



(86) Date de dépôt PCT/PCT Filing Date: 2000/02/26

(87) Date publication PCT/PCT Publication Date: 2000/09/21

(45) Date de délivrance/Issue Date: 2010/08/03

(85) Entrée phase nationale/National Entry: 2001/11/19

(86) N° demande PCT/PCT Application No.: EP 2000/001611

(87) N° publication PCT/PCT Publication No.: 2000/055935

(30) Priorité/Priority: 1999/03/12 (DE199 10 968.0)

(51) Cl.Int./Int.Cl. *H01M 10/056* (2010.01)

(72) Inventeurs/Inventors:

HEIDER, UDO, DE;
SCHMIDT, MICHAEL, DE;
AMANN, ANJA, DE;
NIEMANN, MARLIES, DE;
KUHNER, ANDREAS, DE

(73) Propriétaire/Owner:

MERCK PATENT GESELLSCHAFT MIT
BESCHRAENKTER HAFTUNG, DE

(74) Agent: FETHERSTONHAUGH & CO.

(54) Titre : UTILISATION D'ADDITIFS DANS DES ELECTROLYTES POUR CELLULES ELECTROCHIMIQUES

(54) Title: USE OF ADDITIVES IN ELECTROLYTE FOR ELECTROCHEMICAL CELLS

(57) **Abrégé/Abstract:**

The invention relates to an electrolyte comprising a lithium-containing inorganic conductive salt or lithium-containing organic conductive salt, dissolved in an aprotic solvent, from the group of the methanides, triflates and imides and at least one organic salt of the formula (I) as additive $\text{Li}^+\text{B}^-(\text{OR}^1)_m(\text{OR}^2)_p$ (I), where the additive is present in a concentration between 0.1 and 10%, based on the total weight of conductive salt.

26474-585

Abstract

The invention relates to an electrolyte comprising a lithium-containing inorganic conductive salt or lithium-containing organic conductive salt, dissolved in an aprotic solvent, from the group of the methanides, triflates and imides and at least one organic salt of the formula (I) as additive



where the additive is present in a concentration between 0.1 and 10%, based on the total weight of conductive salt.

**Use of additives in electrolyte
for electrochemical cells**

The invention relates to the use of salt-based
5 compounds as additives in electrolytes for improving
the properties of electrochemical cells.

Lithium ion batteries are amongst the most promising
systems for mobile applications. The areas of
10 application extend from high-quality electronic
equipment (for example mobile telephones, camcorders)
to batteries for electrically driven vehicles.

These batteries consist of a cathode, an anode, a
15 separator and a non-aqueous electrolyte. The cathode is
typically $\text{Li}(\text{MnMe}_2)_2\text{O}_4$, $\text{Li}(\text{CoMe}_2)\text{O}_2$, $\text{Li}(\text{CoNi}_x\text{Me}_2)\text{O}_2$ or
other lithium intercalation and insertion compounds.
Anodes can consist of lithium metal, carbon, graphite,
graphitic carbon or other lithium intercalation and
20 insertion compounds or alloy compounds. The electrolyte
can be a solution containing lithium salts, such as
 LiPF_6 , LiBF_4 , LiClO_4 , LiAsF_6 , LiCF_3SO_3 , $\text{LiN}(\text{CF}_3\text{SO}_2)_2$ or
 $\text{LiC}(\text{CF}_3\text{SO}_2)_3$ and mixtures thereof, in aprotic solvents.

25 A multiplicity of additives for use in lithium ion
batteries is mentioned in the literature. For example,
in EP 0759641 and US 5776627, organic aromatic
compounds, such as biphenyl, substituted thiophenes and
furans, and in EP 0746050 and EP 0851524, substituted
30 anisole, mesitylene and xylene derivatives are added to
the electrolyte in order to increase the safety of the
battery in the case of overcharging. For the same
purpose, US 5753389 uses organic carbonates as
additives. In order to improve the cycle stability,
35 organic boroxines are added in EP 0856901. However, all
these additives have some crucial disadvantages.
Organic substances, as used in the specifications
mentioned here, generally have low flash points and low
explosion limits.

Additive	Explosion limit [%]	Flash point [°C]
Thiophene	1.5-12	-9
Anisole	0.34-6.3	43
Mesitylene	1-6	54
Furan	2.3-14.3	-35

Since the use of electrochemical cells and in particular the occurrence of faults (for example short-circuiting, mechanical damage) is always accompanied by warming, escape of the electrolyte represents an additional source of danger.

The object of the present invention is therefore to provide additives whose volatility is low and whose flash points are relatively high.

The object according to the invention is achieved by the use of organic alkali metal or tetraalkylammonium salts as additive.

The organic alkali metal salts are dissolved in electrolytes which are usually employed in non-aqueous secondary lithium batteries.

It has been found that the additives participate in formation of the coating layer on the anode and cathode. The coating layer results in passivation of the electrodes and thus in an increase in the cyclability of the electrodes. Film formation on the cathode can in addition serve increase safety in the event of overcharging, since after release of a mechanical safety means, for example by a disconnecter, as described in US 5741606, the voltage is dissipated by "internal self-discharge".

The additives are distinguished by very high thermal decomposition points. A crucial advantage over the additives used hitherto is the formation of a glass-

26474-585

- 3 -

like, polymeric layer on thermal decomposition, which can be caused, for example, by a short-circuit.

The invention therefore relates to an electrolyte for non-aqueous secondary lithium batteries which improves the performance, such as, for example, the coating layer formation, cyclability, safety, conductivity and low-temperature behaviour, through the addition of specific additives.

According to another aspect of the present invention, there is provided electrolyte comprising a lithium-containing inorganic conductive salt or lithium-containing organic conductive salt, dissolved in an aprotic solvent, from the group of the methanides, triflates and imides and at least one organic salt of the formula (I) as additive



in which m and p denote 0, 1, 2, 3 or 4, where m+p=4 and R¹ and R² are identical or different, are optionally bonded directly to one another by a single or double bond, each, individually or together, have the meaning of an aromatic or aliphatic carboxylic or sulfonic acid, or each, individually or together, have the meaning of an aromatic ring from the group phenyl, naphthyl, anthra-cenyl or phenanthrenyl, which are unsubstituted or mono- to tetrasubstituted by A or Hal, or each, individually or together, have the meaning of a heterocyclic aromatic ring from the group pyridyl, pyrazyl or bipyridyl, which are unsubstituted or mono- to trisubstituted by A or Hal, or each, individually or together, have the meaning of an aromatic hydroxy acid from the group of aromatic hydroxycarboxylic acids or aromatic hydroxysulfonic acids, which are unsubstituted or mono- to

26474-585

- 3a -

tetrasubstituted by A or Hal, and Hal denotes F, Cl or Br and A denotes alkyl having 1 to 6 C atoms, which is optionally mono- to trihalogenated, where the additive is present in a concentration between 0.1 and 10%, based on the
5 total weight of conductive salt.

Brief Description of the Drawings

Embodiments of the invention will be discussed with reference to the following Figures:

Fig. 1: Cyclic voltammogram of LiPF_6 in
10 EC/DMC (1:1) in a measurement cell containing a graphite anode, a lithium counter electrode and a lithium reference electrode.

Fig. 2: Cyclic voltammogram of LiPF_6
in EC/DMC (1:1) with 1% Lithium-
15 bis[salicylato(2-)O-,O']borate[1-] in a measurement cell containing a graphite anode, a lithium counter electrode and a lithium reference electrode.

Fig. 3: Cyclic voltammogram of LiPF_6
in EC/DMC (1:1) with 5% Lithium-
20 bis[salicylato(2-)O-,O']borate[1-] in a measurement cell containing a graphite anode, a lithium counter electrode and a lithium reference electrode.

Fig. 4: 5 Successive cyclic voltammograms of LiPF_6
in EC/DMC (1:1) with Lithium-
25 bis[salicylato(2-)O-,O']borate[1-] in a measurement cell containing a platinum working electrode, a lithium counter electrode and a lithium reference electrode.

Fig. 5: 5 Successive cyclic voltammograms of LiPF_6
in EC/DMC (1:1) with Lithium-
30 bis[salicylato(2-)O-,O']borate[1-] in a measurement cell

26474-585

- 3b -

containing a stainless-steel working electrode, a lithium counter electrode and a lithium reference electrode.

Fig. 6: Cyclic voltammogram (charging) of LiPF_6 in EC/DMC (1:1) with and without 1.5% Lithium-
5 bis[salicylato(2-)-O-,O']borate[1-] in a measurement cell containing a LiCoO_2 working electrode, a lithium counter electrode and a lithium reference electrode.

Fig. 7: Cyclic voltammogram (discharging) of LiPF_6 in EC/DMC (1:1) with and without 1.5% Lithium-
10 bis[salicylato(2-)-O-,O']borate[1-] in a measurement cell containing a LiCoO_2 working electrode, a lithium counter electrode and a lithium reference electrode.

Brief Description of the Invention

26474-585

- 3c -

Surprisingly, it has been found that lithium salts which actively participate in the formation of a passivating coating layer on the graphite electrode are suitable for improving the passivation of the anode. It
15 has been found that the quality of the coating layer is crucially improved. Reduction of the additive gives a film which is permeable to lithium ions on the anode. This film leads to improved cyclability of the anode from only the second cycle.

20

It has furthermore been found that these additives decompose oxidatively at potentials above the charge potential of the selected cathode and thus form a passivating film on the cathode. These films are
25 permeable to lithium ions and protect the selected solvent and conductive salt against oxidative decomposition.

Use in battery systems based on LiCoO_2 and LiNiO_2
30 appears particularly interesting. It is known that these electrode materials are unstable in the overcharged state. This can result in a vigorous reaction with the electrolyte, with corresponding safety risks occurring. The state of the art consists
35 of internal safety mechanisms, such as, for example, so-called disconnectors. On overcharging of a battery, gaseous components are generally liberated, with evolution of heat. The resultant pressure increase results in the disconnector breaking the contact

- 4 -

between electrode and current conductor and thus preventing further overcharging of the battery. A problem here is that the battery remains in the charged, unstable state.

5

External discharge is no longer possible owing to the irreversible breaking of contact.

10 The aim is, through addition of selected additives, to apply a film to the cathode in the event of overcharging, i.e. at potentials above the charge voltage, which film reacts with the cathode in a controlled manner after addressing of the disconnecter and thus dissipates the "excess potential" through
15 internal self-discharge.

A general example of the invention is explained in greater detail below.

