

The logo of the University of Bordeaux is displayed against a background with a blue diagonal stripe in the top left and a dark grey diagonal stripe in the bottom right. The text 'université' is in a dark brown, lowercase, sans-serif font, with a blue square above the 'i' and a blue 'e'. Below it, the word 'de' is in a smaller, dark brown, lowercase, sans-serif font. To the right of 'de', the word 'BORDEAUX' is in a bold, dark brown, uppercase, sans-serif font.

université
de **BORDEAUX**

9h30 – 12 h15

Histoire et concept électricité et stimulations cérébrales
non invasives

14h00 – 16h00

Rappel sur l'électrophysiologie
Lecture EEG lors ECT
Machines ECT, quels choix

DIU Pratique et théorie de l'électroconvulsivothérapie

Diplôme Inter-Universitaire « PRATIQUE ET THÉORIE DE L'ÉLECTROCONVULSIVOTHÉRAPIE »
Année universitaire 2021-2022 / Université de Bordeaux – Nantes – Lyon

SEMINAIRE 1 BORDEAUX du 31 janvier au 2 février 2022

<https://formatoile2.u-bordeaux.fr/course/view.php?id=3366>

<https://u-bordeaux-fr.zoom.us/j/87470030133?pwd=M2ZEOU9YTakyYnJGbmZlcVRnNXZYdz09>
ID de réunion : 874 7003 0133
Code secret : 781942

Lundi 31 janvier 2022	Mardi 1 ^{er} février 2022	Mercredi 2 février 2022
Cours enregistré, vidéo en différé (lien à venir) : Informations générales / le DIU ECT en 60 mn (Marc AURIACOMBE) 13 h 00 – 14 h 00 Session en direct : questions et réponses au sujet du DIU ECT	9 h 30 – 12 h 15 (intervention en direct) Histoire et concept électricité et stimulations cérébrales non invasives (Christophe DAUDET) (Jean-Arthur MICOULAUD-FRANCHI)	10 h00 – 11 h 30 (intervention en direct) Stimulation cérébrale non invasive et réseaux neuronaux (Christophe DAUDET) 11 h 30 - 12 h 30 Mémoires Discussions des sujets
14 h 00 – 16 h 45 (intervention en direct) Mémoire et dépression, Comment les mesurer avec quels outils ? (Sophie AURIACOMBE) Les effets de l'ECT sur la cognition : rappels sur la mémoire, Mémoire et dépression, (Sophie AURIACOMBE)	14 h 00 – 16 h 00 (intervention en direct) Rappel sur l'électrophysiologie Lecture EEG lors ECT Machines ECT, quels choix (Jean-Arthur MICOULAUD-FRANCHI) (Christophe DAUDET)	14 h 30 – 16 h 30 (intervention en direct) Phénoménologie de l'ECT. Rôle de la crise et de l'électricité (Marc AURIACOMBE) Cours enregistré, vidéo en différé (lien à venir) : Données de neuroimagerie dans la dépression dans le cadre de l'utilisation des ECT et de la rTMS

Renseignements pédagogiques : Pr Marc Auriacombe : marc.auriacombe@u-bordeaux.fr; Pr Emmanuel Poulet : emmanuel.poulet@chu-lyon.fr;
Pr Anne Sauvaget : Anne.sauvaget@chu-nantes.fr

Renseignements administratifs pour Lyon : claudine.martinenghi@chu-lyon.fr
pour Nantes : anne.sauvaget@chu-nantes.fr
pour Bordeaux : alexia.larran@u-bordeaux.fr

Examen écrit : 1^{ère} session
2^{ème} session (si échec à la 1^{ère} session)
Date limite remise mémoire
Soutenance des mémoires validé

juin 2022 à Bordeaux, Lyon, Nantes
septembre 2022 à Bordeaux, Lyon, Nantes
octobre 2022 (validation sujet avril 2022)
décembre 2022, en visio Bordeaux, Lyon, Nantes

11 h - 13 h
11 h - 13 h



Programme

1. Evolutions des objets technologiques de stimulations électriques cérébrales - 3h

- *En lien avec les évolutions sociales de l'électricité: entre vie et mort - 1h30*
 - *Evolution des technologies électriques : une vieille histoire (jam)*
 - *Deux types d'électricité : galvanisation (continu) / faradisation (alternatif) (cd)*
- *En lien avec le monde médicale : la recherche de la preuve - 1h30*
 - *Le cas Mesmer : naissance de l'essai contrôlé randomisé (jam)*
 - *Le cas des thérapies électriques et magnétiques : des fièvres aux chocs (cd)*

2. La mesure électro-physiologique EEG pendant les ECT - 1h

- *Enregistrement EEG pendant une séance d'ECT*
- *Lecture EEG*
- *Adaptation de la cure en fonction de l'EEG*

3. Choix d'une machine d'ECT - 1h

- *Evolution des machines*
- *Critères de comparaisons et de choix*
- *La recherche sur les machines ECT*



L'EEG

université
de **BORDEAUX**

Les machines d'ECT actuel
sont aussi des machines d'EEG

**LES ECT SONT UNE TECHNIQUE
DE NEUROPHYSIOLOGIE INTERVENTIONNELLE !**

Qui a réalisé de la neurophysiologie interventionnelle en premier ?

LES PSYCHIATRES !

Le début des ECT, n'est pas le début de l'électrothérapie



Cerletti et Beneti
ECT
1938

L'électrophysiologie interventionnelle

ÉLECTROPHYSIOLOGIE INTERVENTIONNELLE

La technique se propose de repérer par cartographie endocavitaire, puis de détruire la zone de myocarde arythmogène, les voies accessoires ou le tissu de conduction pour guérir définitivement le patient de son arythmie ou pour améliorer la tolérance fonctionnelle des tachycardies récidivantes [7-21]. La destruction est opérée par le cathéter d'ablation au travers duquel une source d'énergie est appliquée sur le substrat arythmogène.

