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CA 2459111 C 2014/11/18

(11)(21) **2 459 111**

(12) **BREVET CANADIEN**
CANADIAN PATENT

(13) **C**

(86) Date de dépôt PCT/PCT Filing Date: 2002/09/30
(87) Date publication PCT/PCT Publication Date: 2003/04/10
(45) Date de délivrance/Issue Date: 2014/11/18
(85) Entrée phase nationale/National Entry: 2004/02/27
(86) N° demande PCT/PCT Application No.: US 2002/031048
(87) N° publication PCT/PCT Publication No.: 2003/030279
(30) Priorité/Priority: 2001/10/01 (US09/968,112)

(51) Cl.Int./Int.Cl. *H01M 6/16* (2006.01),
H01M 10/0564 (2010.01), *H01M 4/136* (2010.01),
H01M 4/134 (2010.01)
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(54) Titre : ADJONCTION DIRECTE DE BETA-AMINOENONES DANS DES ELECTROLYTES ORGANIQUES DE
CELLULES NON AQUEUSES UTILISANT DES CATHODES SOLIDES
(54) Title: DIRECT ADDITION OF BETA-AMINOENONES IN ORGANIC ELECTROLYTES OF NONAQUEOUS CELLS
EMPLOYING SOLID CATHODES

(57) Abrégé/Abstract:

A method for manufacturing a nonaqueous cell employing an anode, a cathode and an organic electrolyte wherein the cathode comprises an active cathode material, and the organic electrolyte contains a beta-aminoenone in a range of 0.1 to 5.0 volume percent based on the volume of the electrolyte solvent so as to aid in reducing the undesirably high initial open circuit voltage normally observed with the use of cathodes such as FeS_2 .



(12) INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(19) World Intellectual Property Organization
International Bureau(43) International Publication Date
10 April 2003 (10.04.2003)

PCT

(10) International Publication Number
WO 03/030279 A3

(51) International Patent Classification⁷: H01M 6/16, 10/40

(21) International Application Number: PCT/US02/31048

(22) International Filing Date: 30 September 2002 (30.09.2002)

(25) Filing Language: English

(26) Publication Language: English

(30) Priority Data: 09/968,112 1 October 2001 (01.10.2001) US

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(81) Designated States (national): AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZM, ZW.

(84) Designated States (regional): ARIPO patent (GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, SK, TR), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG).

Published:
— with international search report

(88) Date of publication of the international search report: 3 July 2003

For two-letter codes and other abbreviations, refer to the "Guidance Notes on Codes and Abbreviations" appearing at the beginning of each regular issue of the PCT Gazette.

WO 03/030279 A3

(54) Title: DIRECT ADDITION OF BETA-AMINOENONES IN ORGANIC ELECTROLYTES OF NONAQUEOUS CELLS EMPLOYING SOLID CATHODES

(57) Abstract: A method for manufacturing a nonaqueous cell employing an anode, a cathode and an organic electrolyte wherein the cathode comprises an active cathode material, and the organic electrolyte contains a beta-aminoenone in a range of 0.1 to 5.0 volume percent based on the volume of the electrolyte solvent so as to aid in reducing the undesirably high initial open circuit voltage normally observed with the use of cathodes such as FeS₂.

DIRECT ADDITION OF BETA-AMINOENONES IN ORGANIC ELECTROLYTES OF NONAQUEOUS CELLS EMPLOYING SOLID CATHODES

DESCRIPTION

Technical Field

The invention relates to a method for manufacturing a nonaqueous cell employing an anode, a cathode comprising an active cathode material and an organic electrolyte. The method involves the direct addition of an additive such as beta-aminoenone, preferably 4-amino-3-penten-2-one, to the electrolyte to aid in reducing any undesirably high initial open circuit voltage such as that normally observed with cathodes such as FeS_2 .