20 a) Behaviour of the additives at low potentials

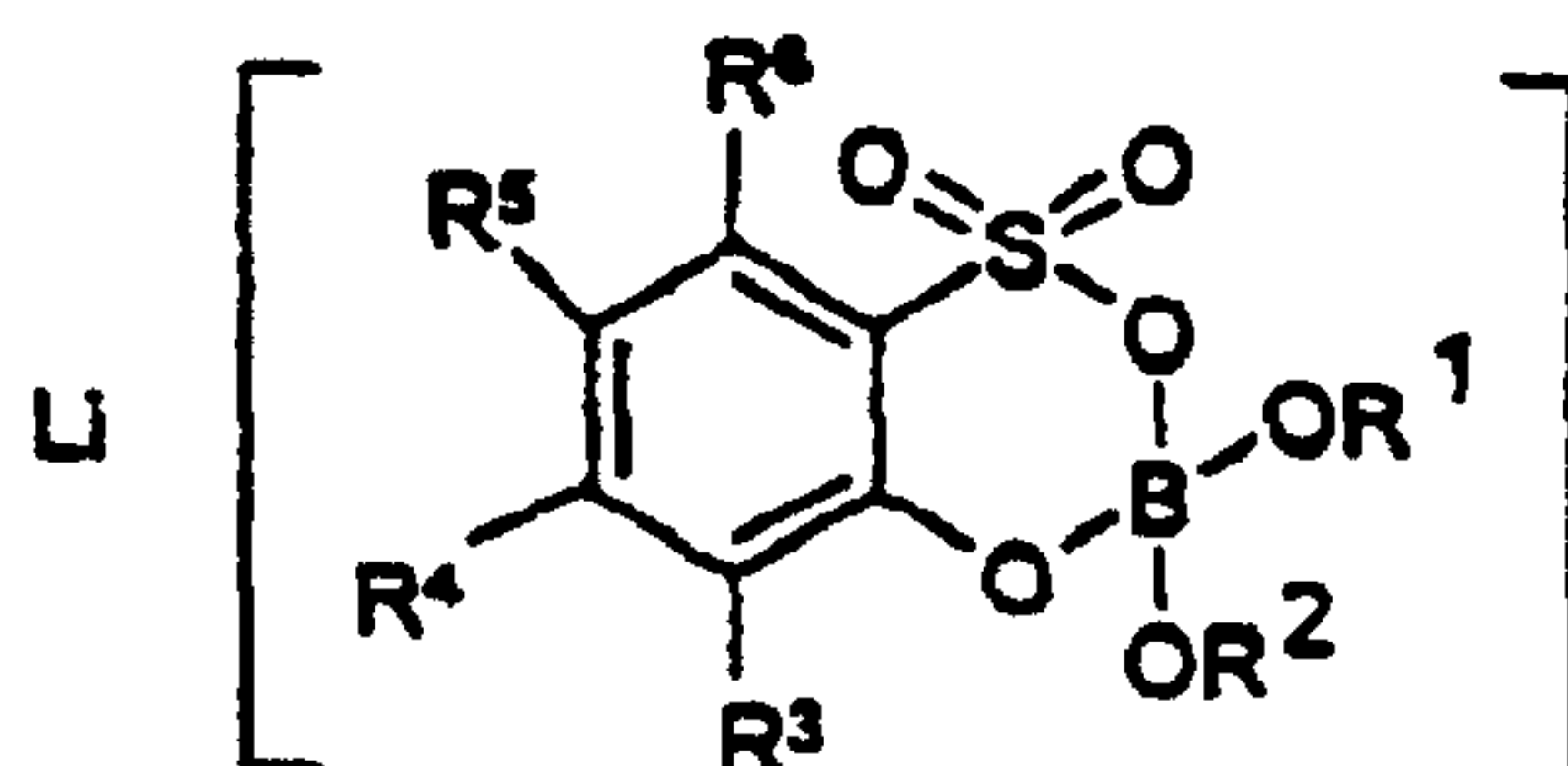
In each case, 3-5 cyclic voltammograms are recorded successively in a measurement cell containing an electrode of lithium metal, carbon, graphite, graphitic
25 carbon or other lithium intercalation and insertion compounds or alloy compounds, a lithium counterelectrode and a lithium reference electrode. Starting from the rest potential, the potential is lowered at a rate of 0.01 - 1 mV/s to 0 V against
30 Li/Li⁺ and then moved back to the rest potential.

The charge and discharge capacities Q_c and Q_d respectively are given by numerical integration of the $i(t)$ curves obtained. The cycling yield is obtained
35 from the quotient Q_c/Q_d .

Electrolytes which can be used are solutions of LiPF₆, LiBF₄, LiClO₄, LiAsF₆, LiCF₃SO₃, LiN(CF₃SO₂)₂ or LiC(CF₃SO₂)₃ and mixtures thereof, in aprotic solvents,

- 5 -

such as EC, DMC, PC, DEC, BC, VC, cyclopentanone, sulfolane, DMS, 3-methyl-1,3-oxazolidin-2-one, γ -butyrolactone, EMC, MPC, BMC, EPC, BEC, DPC, 1,2-diethoxymethane, THF, 2-methyltetrahydrofuran, 1,3-dioxolane, methyl acetate, ethyl acetate and mixtures thereof. The electrolytes can also comprise organic isocyanates (DE 199 44 603) for reducing the water content. Lithium complex salts of the formula



where

R^1 and R^2 are identical or different, are optionally directly bonded to one another via a single or double bond, and are each, individually or together, an aromatic ring from the group consisting of phenyl, naphthyl, anthracenyl and phenanthrenyl, which may be unsubstituted or mono- to hexasubstituted by alkyl (C_1 to C_6), alkoxy groups (C_1 to C_6) or halogen (F, Cl or Br),

or are each, individually or together, an aromatic heterocyclic ring from the group consisting of pyridyl, pyrazyl and pyrimidyl, which may be unsubstituted or mono- to tetrasubstituted by alkyl (C_1 to C_6), alkoxy groups (C_1 to C_6) or halogen (F, Cl or Br),

or are each, individually or together, an aromatic ring from the group consisting of hydroxybenzocarboxyl, hydroxynaphthalenecarboxyl, hydroxybenzosulfonyl and hydroxynaphthalenesulfonyl, which may be unsubstituted or mono- to tetrasubstituted by alkyl (C_1 to C_6), alkoxy groups (C_1 to C_6) or halogen (F, Cl or Br),

- 6 -

R^3 - R^6 can each, individually or in pairs and optionally bonded directly to one another by a single or double bond, have the following meanings:

- 5 1. alkyl (C_1 to C_6), alkoxy (C_1 to C_6) or halogen (F, Cl or Br)
2. an aromatic ring from the groups
- 10 phenyl, naphthyl, anthracenyl and phenanthrenyl, which may be unsubstituted or mono- to hexasubstituted by alkyl (C_1 to C_6), alkoxy groups (C_1 to C_6) or halogen (F, Cl or Br),
- 15 pyridyl, pyrazyl and pyrimidyl, which may be unsubstituted or mono- to tetrasubstituted by alkyl (C_1 to C_6), alkoxy groups (C_1 to C_6) or halogen (F, Cl or Br),
- 20 which are prepared by the following process (DE 199 32 317):
 - a) chlorosulfonic acid is added to 3-, 4-, 5- or 6-substituted phenol (III) in a suitable solvent,
 - 25 b) the intermediate (IV) from a) is reacted with chlorotrimethylsilane, and the product is filtered and subjected to fractional distillation,
 - 30 c) the intermediate (II) from b) is reacted with lithium tetramethoxyborate(1-) in a suitable solvent, and the end product (I) is isolated therefrom,
- 35 can also be present in the electrolyte.

The electrolytes can likewise comprise compounds of the following formula (DE 199 41 566):

- 7 -



where

5 Kt = N, P, As, Sb, S or Se,

A = N, P, P(O), O, S, S(O), SO₂, As, As(O), Sb or
Sb(O),

10 R¹, R² and R³

are identical or different

and are H, halogen, substituted and/or unsubstituted
15 alkyl C_nH_{2n+1}, substituted and/or unsubstituted alkenyl
having 1-18 carbon atoms and one or more double bonds,
substituted and/or unsubstituted alkynyl having 1-18
carbon atoms and one or more triple bonds, substituted
and/or unsubstituted cycloalkyl C_mH_{2m-1}, mono- or
20 polysubstituted and/or unsubstituted phenyl, substitu-
ted and/or unsubstituted heteroaryl,

A can be included in R¹, R² and/or R³ in various
positions,

25

Kt can be included in a cyclic or heterocyclic ring,

the groups bonded to Kt may be identical or different,

30 where

n = 1-18

m = 3-7

35

k = 0 or 1-6

l = 1 or 2 in the case where x = 1 and 1 in the case
where x = 0

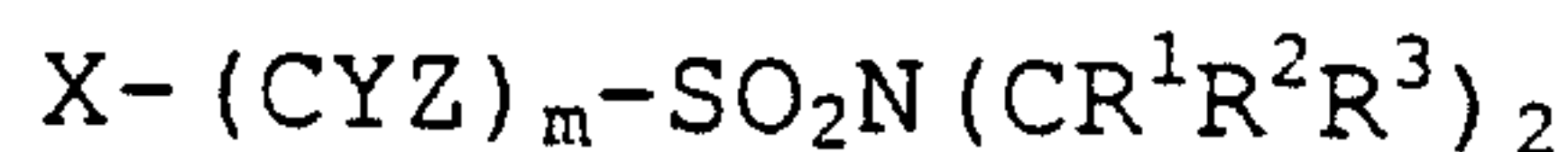
x = 0 or 1

y = 1-4.

5

However, use can also be made of electrolytes comprising compounds of the general formula (DE 199 53 638)

10



where

15

X is H, F, Cl, C_nF_{2n+1} , C_nF_{2n-1} or $(SO_2)_kN(CR^1R^2R^3)_2$,

Y is H, F or Cl

Z is H, F or Cl

20

R^1 , R^2 and R^3 are H and/or alkyl, fluoroalkyl or cycloalkyl

m is 0-9 and, if X = H, $m \neq 0$

25

n is 1-9

k is 0 if $m = 0$ and $k = 1$ if $m = 1-9$,

30

and complex salts of the general formula (DE 199 51 804)



in which

35

x and y are 1, 2, 3, 4, 5 or 6

M^{x+} is a metal ion

E is a Lewis acid selected from the group consisting of

$BR^1R^2R^3$, $AlR^1R^2R^3$, $PR^1R^2R^3R^4R^5$, $AsR^1R^2R^3R^4R^5$ and $VR^1R^2R^3R^4R^5$,

R^1 to R^5 are identical or different, are optionally bonded directly to one another by a single or double bond, and each, individually or together, has the following meaning:

a halogen (F, Cl or Br),

an alkyl or alkoxy radical (C_1 to C_8), which can be partially or fully substituted by F, Cl or Br,

an aromatic ring, optionally bonded via oxygen, from the group consisting of phenyl, naphthyl, anthracenyl and phenanthrenyl, which may be unsubstituted or mono- to hexasubstituted by alkyl (C_1 to C_8) or F, Cl or Br,

an aromatic heterocyclic ring, optionally bonded via oxygen, from the group consisting of pyridyl, pyrazyl and pyrimidyl, which may be unsubstituted or mono- to tetrasubstituted by alkyl (C_1 to C_8) or F, Cl or Br, and

Z is OR^6 , NR^6R^7 , $CR^6R^7R^8$, OSO_2R^6 , $N(SO_2R^6)(SO_2R^7)$, $C(SO_2R^6)(SO_2R^7)(SO_2R^8)$ or $OCOR^6$, where

R^6 to R^8 are identical or different, are optionally bonded directly to one another by a single or double bond and are each, individually or together,

hydrogen or are as defined for R^1 to R^5 .

