivity solvents include cyclic carbonates, cyclic esters and cyclic amides such as propylene carbonate (PC), ethylene carbonate (EC), butylene carbonate (BC), acetonitrile, dimethyl sulfoxide, dimethyl formamide, dimethyl acetamide, γ-valerolactone, γ-butyrolactone (GBL), N-methyl-pyrrolidinone (NMP), and mixtures thereof. In the present invention, the preferred anode is lithium metal and the preferred electrolyte is 0.8M to 1.5M LiAsF₆ or LiPF₆ dissolved in 50:50 mixture, volume, of propylene carbonate as the preferred high permittivity solvent and 1,2-dimethoxyethane as the preferred low viscosity solvent.

[0038] The preferred form of a primary alkali metal/solid cathode electrochemical cell is a case-negative design wherein the anode is in contact with a conductive metal casing and the cathode contacted to a current collector is the positive terminal. The cathode current collector is in contact with a positive terminal pin via a lead of the same material as the current collector. The lead is welded to both the current collector and the positive terminal pin for electrical contact.

[0039] A preferred material for the casing is titanium although stainless steel, mild steel, nickel-plated mild steel and aluminum are also suitable. The casing header comprises a metallic lid having an opening to accommodate the glass-to-metal seal/terminal pin feedthrough for the cathode electrode. The anode electrode is preferably connected to the case or the lid. An additional opening is provided for electrolyte filling. The casing header comprises elements having compatibility with the other components of the electrochemical cell and is resistant to corrosion. The cell is thereafter filled with the electrolyte solution described hereinabove and hermetically sealed such as by close-welding a titanium plug over the fill hole, but not limited thereto. The cell of the present invention can also be constructed in a case-positive design.

[0040] It is appreciated that various modifications to the inventive concepts described herein may be apparent to those of ordinary skill in the art without departing from the spirit and scope of the present invention as defined by the appended claims.

What is claimed is:

1. A cathode for an electrochemical cell, the cathode comprising an ε-phase silver vanadium oxide characterized as prepared by heating a silver vanadium compound mixed with a metal salt to form a reaction mixture heated to at least
one reaction temperature in an oxidizing atmosphere to produce the e-phase silver vanadium oxide having the formula Ag$_3$V$_2$O$_{11}$.

2. The cathode of claim 1 wherein the silver vanadium compound is γ-phase silver vanadium oxide having the formula Ag$_{1.2}$V$_2$O$_{8.1}$.

3. The cathode of claim 1 wherein the metal salt is selected from the group consisting of silver lactate, silver trilate, silver pentahydropropionate, silver laurate, silver myristate, silver palmitate, silver stearate, silver vanadate, silver oxide, silver carbonate, copper oxide, copper carbonate, manganese carbonate, manganese oxide, magnesium carbonate, magnesium oxide, and combinations and mixtures thereof.

4. The cathode of claim 1 wherein the metal salt is Ag$_2$O and the e-phase silver vanadium oxide has a BET surface area of about 0.54 m$^2$/g.

5. The cathode of claim 1 wherein the metal salt is Ag$_2$CO$_3$ and the e-phase silver vanadium oxide has a BET surface area of about 0.44 m$^2$/g.

6. The cathode of claim 1 wherein the reaction mixture is heated to the at least one reaction temperature in a range from about 300$^\circ$ C to about 550$^\circ$ C.

7. The cathode of claim 1 wherein the reaction mixture is heated to the at least one reaction temperature for about 5 hours to about 30 hours.

8. The cathode of claim 1 further comprising a binder and a conductive material.

9. A cathode for an electrochemical cell, the cathode comprising an electrode active material characterized as prepared from γ-phase silver vanadium oxide having the formula Ag$_{1.2}$V$_2$O$_{8.1}$ mixed with a metal salt compound to form a reaction mixture heated to at least one reaction temperature in an oxidizing atmosphere to produce the electrode active material selected from the group consisting of Ag$_2$V$_2$O$_{11}$, Cu$_{0.2}$Ag$_{0.8}$V$_2$O$_{5.6}$, Mn$_{0.2}$Ag$_{0.8}$V$_2$O$_{5.6}$, and Mg$_{0.2}$Ag$_{0.8}$V$_2$O$_{5.6}$.

10. The cathode of claim 9 wherein the metal salt is selected from the group consisting of silver lactate, silver trilate, silver pentahydropropionate, silver laurate, silver myristate, silver palmitate, silver stearate, silver vanadate, silver oxide, silver carbonate, copper oxide, copper carbonate, manganese carbonate, manganese oxide, magnesium carbonate, magnesium oxide, and combinations and mixtures thereof.

11. The cathode of claim 9 wherein the metal salt is Ag$_2$O such that the product electrode active material having the formula Ag$_2$V$_2$O$_{11}$ has a BET surface area of about 0.54 m$^2$/g.

12. The cathode of claim 9 wherein the metal salt is Ag$_2$CO$_3$ such that the product electrode active material having the formula Ag$_2$V$_2$O$_{11}$ has a BET surface area of about 0.44 m$^2$/g.

13. The cathode of claim 9 wherein the metal salt is CuO such that the product electrode active material having the formula Cu$_{0.2}$Ag$_{0.8}$V$_2$O$_{5.6}$ has a BET surface area of about 0.31 m$^2$/g.

14. A nonaqueous electrochemical cell, comprising:

a) an anode;

b) a cathode containing an active material comprising an e-phase silver vanadium oxide compound characterized as having been prepared from a mixture of a silver vanadium compound and a metal salt forming a reaction mixture heated to at least one reaction temperature in an oxidizing atmosphere to produce the e-phase silver vanadium oxide having the formula Ag$_3$V$_2$O$_{11}$;

c) a non-aqueous electrolyte activating the anode and the cathode; and

d) a separator material electrically insulating the anode from the cathode, and of a porosity to allow for electrolyte flow.

15. The electrochemical cell of claim 14 wherein the anode is comprised of lithium.

16. The electrochemical cell of claim 14 wherein the silver vanadium containing compound is γ-phase silver vanadium oxide having the formula Ag$_{1.2}$V$_2$O$_{8.1}$.

17. The electrochemical cell of claim 14 wherein the metal salt is selected from the group consisting of silver lactate, silver trilate, silver pentahydropropionate, silver laurate, silver myristate, silver palmitate, silver stearate, silver vanadate, silver oxide, silver carbonate, copper oxide, copper carbonate, magnesium carbonate, manganese carbonate, manganese oxide, magnesium carbonate, magnesium oxide, and combinations and mixtures thereof.

18. The electrochemical cell of claim 14 wherein the metal salt is Ag$_2$O and the e-phase silver vanadium oxide has a BET surface area of about 0.54 m$^2$/g.

19. The electrochemical cell of claim 14 wherein the metal salt is Ag$_2$CO$_3$ and the e-phase silver vanadium oxide has a BET surface area of about 0.44 m$^2$/g.

20. The electrochemical cell of claim 14 wherein the reaction mixture is heated to the at least one reaction temperature in a range from about 300$^\circ$ C to about 550$^\circ$ C.

21. The electrochemical cell of claim 14 wherein the reaction mixture is heated to the at least one reaction temperature for about 5 hours to about 30 hours.