Background Art

The development of high energy cell systems requires the compatibility of an electrolyte possessing desirable electrochemical properties with highly active anode materials, such as lithium, calcium, sodium and the like, and the efficient use of high energy density cathode materials. Such suitable high energy density cathode materials includes iron sulfides such as FeS_2 and FeS , carbon flourides such as CF_x , metal oxides such as V_2O_5 , WO_3 , MoO_3 , lead oxides such as Pb_3O_4 , PbO_2 and PbO , cobalt oxides such as Co_3O_4 , manganese oxides such as MnO_2 , In_2S_3 , NiS , metal chromates such as Ag_2CrO_4 , metal phosphates such as Ag_3PO_4 , LiCoO_2 , LiMn_2O_4 , Bi_2O_3 , CuO and Cu_2O and metal sulfates such as CuSO_4 . The use of aqueous electrolytes is precluded in these systems since the anode materials are sufficiently active to react with water chemically. Therefore, in order to realize the high energy density obtainable through use of these highly reactive anodes and high energy density cathodes, it is necessary to use a nonaqueous electrolyte system.

Many cell or battery applications, particularly in transistorized devices, such as hearing aids, watches, calculators, and the like, require a substantially unipotential discharge source for proper operation. However, it has been found that in many nonaqueous cells employing positive active materials, which include conductive additives such as graphite and/or carbon, the cell upon initially being discharged, exhibits a high voltage whereupon the cell then proceeds to reach its lower operating discharge voltage level only after a certain time period has elapsed. The time period for the cell to reach its intended operating discharge voltage level usually depends on the discharge rate through the load and thus, depending on the apparatus it is to power, could result in a period extending up to several hours or even days. This phenomenon has serious drawbacks when a cell is intended to be used in electronic devices requiring a substantially unipotential discharge source for proper operation. In some of these electronic devices, any initial voltage peak substantially exceeding the intended operating voltage for the device could result in serious damage to the electronic components of the device.

One approach to protect devices from batteries exhibiting high voltages prior to leveling off to their desired operating voltage level is to add additional electronic circuit components to protect the main operating components of the device. However, this not only adds to the expense of the device but also would result in enlarging the device to accommodate the protective circuitry. Larger devices run counter to the current emphasis placed on miniaturization, where it has become necessary for the battery industry to design smaller and smaller electrochemically powered cells.

Another approach designed to reduce an initially high and potentially disadvantageous voltage is to subject such cells to a pre-discharge regimen prior to initial use by an end user. But such pre-discharge regimens are time consuming and costly, and

reduce the capacity of the cell. Therefore, it is desirable to reduce or eliminate any required pre-discharge.

It has been suggested in U.S. Pat. No. 4489144 that the addition of an isoxazole derivative, such as 3,5-dimethylisoxazole, to the electrolyte may compensate for or eliminate this initial high voltage during discharge of nonaqueous cells. It was believed that the isoxazole derivative reacted with unwanted species that presumably caused the initial high voltage by reducing these species. The entire disclosure of U.S. Pat. No. 4489144 may be referred to for further details.

It has now been determined that a favorable reduction in initial open circuit voltage in a nonaqueous cell can be achieved by the addition of an additive as defined below to the electrolyte. Accordingly, it is an aspect of the within invention to provide a method for manufacturing a cell comprising a highly active anode such as lithium, a cathode comprising an active material such as FeS₂, and a nonaqueous electrolyte comprising an additive, such as 4-amino-3-penten-2-one.

In a broad aspect the invention comprehends a method for manufacturing a nonaqueous cell comprising the steps of providing an anode, an organic electrolyte solution comprising a solute dissolved in an organic solvent, and a solid cathode, and adding an additive to the electrolyte solution, the additive added to the electrolyte solution in a range of 0.1 to 5.0 volume percent based on the volume of the organic solvent and defined as any one of:



where R is hydrogen or an alkyl group; or



where R, R₂ and R₃ are hydrogen or an alkyl group; or related azirenes of the formula



where R, R', R'', R² and R³ are hydrogen or an alkyl group.

Detailed Description Of The Invention

The addition of the additive, such as 4-amino-3-penten-2-one, to nonaqueous organic electrolytes is effective in reducing and stabilizing the undesirably high initial open circuit voltages observed with cathode materials such as FeS₂. As used herein, the term "additive" will be used to describe a material containing any of the above structures.

A preferred additive is a beta-aminoenone such as 4-amino-3-penten-2-one.

The additive is preferably employed in the concentration range of between 0.1 and 5.0 volume percent based on the volume of the electrolyte solvent, preferably between 0.2 and 4.0 volume percent, and still more preferably between 0.2 and 2.0 volume percent, for most nonaqueous electrochemical couples.