These electrolytes can be employed in electrochemical cells containing cathodes made from customary lithium intercalation and insertion compounds, but also containing cathode materials consisting of lithium

- 10 -

mixed oxide particles coated with one or more metal oxides (DE 199 22 522) or polymers (DE 199 46 066).

0% for the control and from 0.1 to 10% (based on the total weight of conductive salt) of additives from the group consisting of organic alkali metal salts are added. Particular preference is given to additives from the group consisting of organic alkali metal borates and alkali metal alkoxides or tetraalkylammonium borates and alkoxides. From 0.1 to 7% of additives are preferably added to the conductive salt.

On evaluation of the measurement curves, it becomes clear that the additive decomposes by reduction at potentials of about 900 - 1000 mV against Li/Li⁺.

Through the reduction of the additive, more capacity, compared with conventional systems, is consumed in the 1st cycle. At the latest after the 3rd cycle, however, significantly higher cycling yields are obtained than without additive.

b) Behaviour of the additives at high potentials

In each case, 3-5 cyclic voltammograms were recorded successively in a measurement cell containing a stainless-steel, platinum or gold working electrode, a lithium counterelectrode and a lithium reference electrode. To this end, starting from the rest potential, the potential was firstly increased at a rate of from 1 mV/s to 100 mV/s to voltages above the respective decomposition potential of the corresponding additive against Li/Li⁺, and then moved back to the rest potential.

35

Depending on the oxidation potential, the additives are oxidized in the first cycle at potentials of from 3 V to 5 V against Li/Li⁺. However, this oxidation does not result in a lasting current increase, as in

- 11 -

conventional salts, such as LiPF_6 , Li imide or Li methanide, but, after passing through a maximum with relatively low currents, results in the formation of a passivating coating layer on the working electrode.

5 Electrolytes which can be used are solutions of LiPF_6 , LiBF_4 , LiClO_4 , LiAsF_6 , LiCF_3SO_3 , $\text{LiN}(\text{CF}_3\text{SO}_2)_2$ or $\text{LiC}(\text{CF}_3\text{SO}_2)_3$, and mixtures thereof, in aprotic solvents, such as EC, DMC, PC, DEC, BC, VC, cyclopentanone,
10 sulfolane, DMS, 3-methyl-1,3-oxazolidin-2-one, γ -butyrolactone, EMC, MPC, BMC, EPC, BEC, DPC, 1,2-diethoxymethane, THF, 2-methyltetrahydrofuran, 1,3-dioxolane, methyl acetate, ethyl acetate and mixtures thereof. 0% for the control and from 0.1 to 10% (based
15 on the total weight of conductive salt) of additives from the group consisting of organic alkali metal salts are added. Particular preference is given to additives from the group consisting of organic alkali metal borates and alkali metal alkoxides. From 0.1 to 7% of
20 additives are particularly preferably added to the conductive salt.

c) Properties of the coating layer formed by oxidation

25 In each case, 3-5 cyclic voltammograms are recorded successively in a measurement cell containing a stainless-steel working electrode, a lithium counterelectrode and a lithium reference electrode. Starting from the rest potential, the potential is
30 firstly increased at a rate of 10 mV/s - 20 mV/s to values above the respective decomposition potential of the corresponding additives. The coating layer is deposited on the electrode in the process. The potential is then lowered to values below 0 V against
35 Li/Li^+ , thus initiating lithium deposition on the stainless-steel electrode. To this end, lithium ions must migrate through the film formed. In order to exclude dissolution of the coating layer during this process, the potential is again increased to values

above the respective decomposition potential of the salts mentioned. Lithium cycling (evident from deposition and dissolution peaks at low potentials) is possible in the electrolyte. Furthermore, the coating layer is not dissolved by the selected process, since otherwise oxidation of the salt used at the above-mentioned potentials would have to be detectable in the second and all subsequent cycles.

10 d) Application of the coating layer to certain cathode materials

LiMn₂O₄, LiCoO₂, LiNiO₂ and LiNi_xCo_{1-x}O₂ cathodes are particularly interesting. In a measurement cell, a working electrode having one of the compositions indicated here, a lithium counterelectrode and a lithium reference electrode are used.

Electrolytes which can be used are solutions of LiPF₆, LiBF₄, LiClO₄, LiAsF₆, LiCF₃SO₃, LiN(CF₃SO₂)₂ or LiC(CF₃SO₂)₃, and mixtures thereof, in aprotic solvents, such as EC, DMC, PC, DEC, BC, VC, cyclopentanone, sulfolane, DMS, 3-methyl-1,3-oxazolidin-2-one, γ-butyrolactone, EMC, MPC, BMC, EPC, BEC, DPC, 1,2-diethoxymethane, THF, 2-methyltetrahydrofuran, 1,3-dioxolane, methyl acetate, ethyl acetate and mixtures thereof. 0% for the control and from 0.1 to 10% (based on the total weight of conductive salt) of additives from the group consisting of organic alkali metal salts are added. Particular preference is given to additives from the group consisting of organic alkali metal borates and alkali metal alkoxides. From 0.1 to 7% of additives are preferably added to the conductive salt.

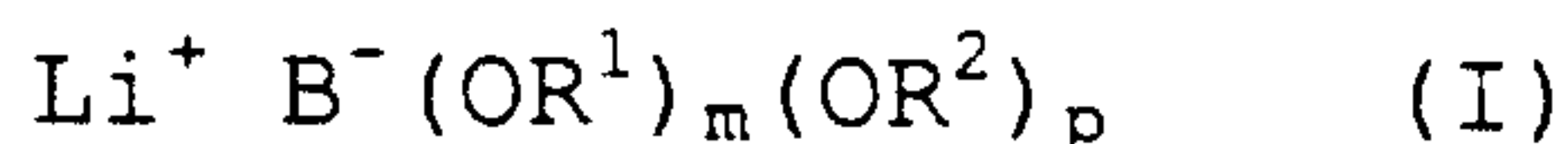
35 Starting from the rest potential, the cathodes are first fully charged against Li/Li⁺.

The cathode is subsequently overcharged. During this operation, the voltage reaches an upper value

- 13 -

determined by the measurement arrangement. If the potentiostat/galvanostat is then switched off, the potential drops very rapidly.

- 5 Compared with the reference, the increase in potential in the range 4.3 - 6 V against Li/Li⁺ is slower in the case of the additive-containing electrolytes according to the invention.
- 10 After the potentiostat/galvanostat has been switched off (release of the disconnecter is simulated), the potential of the cathode drops rapidly in both electrolytes.
- 15 In the case of the electrolyte without additive, the potential oscillates at values around 4.2 - 4.3 V against Li/Li⁺. Accordingly, the cathode remains in the charged, high-energy state.
- 20 By contrast, the addition of additives causes a lowering of the potential. The potential then corresponds to the rest potential of an uncharged electrode. This suggests that additives or the film formed by decomposition of the additive is capable of
- 25 dissipating the "excess potential" in a controlled manner by internal self-discharge and thus converting the battery into a low-energy state after release of the safety means (for example disconnecter).
- 30 Particularly suitable additives according to the invention are compounds of the following formula:



35

in which

m and p are 0, 1, 2, 3 or 4, where $m + p = 4$, and

- 14 -

R¹ and R² are identical or different,

are optionally bonded directly to one another by a single or double bond,

5

and are each, individually or together, an aromatic or aliphatic carboxylic or sulfonic acid, or

10 are each, individually or together, an aromatic ring from the group consisting of phenyl, naphthyl, anthracenyl and phenanthrenyl, which may be unsubstituted or mono- to tetrasubstituted by A or Hal, or,

15 are each, individually or together, a heterocyclic aromatic ring from the group consisting of pyridyl, pyrazyl and bipyridyl, which may be unsubstituted or mono- to trisubstituted by A or Hal, or

20 are each, individually or together, an aromatic hydroxy acid from the group consisting of aromatic hydroxycarboxylic acids and aromatic hydroxysulfonic acids, which may be unsubstituted or mono- to tetrasubstituted by A or Hal,

25

and

Hal is F, Cl or Br

30 and

A is alkyl having 1 to 6 carbon atoms, which may be mono- to trihalogenated.

35 Particularly suitable compounds are also those of the following formula:



in which R

is an aromatic or aliphatic carboxylic or sulfonic acid, or

5

is an aromatic ring from the group consisting of phenyl, naphthyl, anthracenyl and phenanthrenyl, which may be unsubstituted or mono- to tetrasubstituted by A or Hal, or,

10

is an aromatic heterocyclic ring from the group consisting of pyridyl, pyrazyl and bipyridyl, which may be unsubstituted or mono- to trisubstituted by A or Hal, or

15

is an aromatic hydroxy acid from the group consisting of aromatic hydroxycarboxylic acids and aromatic hydroxysulfonic acids, which may be unsubstituted or mono- to tetrasubstituted by A or Hal,

20

and

Hal is F, Cl or Br

25 and

A is alkyl having 1 to 6 carbon atoms, which may be mono- to trihalogenated.

30 Particularly preferred additives are lithium bis[1,2-benzenediolato(2-)O,O']borate(1-), lithium bis[3-fluoro-1,2-benzenediolato(2-)O,O']borate(1-), lithium bis[2,3-naphthalenediolato(2-)O,O']borate(1-), lithium bis[2,2'-biphenyldiolato(2-)O,O']borate(1-), lithium
35 bis[salicylato(2-)O,O']borate(1-), lithium bis[2-olato-benzenesulfonato(2-)O,O']borate(1-), lithium bis[5-fluoro-2-olatobenzenesulfonato(2-)O,O']borate, lithium phenoxide and lithium 2,2-biphenoxide.

- 16 -

Suitable additives according to the invention are also compounds of the formula (III), which exhibit similar properties to the compounds of the formulae (I) and (II):

5



where

10 w, x, y and z can be 0, 1, 2, 3 or 4, where $w + x + y + z = 4$, and

R'_w, R''_x, R'''_y and R''''_z are identical or different and are each alkyl having 1 to 8 carbon atoms which may in each case be mono- to trihalogenated, and

15 A⁻ is OR¹ or B(OR¹)_m(OR²)_p, in which m and p are 0, 1, 2, 3 or 4, where $m + p = 4$, and R¹ and R² are identical or different,

are optionally bonded directly to one another by a single or double bond,

20

and are each, individually or together, an aromatic or aliphatic carboxylic, dicarboxylic or sulfonic acid radical, or

25 are each, individually or together, an aromatic ring from the group consisting of phenyl, naphthyl, anthracenyl and phenanthrenyl, which may be unsubstituted or mono- to tetrasubstituted by A or Hal, or,

30 are each, individually or together, a heterocyclic aromatic ring from the group consisting of pyridyl, pyrazyl and bipyridyl, which may be unsubstituted or mono- to trisubstituted by A or Hal, or

35 are each, individually or together, an aromatic hydroxy acid from the group consisting of aromatic hydroxycarboxylic acids and aromatic hydroxysulfonic acids, which may be unsubstituted or mono- to tetrasubstituted by A or Hal,

and

Hal is F, Cl or Br

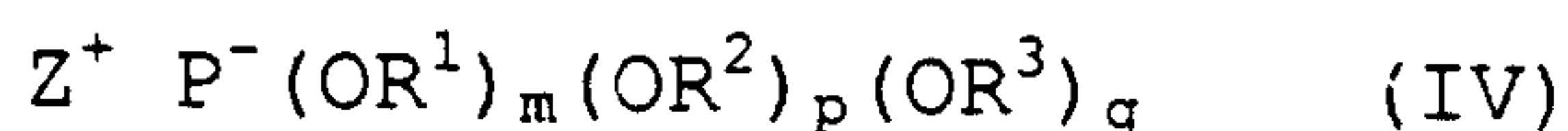
5

and

A is alkyl having 1 to 6 carbon atoms, which may in each case be mono- to trihalogenated.