The effectiveness of such additive in controlling initially high open circuit voltage was first discovered while examining the electrolyte solution of nonaqueous cells employing carbon and/or graphite-containing FeS₂ cathodes and lithium anodes. These cells were manufactured with an electrolyte solvent that included 3,5-dimethylisoxazole (DMI), an isoxazole derivative, as described in U.S. Pat. No. 4489144. Upon aging, gas chromatography (GC) studies on the electrolyte from the undischarged cells revealed that the DMI was no longer detectable. Instead, an amine was detected which was not initially present in the electrolyte. The amine was identified as 4-amino-3-penten-2-one (AP), one member of a class of beta-aminoenones. A second compound was also detected which was not initially present in the electrolyte: 2,4-pentanedione (PD). It is believed that the presence of PD is at least in part an artifact produced by exposure of the electrolyte to water during the GC sample preparation.

The reaction mechanism involving DMI and AP was further investigated by GC analysis of electrolytes from vial storage studies. In these studies, various components of a Li/FeS₂ cell were isolated and combined with electrolyte containing DMI. The procedure in general was as follows: A 3/4 inch square piece of cell component sample was placed in a 1 ounce jar. An electrolyte was prepared containing 1 mole of LiCF₃SO₃ added to 1 liter of 25:75:0.2 DIOX:DME:DMI by volume, where DIOX is 1,3-dioxolane and DME is 1,2-dimethoxyethane. One hundred micro liters of electrolyte was added to a vial. The vial was then capped and stored in the dry box for three hours. At the end of this period, the electrolyte was recovered in methylene chloride, the solution was extracted with water and the methylene chloride layer was analyzed.

Initially, electrolyte reaction with the lithium, separator material and cathode stock was examined. The results are presented in Table 1.

TABLE 1:

REACTANTS	PERCENT OR PPM BY GC PEAK AREA			
	% DIOX	% DME	% DMI	ppm AP
Electrolyte + lithium foil	23.42	76.14	.44	ND*
Electrolyte + separator	23.66	75.91	.44	ND*
Electrolyte + cathode stock	21.97	77.87	.09	614
Electrolyte	25.48	74.14	.38	ND*

*None detected.

The above data indicate that DMI is degraded in a reaction involving the cathode. To further isolate the point of DMI degradation, cathode mix was isolated from the aluminum foil substrate and the vial test was run on the cathode mix and this aluminum separately. The mix and the aluminum substrate were recombined and another vial test was run. A

Further vial test was run on cathode mix and fresh aluminum. The results are presented in Table 2.

TABLE 2:

REACTANTS	PERCENT OR PPM BY GC PEAK AREA				
	% DIOX	% DME	% DMI	ppm AP	ppm PD
Electrolyte + cathode mix	23.88	75.52	.44	ND*	ND*
Electrolyte + alum. substrate	24.04	75.46	.43	ND*	ND*
Electrolyte + cathode mix + alum. substrate	23.61	76.06	.25	362	534
Electrolyte + cathode mix + fresh aluminum	24.18	75.45	.31	47	477

*None detected.

In a further attempt to isolate the reactive species in the degradation of DMI and the generation of AP, cathode mix components were individually subjected to the vial test in combination with fresh substrate aluminum. These results are presented in Table 3.

TABLE 3:

REACTANTS	PERCENT OR PPM BY GC PEAK AREA				
	% DIOX	% DME	% DMI	ppm AP	ppm PD
Electrolyte + FeS2 (undried)	20.27	78.90	.42	ND*	ND*
Electrolyte + Acetylene black	25.24	74.38	.38	ND*	ND*
Electrolyte + polyethylene /polypropylene copolymer	25.32	74.27	.41	ND*	ND*
Electrolyte + polyethylene oxide	25.13	74.47	.38	ND*	ND*

*None detected.

The above data indicates that the DMI degradation reaction to produce AP only takes place in the presence of both the cathode mix and an aluminum substrate current carrier, and suggests that the degradation reaction involves a possible interaction of the cathode mix with the aluminum cathode current carrier.