10

Suitable additives according to the invention are also compounds of the following formula:



15

where

Z^+ is Li^+ or $[NR'_wR''_xR'''_yR''''_z]^+$,

where

20 w, x, y and z can be 0, 1, 2, 3 or 4, where $w + x + y + z = 4$, and

R'_w , R''_x , R'''_y and R''''_z are identical or different and are each alkyl having 1 to 8 carbon atoms which can in each case be mono- to trihalogenated, and

25 where

m, p and q are 0, 1, 2, 3, 4, 5 or 6, where $m + p + q = 6$, and

R^1 , R^2 and R^3 are identical or different,

30 are optionally bonded directly to one another by a single or double bond,

and are each, individually or together, an aromatic or aliphatic carboxylic, dicarboxylic or sulfonic acid radical, or

35

are each, individually or together, an aromatic ring from the group consisting of phenyl, naphthyl, anthracenyl and phenanthrenyl, which may be

- 18 -

unsubstituted or mono- to tetrasubstituted by A or Hal,
or,

are each, individually or together, a heterocyclic
5 aromatic ring from the group consisting of pyridyl,
pyrazyl and bipyridyl, which may be unsubstituted or
mono- to trisubstituted by A or Hal, or

are each, individually or together, an aromatic hydroxy
10 acid from the group consisting of aromatic
hydroxycarboxylic acids and aromatic hydroxysulfonic
acids, which may be unsubstituted or mono- to
tetrasubstituted by A or Hal,

15 and

Hal is F, Cl or Br

and

20

A is alkyl having 1 to 6 carbon atoms, which may be
mono- to trihalogenated.

The following examples are intended to illustrate the
25 invention in greater detail, but without representing a
limitation.

Examples

30 Example 1

In each case, 3 cyclic voltammograms were recorded
successively in a measurement cell containing a
graphite anode (SFG 44 with PVDF binder), a lithium
35 counterelectrode and a lithium reference electrode. To
this end, starting from the rest potential, the
potential was firstly lowered at a rate of 0.1 mV/s to
0 V against Li/Li⁺ and then moved back to the rest
potential.

The electrolytes used were solutions of LiPF_6 in EC/DMC (1:1) containing 0% (control), 1% and 5% (based on the weight of LiPF_6) of lithium bis[salicylato(2-)-O,O']borate(1-) (abbreviated to lithium salborate).

The results are shown in Table 1 and in Figures 1, 2 and 3.

Table 1: Cycling yields on graphite

Electrolyte	Yield 1st cycle	Yield 3rd cycle
1M LiPF_6 in EC/DMC (1:1)	71.7%	90.5%
1M LiPF_6 in EC/DMC (1:1) + 1% lithium bis[salicylato(2-)-O,O']-borate(1-)	69.5%	95.5%
1M LiPF_6 in EC/DMC (1:1) + 5% lithium bis[salicylato(2-)-O,O']-borate(1-)	61.3%	95.1%

It is clearly evident from Figures 1 and 2 that the additive decomposes just before the film formation by ethylene carbonate. The reduction potential can be given as about 900-1000 mV against Li/Li^+ .

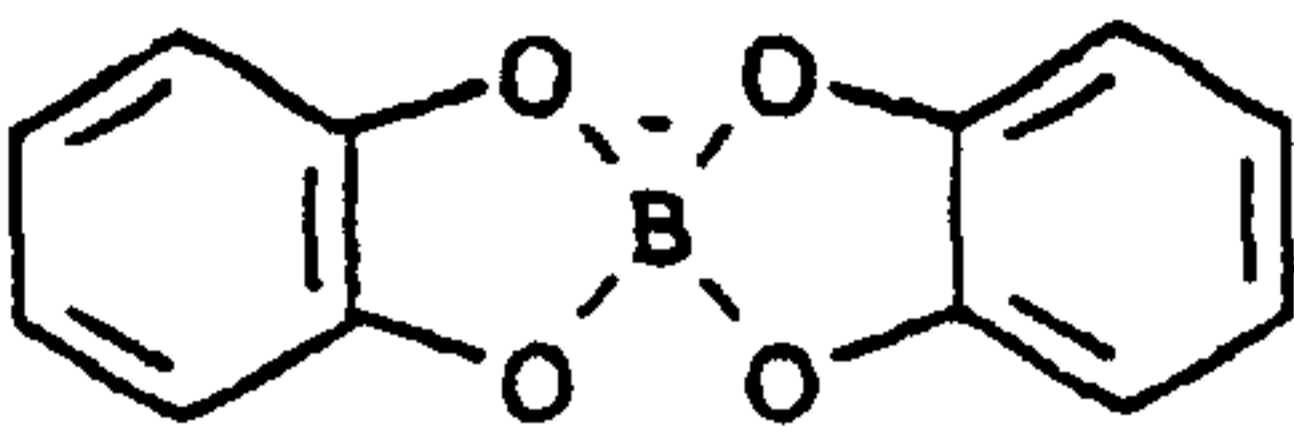
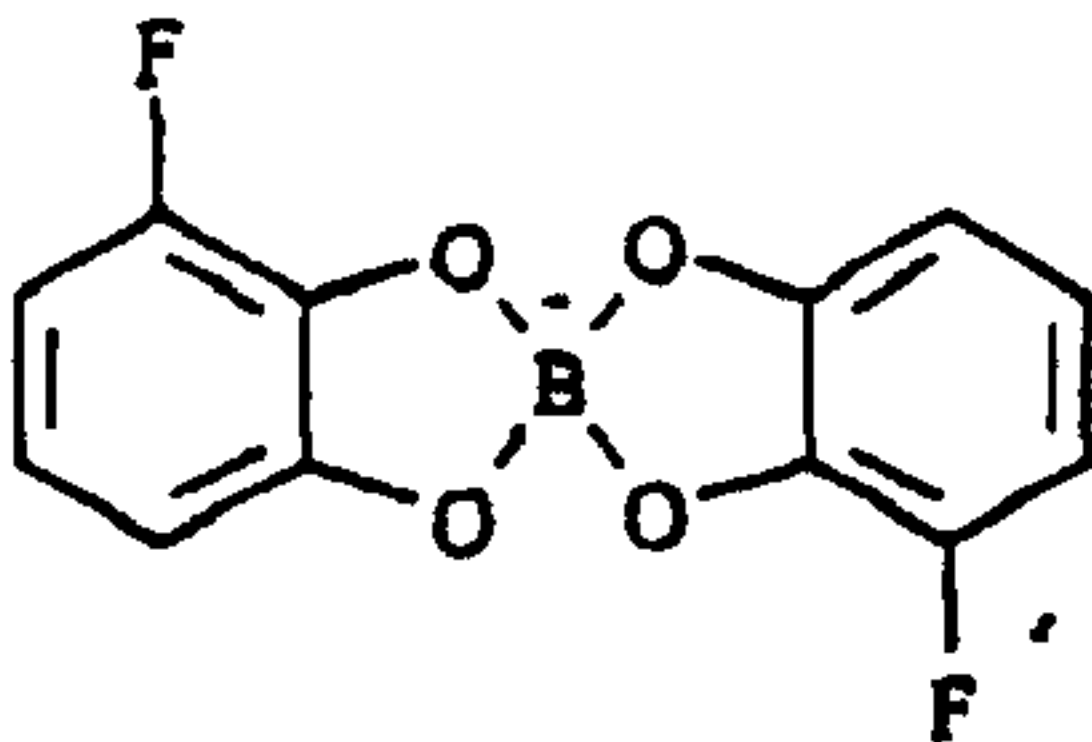
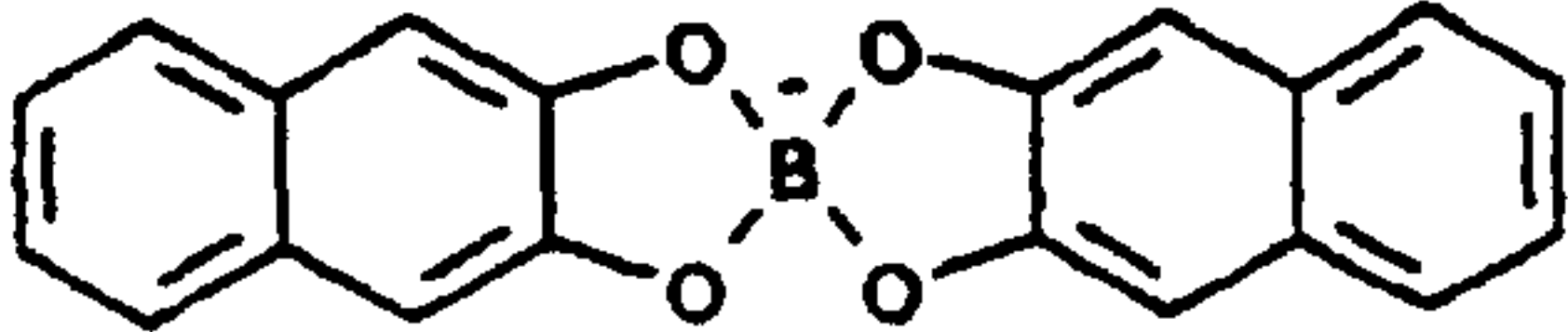
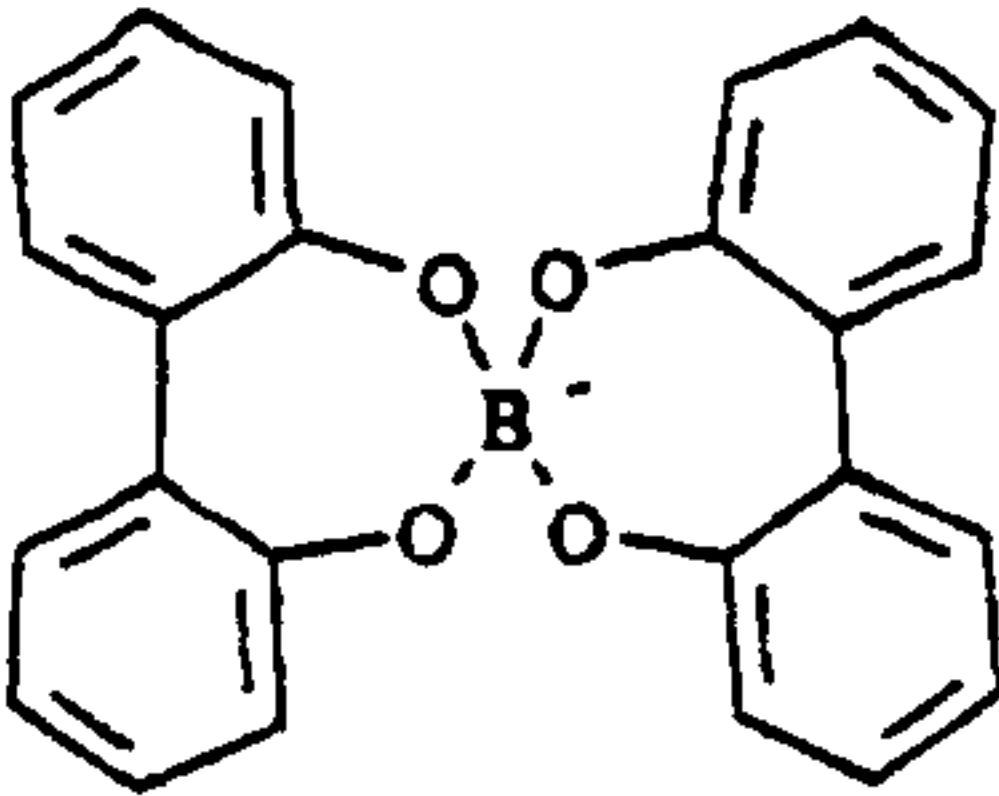
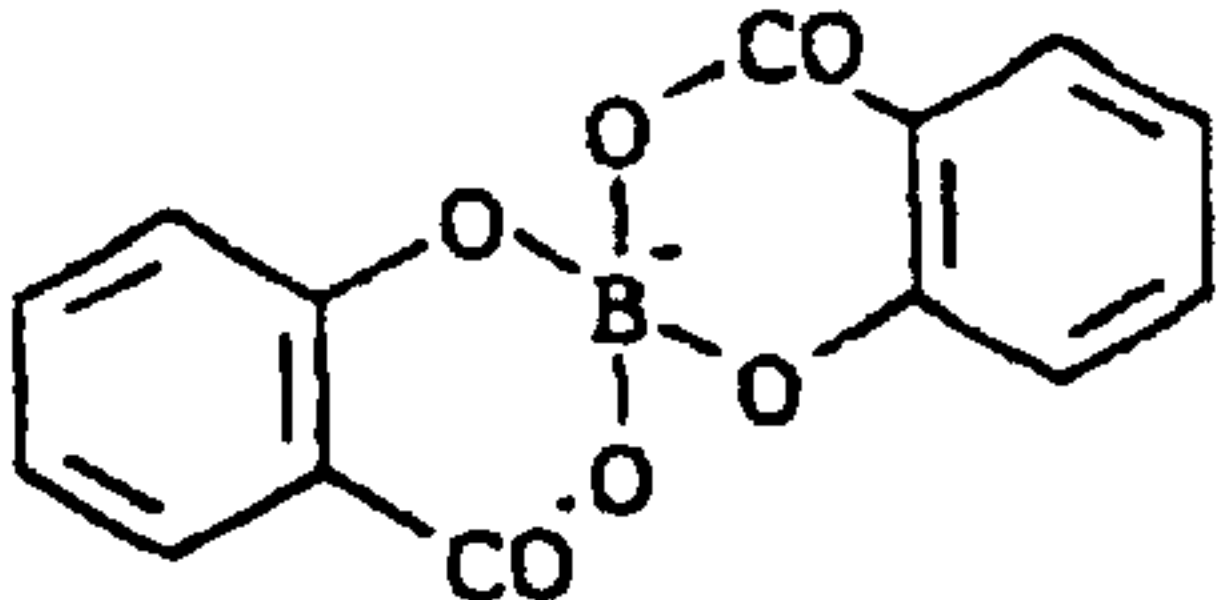
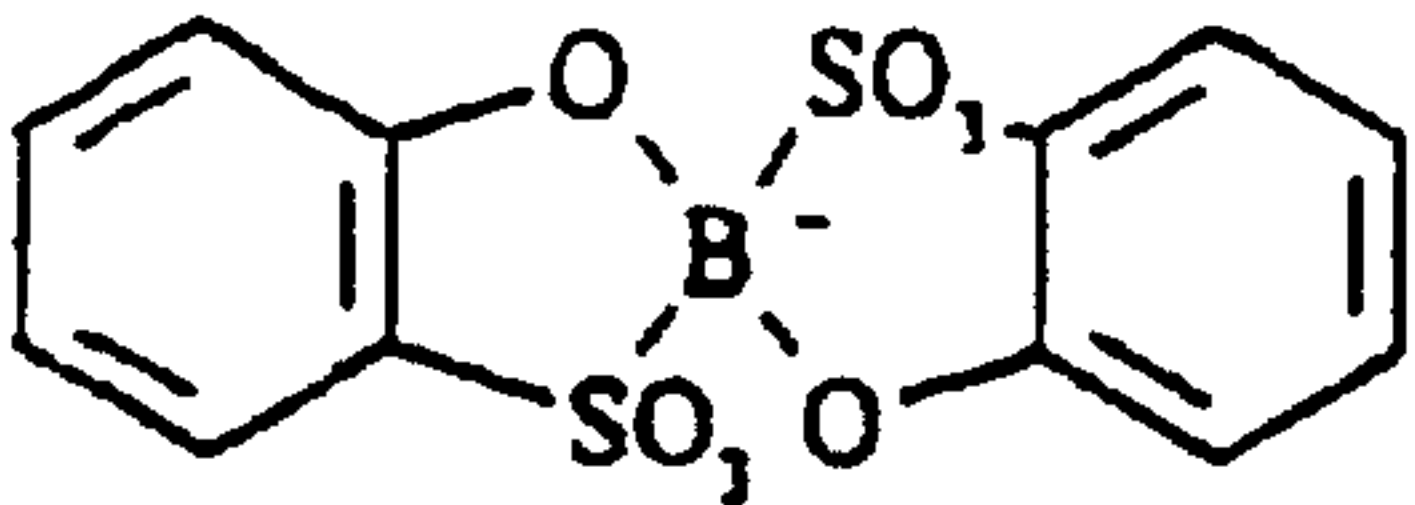
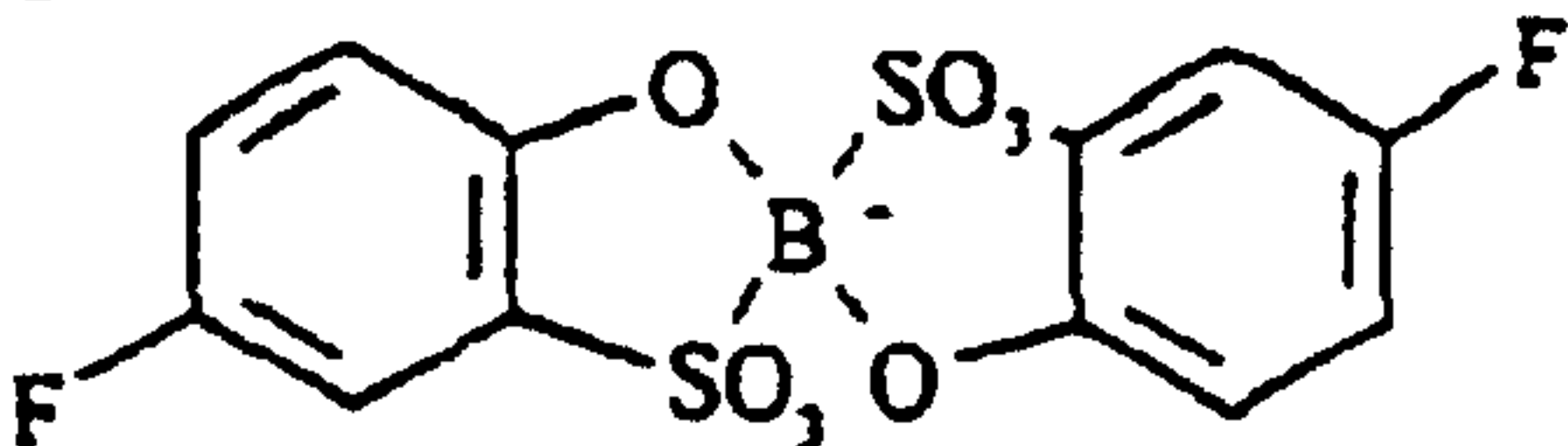
The reduction of the additive causes a somewhat greater consumption of capacity in the first cycle. This disadvantage is compensated for from the third cycle. Significantly higher cycling yields are obtained.

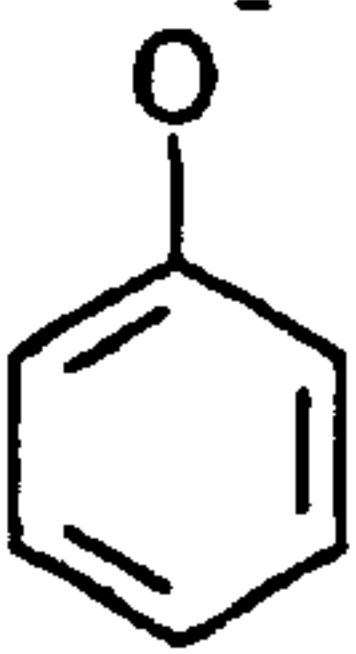
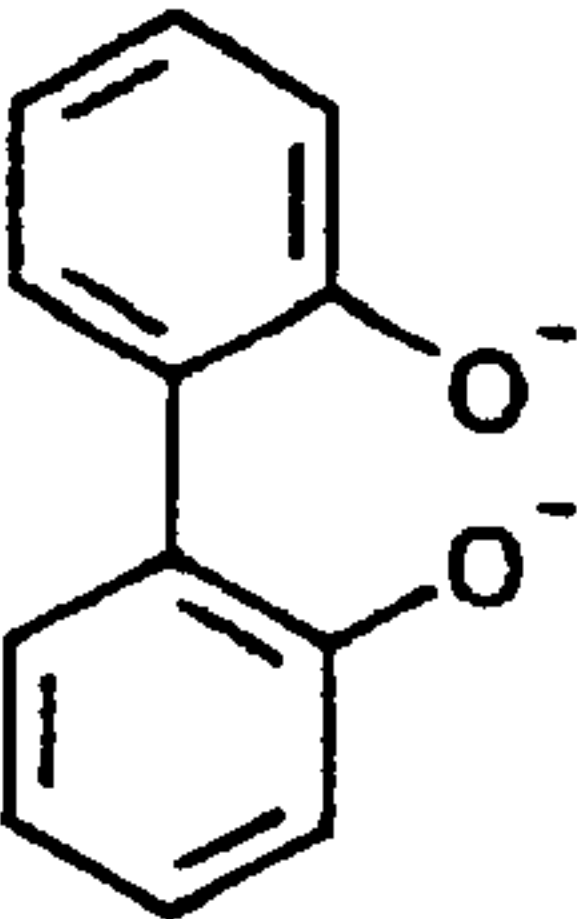
Example 2

Passivation of the cathode

Lithium salts have been found which participate actively in the build-up of a passivating coating layer on the cathode. The coating layer formed is permeable to lithium ions.