Further experiments were carried out to determine whether the direct addition of AP or PD into the electrolyte solvent in place of DMI would show similar advantageous results in reducing the open circuit voltage of an undischarged cell. Several lots of nonaqueous AA size lithium anode iron pyrite cathode cells were constructed using the electrolyte described above with the following variations: a) no DMI, b) 0.2% DMI, c) 0.2% PD (instead of DMI) and d) 0.2% AP (instead of DMI), all by volume percent of the electrolyte solvent. The pyrite for these cells came from different production batches as indicated in the attached tables but was essentially the same material. The cells were stored either at ambient temperature (21°C) or at 60°C. The cells were removed from storage at the indicated period in the tables below and the open circuit voltage for the cell was measured. The cells were then autopsied and GC analysis was performed on the electrolyte. The results for the ambient cells are presented in Table 4 and for the 60°C cells in Table 5. Significantly, none of these cells contained any detectable amounts of DMI even after only two weeks at ambient temperature, indicating that the degradation reaction is completed relatively quickly, even at ambient storage temperatures.

TABLE 4: STORAGE AT 21°C

Electrolyte description/ storage duration	Pyrite lot #	OCV prior to autopsy (Volts)	ppm AP (by GC peak area)	ppm PD (by GC peak area)
No DMI	171			
		1.727	ND*	ND*
		1.757	ND*	ND*
		1.785	ND*	ND*
No DMI	172			

Electrolyte description/ storage duration	Pyrite lot #	OCV prior to autopsy (Volts)	ppm AP (by GC peak area)	ppm PD (by GC peak area)
2 weeks		1.732	ND*	ND*
10 weeks		1.760	ND*	ND*
16 weeks		1.775	ND*	ND*
0.2% DMI	171			
2 weeks		1.725	1441	ND*
7 weeks		1.748	334	ND*
23 weeks		1.748	ND*	ND*
0.2%DMI	172			
2 weeks		1.726	1080	ND*
10 weeks		1.730	51	ND*
16 weeks		1.745	32	ND*
0.2%PD	171			
2 weeks		1.725	ND*	39
7 weeks		1.758	ND*	ND*
23 weeks		1.780	ND*	ND*
0.2%AP (supplied by Lancaster Synthesis)	172			
2 weeks		1.727	1379	ND*
10 weeks		1.730	ND*	ND*
16 weeks		1.740	304	ND*
0.2%AP (supplied by TCI)	172			
2 weeks		1.727	2215	ND*
10 weeks		1.740	192	ND*
16 weeks		1.740	ND*	ND*

*None detected.

TABLE 5: STORAGE AT 60°C

Electrolyte description/ storage duration	Pyrite lot #	OCV PRIOR TO AUTOPSY (Volts)	ppm AP (by GC peak area)	ppm PD (by GC peak area)
No DMI				
4 weeks	171	1.846	ND*	ND*
12 weeks		1.847	ND*	ND*
No DMI	172			
4 weeks		1.830	ND*	ND*
12 weeks		1.830	ND*	ND*
.2% DMI	171			
4 weeks		1.798	ND*	ND*
12 weeks		1.837	ND*	ND*
.2%DMI	172			
4 weeks		1.810	ND*	ND*
12 weeks		1.824	ND*	ND*
.2%PD	171			

Electrolyte description/ storage duration	Pyrite lot #	OCV PRIOR TO AUTOPSY (Volts)	ppm AP (by GC peak area)	ppm PD (by GC peak area)
4 weeks		1.840	ND*	32
12 weeks		1.834	ND*	ND*
.2%AP (supplied by Lancaster Synthesis)	172			
4 weeks		1.810	ND*	ND*
12 weeks		1.750	ND*	ND*
.2%AP (supplied by TCI)	172			
4 weeks		1.760	267	ND*
12 weeks		1.790	ND*	ND*

*None detected

A study of the open circuit voltages for the cells in Table 4 reveals the following: at 16 weeks ambient, the cells with DMI in the electrolyte have a 30 mV lower open circuit voltage than the cells without DMI, confirming the observation disclosed in US Pat. No. 4489144 that DMI aids in reducing undesirably high open circuit voltage in such cells. The cells containing AP have a 35 mV lower open circuit voltage than the cells with no DMI. On the other hand, at 23 weeks ambient there is a 37 mV difference between the DMI cells and the no added DMI cells, but only a 5 mV difference between the PD cells and the cells without DMI. In general, it can be concluded that the cells with AP added to the electrolyte behave in the same manner as do cells with added DMI in reducing open circuit voltage, while the cells with PD added to the electrolyte behave in the same manner as do cells without any DMI added to the electrolyte. The cells represented in Table 5 demonstrate that the addition of AP to the electrolyte reduces open circuit voltage.