Table 2: Selected lithium salts

Anion	E_{ox} vs. Li/Li^+ [V]
Bis[1,2-benzenediolato(2-)O,O']-borate(1-) 	3.6
Bis[3-fluoro-1,2-benzenediolato(2-)O,O']borate(1-) 	3.75
Bis[2,3-naphthalenediolato(2-)O,O']-borate(1-) 	3.8
Bis[2,2'-biphenyldiolato(2-)O,O']-borate(1-) 	4.1
Bis[salicylato(2-)O,O']borate(1-) 	4.5
Bis[2-olatobenzenesulfonato(2-)O,O']borate(1-) 	4.3
Bis[5-fluoro-2-olatobenzene-sulfonato(2-)O,O']borate 	4.5

Phenoxide 	3.5
2,2-Biphenoxide 	3.7

2a) Experiments on platinum electrodes

In each case, 5 cyclic voltammograms were recorded successively in a measurement cell containing a stainless-steel, platinum or gold working electrode, a lithium counterelectrode and a lithium reference electrode. To this end, starting from the rest potential, the potential was firstly increased at a rate of 10 mV/s or 20 mV/s to 5 V against Li/Li⁺ and then moved back to the rest potential.

The salts shown in Table 2 exhibit the following characteristic behaviour. Depending on the oxidation potential, the salts mentioned are oxidized at potentials between 3.5 and 4.5 V against Li/Li⁺. However, this oxidation does not result in a lasting current increase, as in other salts, such as LiPF₆, Li imide or Li methanide, but, after passing through a maximum with relatively low currents, results in the formation of a passivating coating layer on the working electrode. Figure 4 shows this using the example of lithium bis[2-olatobenzenesulfonato(2-)O,O']borate(1-).

2b) Properties of the coating layer formed

In each case, 5 cyclic voltammograms were recorded successively in a measurement cell containing a stainless-steel working electrode, a lithium

- 22 -

counterelectrode and a lithium reference electrode. To this end, starting from the rest potential, the potential was firstly increased at a rate of 10 mV/s - 20 mV/s to values above the respective decomposition potential of the salts mentioned. A coating layer was deposited on the electrode in the process. The potential was then lowered to values below 0 V against Li/Li⁺, initiating deposition of lithium on the stainless-steel electrode. To this end, lithium ions must migrate through the film formed. In order to exclude dissolution of the coating layer during this operation, the potential was again increased to values above the respective decomposition potential of the salts mentioned. Figure 5 shows, in representative terms, the results obtained for lithium bis[2-olatobenzenesulfonato(2-)-O,O']borate(1-). Lithium cycling is possible in the electrolyte. This is evident from the deposition and dissolution peaks at low potentials. Furthermore, the coating layer is not dissolved by the selected process, since otherwise oxidation of the salt used at the abovementioned potentials would have to be detectable in the second and all subsequent cycles.

25 Example 3

Cycling experiments are carried out in button cells containing a metallic lithium anode and LiCoO₂. The electrolytes used were solutions of LiPF₆ in EC/DMC (1:1) containing 0% (control), 1% and 5% (based on the weight of LiPF₆) of lithium bis[salicylato(2-)-O,O']borate(1-).

The results are shown in Tables 3, 4 and 5.

Table 3: System 1M LiPF₆ in EC/DMC (1:1)

Cycle number	Charge capacity [mAh/g]	Discharge capacity [mAh/g]
1	164.6	153.4
2	155.1	153.6
3	155.0	153.8

Table 4: System 1M LiPF₆ in EC/DMC (1:1) + 1% lithium
5 bis[salicylato(2-)-O,O']borate(1-)

Cycle number	Charge capacity [mAh/g]	Discharge capacity [mAh/g]
1	164.0	153.7
2	155.2	153.0
3	154.0	152.8
4	153.4	152.5

Table 5: System 1M LiPF₆ in EC/DMC (1:1) + 5% lithium
10 bis[salicylato(2-)-O,O']borate(1-)

Cycle number	Charge capacity [mAh/g]	Discharge capacity [mAh/g]
1	163.4	149.8
2	151.0	148.9
3	149.5	148.1
4	148.4	147.1

It can be seen from the values shown that addition of
1% of borate does not have an adverse effect on the
performance of the cathode used.

15 Example 4

Behaviour of the additive lithium bis[salicylato(2-)-
O,O']borate(1-) on overcharging

20 The following measurement cycle was recorded in a
measurement cell containing a LiCoO₂ working electrode,

- 24 -

a lithium counterelectrode and a lithium reference electrode.

5 The electrolyte used was solutions of LiPF_6 in EC/DMC (1:1) containing 0% (reference) or 1.5% of lithium bis[salicylato(2-)-O,O']borate(1-).

10 Starting from the rest potential, the cathode was firstly charged at a charge rate of C/15 - C/18 to 4.3 V against Li/Li^+ .

15 The cathode was subsequently overcharged at a charge rate of C/5. During this operation, the voltage reached a specified maximum value of 6 V against Li/Li^+ . If the potentiostat/galvanostat is then switched off, the potential drops very rapidly.

Comparison of the curves (Figures 6 and 7):

20 Compared with the reference, the potential increase in the region 4.3 - 6 V against Li/Li^+ is slower in the case of the electrolyte containing lithium bis[salicylato(2-)-O,O']borate(1-). This can be explained by a desired decomposition of the additive
25 and the consequent formation of a coating layer.

After the external potentiostat/galvanostat has been switched off, thus simulating release of the disconnecter, the potential of the cathode in both
30 electrolytes drops rapidly.

In the case of the electrolyte with no additive, the potential oscillates about values of 4.2 - 4.3 V against Li/Li^+ . The cathode remains in the charged,
35 high-energy state.

By contrast, addition of lithium bis[salicylato(2-)-O,O']borate(1-) causes the potential to drop to values of about 3.7 V against Li/Li^+ . This corresponds to the

rest potential of an uncharged LiCoO_2 electrode. This suggests that lithium bis[salicylato(2-)-O,O']borate(1-) or the film formed by decomposition of the additive is capable of dissipating the "excess potential" in a controlled manner by internal self-discharge and thus converting the battery into a low-energy state after release of a safety means (for example disconnecter).

10 Example 5

Cycling experiments are carried out in button cells containing a metallic lithium anode and LiMn_2O_4 . The electrolytes used are solutions of LiPF_6 in EC/DMC (1:1) containing 0% (control) and 0.2% (based on the weight of the electrolyte) of lithium bis[2,2'-biphenyldiolato(2-)-O,O']borate(1-).

Table 6: System 1M LiPF_6 in EC/DMC (1:1)

Cycle number	Charge capacity [mAh/g]	Discharge capacity [mAh/g]
1	119.8	112.5
2	111.8	111.6
3	111.0	110.3
4	109.8	110.2
5	111.5	110.8
6	109.6	109.0
7	108.7	108.8
8	108.7	108.8
9	109.0	108.7
10	108.1	107.7

Table 7: System 1M LiPF₆ in EC/DMC (1:1) + 0.2% of lithium bis[2,2'-biphenyldiolato(2-)O,O']borate(1-)

Cycle number	Charge capacity [mAh/g]	Discharge capacity [mAh/g]
1	130.5	119.8
2	118.4	117.0
3	118.6	117.4
4	117.7	116.5
5	116.5	115.5
6	116.9	114.8
7	116.4	114.4
8	114.5	113.4
9	113.8	112.8
10	113.9	113.6

- 5 It can be seen from the values shown that addition of 0.2% of lithium bis[2,2'-biphenyldiolato(2-)-O,O']borate(1-) has a significant, positive effect on the performance of the cathode used.

26474-585

- 27 -

CLAIMS:

1. Electrolyte comprising a lithium-containing inorganic conductive salt or lithium-containing organic conductive salt, dissolved in an aprotic solvent, from the group of the methanides, triflates and imides and at least one organic salt of the formula (I) as additive



in which

m and p denote 0, 1, 2, 3 or 4, where m+p=4 and

- 10 R^1 and R^2 are identical or different, are optionally bonded directly to one another by a single or double bond, each, individually or together, have the meaning of an aromatic or aliphatic carboxylic or sulfonic acid, or each, individually or together, have the meaning of an aromatic ring from the group phenyl, naphthyl, anthra-cenyl or phenanthrenyl, which are unsubstituted or mono- to tetrasubstituted by A or Hal, or

- each, individually or together, have the meaning of a heterocyclic aromatic ring from the group pyridyl, pyrazyl or bipyridyl, which are unsubstituted or mono- to trisubstituted by A or Hal, or each, individually or together, have the meaning of an aromatic hydroxy acid from the group of aromatic hydroxycarboxylic acids or aromatic hydroxysulfonic acids, which are unsubstituted or mono- to tetrasubstituted by A or Hal, and

Hal denotes F, Cl or Br and

A denotes alkyl having 1 to 6 C atoms, which is optionally mono- to trihalogenated,

26474-585

- 28 -

where the additive is present in a concentration between 0.1 and 10%, based on the total weight of conductive salt.

2. Electrolyte according to claim 1, wherein the additive is selected from the group:

- 5 lithium bis[1,2-benzenediolato(2-)O,O']borate(1-),
 lithium bis[3-fluoro-1,2-benzenediolato(2-)O,O']borate(1-),
 lithium bis[2,3-naphthalenediolato(2-)O,O']borate(1-),
 lithium bis[2,2'-biphenyldiolato(2-)O,O']borate(1-),
 lithium bis[salicylato(2-)O,O']borate(1-),
 10 lithium bis[2-olatobenzenesulfonato(2-)O,O']borate(1-),
 lithium bis[5-fluoro-2-
 olatobenzenesulfonato(2-)O,O']borate(1-),
 lithium bis[oxalato(2-)O,O']borate(1-), and
 lithium bis[malonato(2-)O,O']borate(1-).

15 3. An electrochemical cell comprising a cathode, anode, separator and electrolyte, wherein the electrolyte is as defined in claim 1 or 2.

4. Use of an organic salt of the formula (I), as defined in claim 1, as an additive in the electrolyte of
 20 electrochemical cells, batteries and secondary lithium batteries, wherein the additive has a concentration which is at least ten fold smaller than the concentration of the conductive salt.