Literature studies show that AP can be obtained by the partial reduction of DMI. However, it is unexpected that DMI would be reduced in a reaction with the cathode, in view of the presence of an even stronger reducing agent, the lithium metal anode. However, when DMI is reduced with butyl lithium (often used to mimic the reducing power of lithium

metal), the reaction product is 3-methyl-5-hydroxymethyl isoxazole, a compound that is not a beta-aminoenone as defined herein. The fact that this reaction product is not seen in the studies represented in Tables 1 through 5, together with the fact that AP does react readily with lithium metal, further supports the proposition that DMI is reduced in a reaction with the cathode to lower the cathode voltage and the resulting AP is consumed by the lithium anode. Based on this information, it would not be expected that AP would react with the cathode to lower its voltage. Lowering the cathode voltage presumably involves reducing the cathode. Thus, the more reducing power the additive has the more likely it is to lower the cathode voltage. AP, on the other hand, is a reduced form of DMI in that AP has two additional hydrogen atoms. AP, being already partly reduced, has less reducing power than DMI. Consequently, one would not have expected AP to be as effective as DMI or even effective at all in lowering the cathode voltage. However, the results in Tables 1 through 5 demonstrate that AP behaves in the same manner as does DMI and does lower the cathode voltage.

The amount of the additive below 0.1 volume percent based on the volume of the electrolyte solvent would likely not provide sufficient reducing material to quickly and effectively reduce impurities and/or unwanted active species in the cell system. An amount above 5.0 volume percent based on the volume of the electrolyte solvent may provide excessive reducing material that could cause deleterious effects on other desired aspects of the cell. Although one embodiment of the invention, if focusing solely on voltage reduction, would include the use in the cathode of a metallic reducing agent such as zinc in conjunction with the additive in the electrolyte, the additive can be used without the metallic reducing agent in the cell, without departing from the scope of the invention. The omission of such a reducing agent may in fact be preferable, since

such reducing agents may produce other undesirable features within the cell.

If the within invention is practiced in conjunction with the addition of a metallic reducing agent, the metallic reducing agent can either be employed in a mixture in the forming of a cathode or it could be placed in contact with the cathode. Thus, any discrete material such as a layer, a coating, a metallic screen, a metallic strip, a porous metallic strip, or a powder and could be employed as long as it is in electronic and ionic contact with the cathode. The choice of a metallic reducing agent will depend upon the voltage of the cell's active cathode material with reference to the anode of the cell. For example, while zinc may be satisfactory for a nonaqueous lithium/FeS₂ system, it may not be suitable for a nonaqueous lithium/Ag₂O system since, in the latter, a less anodic (negative) metallic reducing agent may be required, such as tin or lead. Thus, once the EMF series is determined for a specific nonaqueous electrolyte and anode/cathode system, one can select which metallic reducing agents can be used in conjunction with the additive.

In the nonaqueous cell systems of this invention, suitable active cathode materials would include iron sulfides such as FeS₂ and FeS, carbon flourides such as CF_x, metal oxides such as V₂O₅, WO₃, MoO₃, lead oxides such as Pb₃O₄, PbO₂ and PbO, cobalt oxides such as Co₃O₄, manganese oxides such as MnO₂, In₂S₃, NiS, metal chromates such as Ag₂CrO₄, metal phosphates such as Ag₃PO₄, LiCoO₂, LiMn₂O₄, Bi₂O₃, CuO and Cu₂O and metal sulfates such as CuSO₄. Highly active anodes for use in nonaqueous systems according to this invention would be consumable metals and include aluminum, the alkali metals, alkaline earth metals and alloys of alkali metals or alkaline earth metals with each other and other metals. The term "alloys" as used herein is intended to include mixtures, solid solutions, such as lithium-magnesium, and the intermetallic compounds, such as

lithium monoaluminide. The preferred anode materials are lithium, sodium, potassium, calcium, magnesium and alloys thereof. Carbon anodes such as are used in lithium ion cells are also suitable anodes for use with the present invention. A preferred anode material to be used in conjunction with an FeS_2 cathode having an aluminum cathode substrate would be a lithium-aluminum alloy such as that disclosed in US Pat. No. 5514491, which may be referred to for further details. One such alloy is 0.5% aluminum by weight and is available from Chemetall-Foote.