1/7

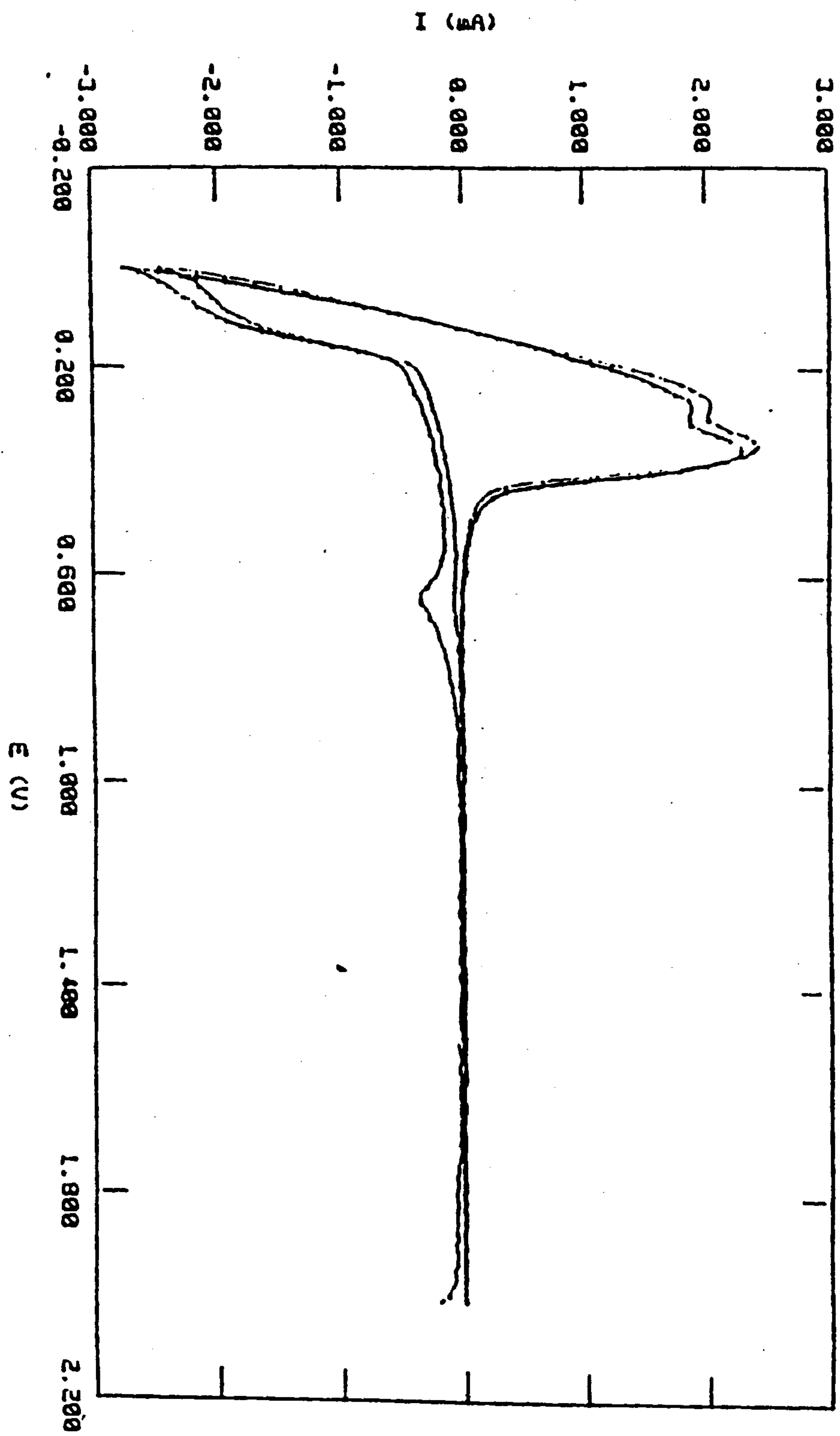


Fig. 1

2/7

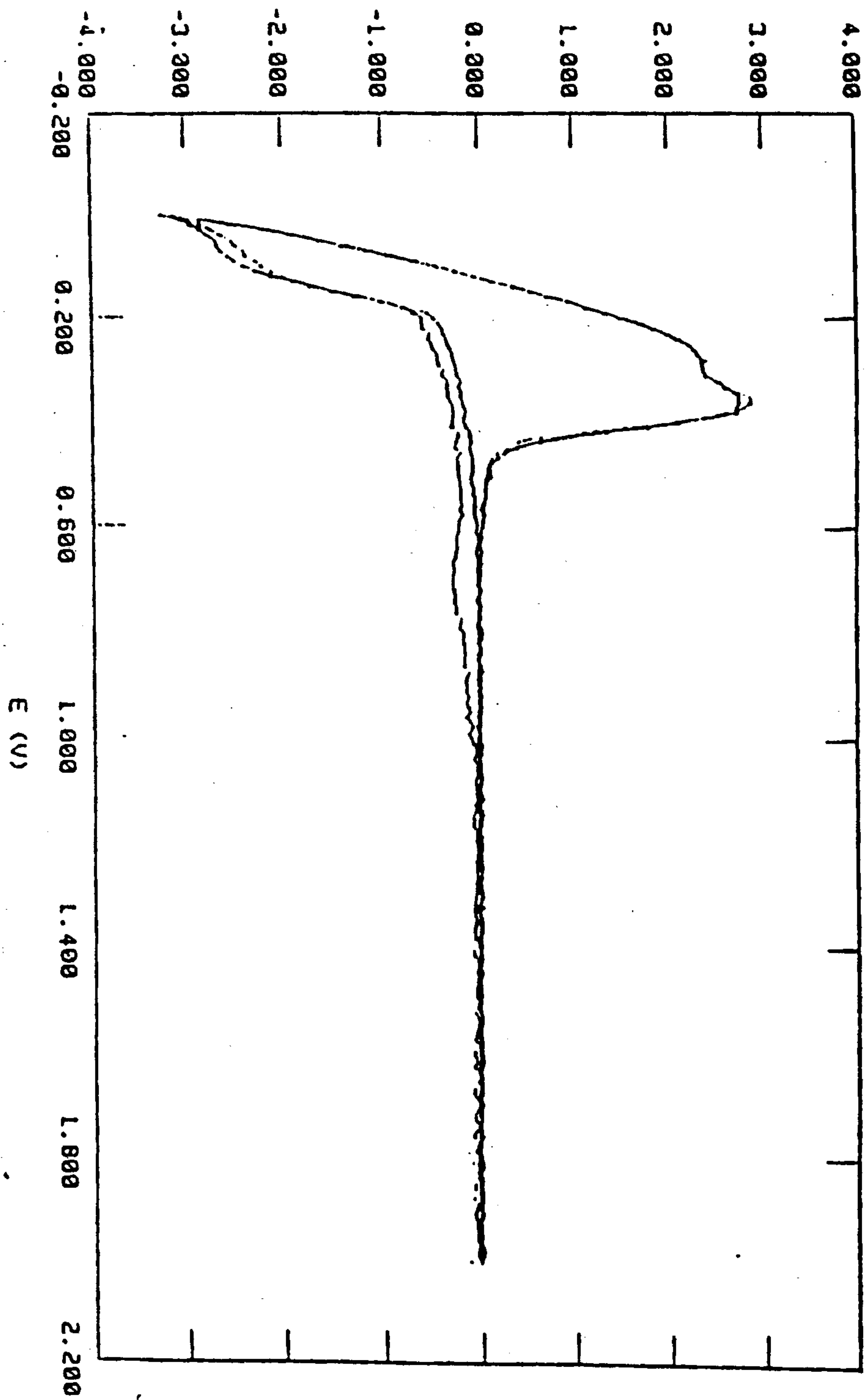


Fig. 2

3/7

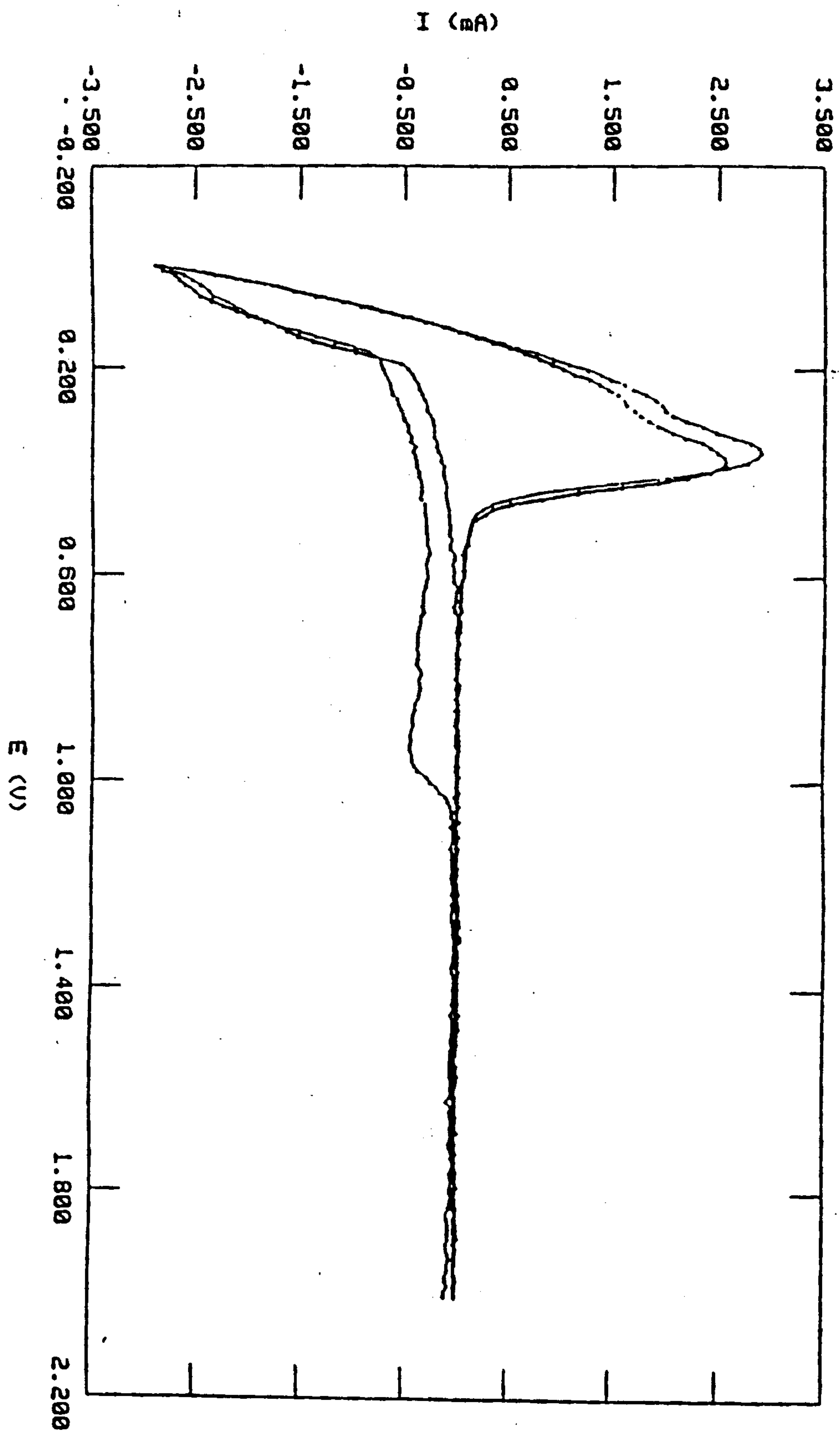


Fig. 3

4/7

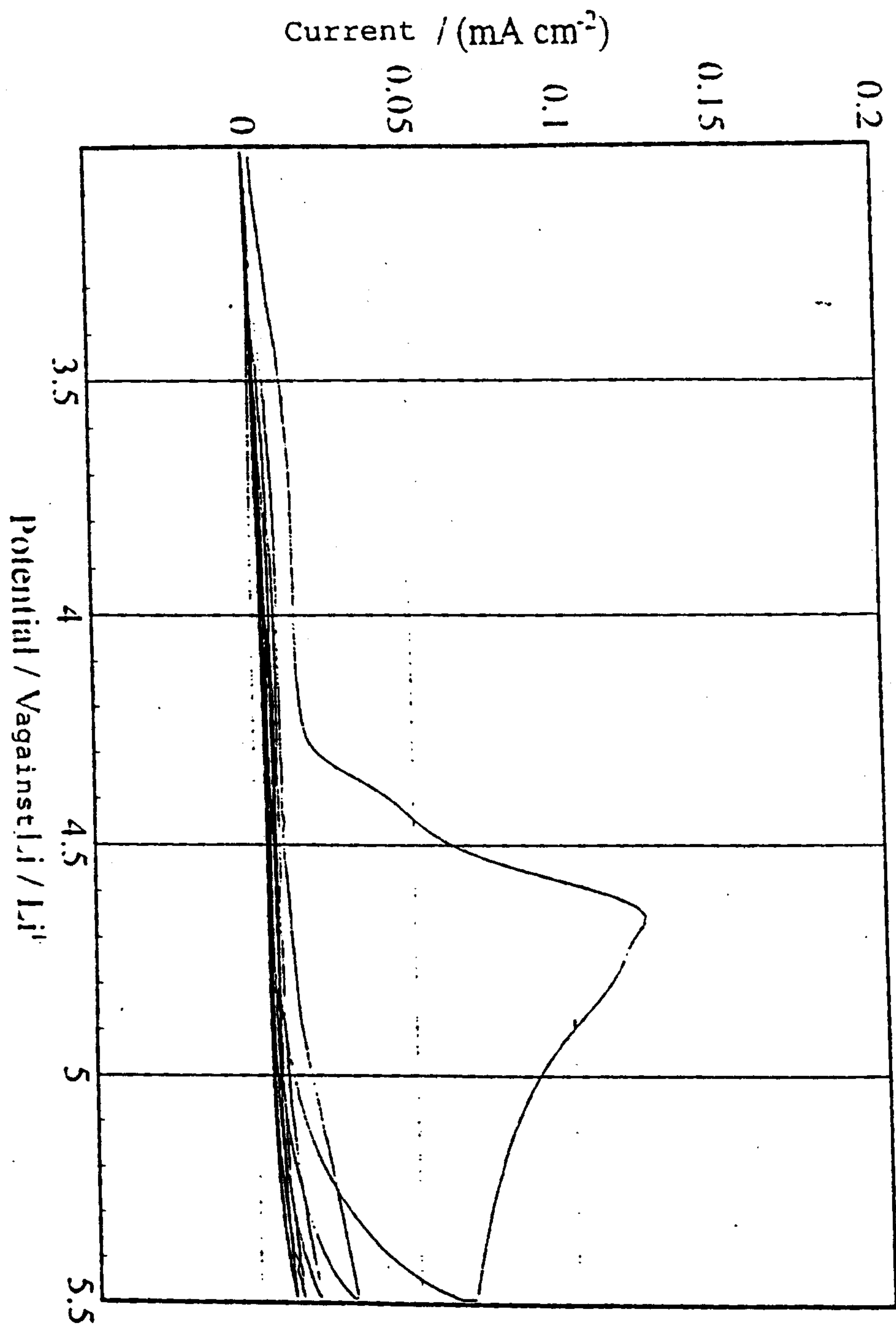


Fig. 4

5/7

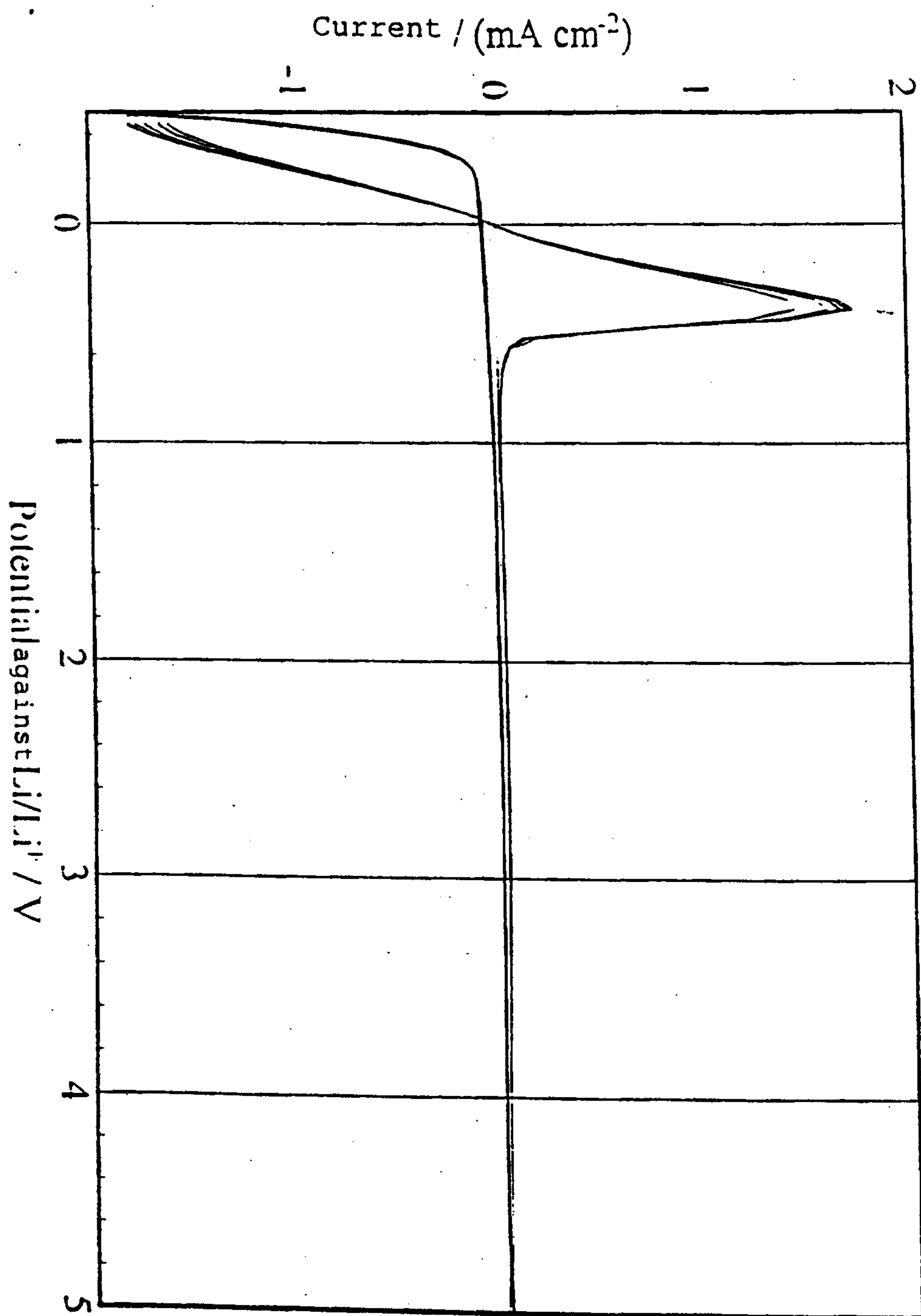
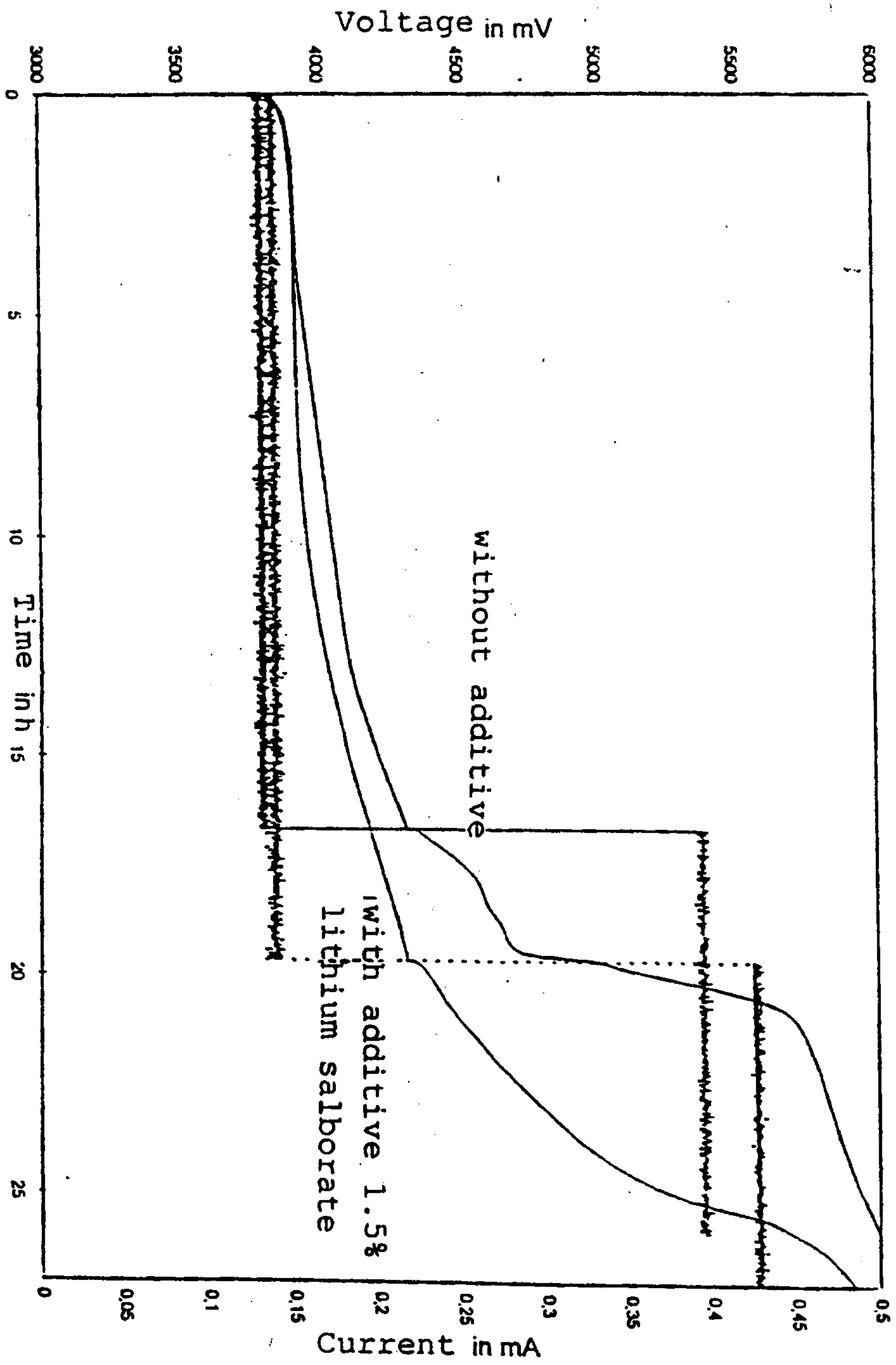


Fig. 5

6/7

Fig. 6



7/7

Fig. 7