In nonaqueous cells, useful organic solvents employed alone or mixed in the cell of this invention include the following classes of compounds: alkylene nitriles: e.g., crotonitrile (liquid ranges, -51.1 to 120°C); trialkyl borates: e.g., trimethyl borate, $(\text{CH}_3\text{O})_3\text{B}$ (liquid range, -29.3°C to 67°C); tetraalkyl silicates: e.g., tetramethyl silicate, $(\text{CH}_3\text{O})_4\text{Si}$ (boiling point, 121°C); nitroalkanes: e.g., nitromethane, CH_3NO_2 (liquid range, -17°C to 100.8°C); alkynitriles: e.g., acetonitrile, CH_3CN (liquid range, -45°C to 81.6°C); dialkylamides: e.g., dimethylformamide, $\text{HCON}(\text{CH}_3)_2$ (liquid range, -60.48°C to 149°C); lactams: e.g., N-methylpyrrolidone, (liquid range, -16°C to 202°C); monocarboxylic acid esters: e.g., ethyl acetate (liquid range, -83.6°C to 77.06°C); orthoesters: e.g., trimethylorthoformate, $\text{HC}(\text{OCH}_3)_3$ (boiling point, 103°C); lactones: e.g., (gamma)butyrolactone, (liquid range, -42°C to 206°C); dialkyl carbonates: e.g., dimethyl-carbonate, $\text{OC}(\text{OCH}_3)_2$ (liquid range, 2°C to 90°C); alkylene carbonates: e.g., propylene carbonate (liquid range, -48°C to 242°C), ethylene carbonate and vinylene carbonate; monoethers: e.g., diethyl ether (liquid range, -116°C to 34.5°C); polyethers: e.g., 1,1-and 1,2-dimethoxyethane (liquid ranges, -113.2°C to 64.5°C and -58°C to 83°C, respectively); cyclic ethers: e.g., tetrahydrofuran (liquid range, -65°C to 67°C), 1,3-dioxolane (liquid range, -95°C to 78°C) and substituted dioxolanes; nitroaromatics: e.g., nitrobenzene (liquid

range, 5.7°C to 210.8°C); cyclic sulfones: e.g., sulfolane, (melting point, 22°C), 3-methylsulfolane (melting point, -1°C); saturated heterocyclics: e.g., tetrahydrothiophene (liquid range, -96°C to 121°C); 3-methyl-2-oxazolidone (melting point, 15.9°C); five-membered unsaturated heterocyclics: e.g. 1-methylpyrrole (boiling point, 114°C), 2,4-dimethylthiazole (boiling point, 144°C), and furan (liquid range, -85.65°C to 31.36°C); dialkyl sulfoxides: e.g., dimethyl sulfoxide (liquid range, 18.4°C to 189°C); dialkyl sulfates: e.g., dimethylsulfate (liquid range, -31.75°C to 188.5°C); dialkyl sulfites: e.g., dimethylsulfite (boiling point, 126°C); alkylene sulfites: e.g., ethylene glycol sulfite (liquid range, -11°C to 173°C); halogenated alkanes: e.g., methylene chloride (liquid range, -95°C to 40°C); 1,3-dichloropropane (liquid range, -99.5°C to 120.4°C). Of the above, the preferred solvents are sulfolane; tetrahydrofuran; methyl-substituted tetrahydrofuran; 1,3-dioxolane; alkyl substituted 1,3-dioxolane; 3-methyl-2-oxazolidone; propylene or ethylene carbonate; (gamma)butyrolactone; ethylene glycol sulfite; dimethylsulfite; dimethyl sulfoxide; and 1,1-, 1,2-dimethoxyethane and glymes. Of the preferred solvents, the best are 3-methyl-2-oxazolidone, propylene or ethylene carbonate, 1,2-dimethoxyethane and 1,3-dioxolane because they appear more chemically inert to battery components and have wide liquid ranges, and especially because they permit highly efficient utilization of the cathode materials.

The ionizing solute for use in the invention may be a simple or double salt or mixtures thereof, e.g., LiCF_3SO_3 or LiClO_4 or LiI , which will produce an ionically conductive solution when dissolved in one or more solvents. Useful solutes include complexes of inorganic or organic Lewis acids and inorganic ionizable salts. The only requirements for utility are that the salts, whether simple or complex, be compatible with the solvent or solvents being employed and that they yield a solution which is sufficiently

ionically conductive. According to the Lewis or electronic concept of acids and bases, many substances, which contain no active hydrogen can act as acids or acceptors of electron doublets. The basic concept is set forth in the chemical literature (Journal of the Franklin Institute, Vol. 226, July/December 1938, pages 293-313 by G. N. Lewis). A suggested reaction mechanism for the manner in which these complexes function in a solvent is described in detail in U.S. Pat. No. 3,542,602 wherein it is suggested that the complex or double salt formed between the Lewis acid and the ionizable salt yields an entity which is more stable than either of the components alone. Typical Lewis acids suitable for use in the present invention include aluminum fluoride, aluminum bromide, aluminum chloride, antimony pentachloride, zirconium tetrachloride, phosphorus pentachloride, boron fluoride, boron chloride and boron bromide. Ionizable salts useful in combination with the Lewis acids include lithium fluoride, lithium chloride, lithium bromide, lithium sulfide, sodium fluoride, sodium chloride, lithium bromide, potassium fluoride, potassium chloride and potassium bromide.

The within invention can be utilized in the manufacture of both primary and secondary nonaqueous cells in a wide variety of cell constructions, including but not limited to cylindrical wound cells, cylindrical bobbin cells, miniature button cells, cells with one or more planar electrodes arranged in a co-facial orientation or in a co-planar orientation and prismatic cells. Further, although the experiments were performed by adding AP to the electrolyte, one of skill in the art will appreciate that the same result can be expected by the addition of AP to the cathode of the cell, without departing from the scope of the within invention.

WHAT IS CLAIMED IS:

1. A method for manufacturing a nonaqueous cell comprising the steps of providing an anode, an organic electrolyte solution comprising a solute dissolved in an organic solvent, and a solid cathode, and adding an additive to the electrolyte solution, the additive added to the electrolyte solution in a range of 0.1 to 5.0 volume percent based on the volume of the organic solvent, wherein the additive is defined as anyone of the following formulas:



where R is hydrogen or an alkyl group; or



where R, R₂ and R₃ are hydrogen or an alkyl group; or



where R, R', R'', R² and R³ are hydrogen or an alkyl group.

2. The method of claim 1, wherin the additive is in a range of from 0.2 to 4.0 volume percent based on the volume of the organic solvent.
3. The method of claim 1, wherein the additive is 4-amino-3-penten-2-one.
4. The method of claim 1, wherein the cathode comprises a material selected from the group consisting of iron sulfides, carbon flourides, V₂O₅, WO₃, MoO₃, Pb₃O₄, PbO₂, PbO, Co₃O₄, manganese oxides, In₂S₃, NiS, Ag₂CrO₄, Ag₃PO₄, LiCoO₂, LiMn₂O₄, Bi₂O₃, CuO, Cu₂O and CuSO₄.
5. The method of claim 1, wherein the anode comprises a material selected from the

group consisting of lithium, sodium, potassium, calcium, magnesium, carbon, alloys of lithium, alloys of sodium, alloys of potassium, alloys of calcium, alloys of magnesium and alloys of carbon.

6. The method of claim **4**, wherein the anode comprises a material selected from the group consisting of lithium, sodium, potassium, calcium, magnesium, carbon, alloys of lithium, alloys of sodium, alloys of potassium, alloys of calcium, alloys of magnesium and alloys of carbon.

7. The method of claim **6**, wherein the cathode comprises FeS₂ and the anode comprises a lithium-aluminum alloy.

8. The method of claim **7**, further comprising the step of adding a polyether to the electrolyte solution.

9. The method of claim **8**, wherein said polyether is 1,2-dimethoxyethane.

10. The method of claim **7**, further comprising the step of adding a cyclic ether to the electrolyte solution.

11. The method of claim **10**, wherein said cyclic ether is 1,3-dioxolane.