RECOMMANDATIONS



Recommandations
de la Société française de cardiologie
concernant l'électrophysiologie
diagnostique et interventionnelle,
la stimulation cardiaque permanente
et la défibrillation automatique
implantable

La neurophysiologie interventionnelle

Neurophysiol Clin 2001 ; 31 : 215-7
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S0987705301002660/EDI
www.elsevier.fr/direct/nc-cn

ÉDITORIAL INVITÉ

Plaidoyer pour une neurophysiologie interventionnelle

La participation de la neurophysiologie clinique prend place non seulement en salle d'opération au cours de l'intervention chirurgicale elle-même – c'est le « *Monitoring* peropératoire »

la neurophysiologie est, non pas seulement utile, mais réellement nécessaire à la thérapeutique. Plus que cela même, elle fait partie de cette thérapeutique et ne peut être dissociée de l'acte chirurgical

Les ECT et la neurophysiologie clinique

*Il s'agit d'un excellent travail mais qui intéressera davantage
**des non-spécialistes en
neurophysiologie clinique**
faisant appel à la méthode, en
particulier, **des psychiatres.***

*Ma suggestion serait dès lors de le
soumettre à une revue psychiatrique,
soit française (**je ne les connais pas
bien**),
soit belge (par exemple, les *Acta
Psychiatrica Belgica*).
**Je suis certain qu'un tel article
pourrait leur être d'une grande
utilité.***

Annales Médico-Psychologiques, xxx (2013) xxx-xxx

DÉVELOPPEMENT PROFESSIONNEL CONTINU

Neurophysiologie clinique en psychiatrie : 3 – Électroencéphalographie pendant les séances d'électroconvulsivothérapie

Clinical neurophysiology in psychiatry: 3 – Electroencephalography during electroconvulsive therapy sessions

Jean-Arthur Micoulaud-Franchi^{a,b,c,d,e}, Raphaëlle Richieri^{a,c}, Clélia Quiles^{d,e},
Céline Baltani^{a,b}, Christophe Lancon^{a,c}, Jean Vion-Dury^{a,b}

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Sainte-Marguerite, 270, boulevard de Sainte-Marguerite, 13009 Marseille, France

^b Laboratoire de neurosciences cognitives (LNC), UMR CNRS 7291, 31 All-Marseille université, site Saint-Charles, 3, place Victor-
Hugo, 13331 Marseille cedex 3, France

^c EA 3279, laboratoire de santé publique, évaluation des systèmes de soins et santé perçue, faculté de médecine, université de la
Méditerranée, 27, boulevard Jean-Moulin, 13385 Marseille cedex 05, France

^d Pôle universitaire de psychiatrie adulte, centre hospitalier Charles-Ferrès, 121, rue de la Béchade, 33076 Bordeaux cedex,
France

^e Université Bordeaux Segalen, 146, rue Léon-Saignes, 33076 Bordeaux cedex, France

Résumé

La surveillance électroencéphalographique (EEG) des séances d'électroconvulsivothérapie (ECT) est indispensable pour deux raisons. Premièrement, la surveillance EEG des séances ECT est la méthode la plus efficace pour détecter une crise épileptique prolongée. Éviter une crise prolongée permet de minimiser les effets secondaires cérébraux des ECT. En effet, une crise épileptique prolongée augmente à court terme le risque d'état de mal épileptique post-ECT, complication rare mais grave des ECT, et à moyen terme augmente le risque de mauvaise tolérance cognitive de la cure ECT sans en augmenter l'efficacité clinique. Deuxièmement, la surveillance EEG des séances ECT est une méthode, complémentaire à la surveillance clinique, pour permettre de confirmer la présence d'une crise épileptique adéquate voire optimale et d'adapter les modalités de la stimulation électrique afin de maximiser l'efficacité des ECT. Cet article propose une aide à la lecture de l'EEG pendant les séances ECT et un arbre décisionnel de la pratique ECT guidée par cette lecture. La volonté n'est pas de résumer l'intérêt de la conduite des cures ECT en psychiatrie, mais de souligner la place centrale que peut prendre la neurophysiologie clinique dans une stratégie thérapeutique psychiatrique.
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Abstract

Electroencephalographic (EEG) monitoring during electroconvulsive therapy (ECT) sessions is essential for two reasons. First, the EEG monitoring during ECT sessions is the most effective method to detect a prolonged seizure. Avoid prolonged crisis allows to minimize the brain side effects of ECT. Indeed, prolonged seizures, in the short-term, increase the risk of

* Auteur correspondant.

Adresse e-mail : jmicoulaudfranchi@gmail.com (J.-A. Micoulaud-Franchi).

¹ Cette unité fait partie du réseau FondeMentel.

Outil de stimulation et d'enregistrement électrique



Gouttière dentaire de protection

Electrodes de stimulation et pâte conductrice

Electrodes d'enregistrement
voie gauche et voie droite

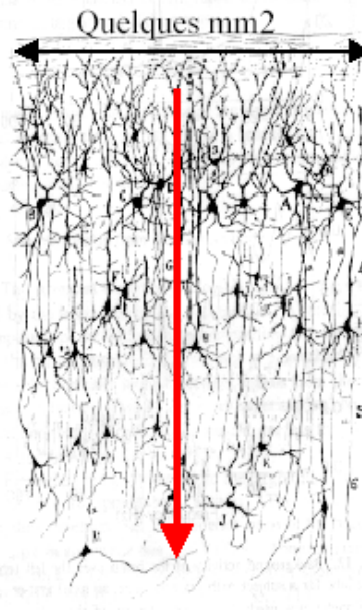
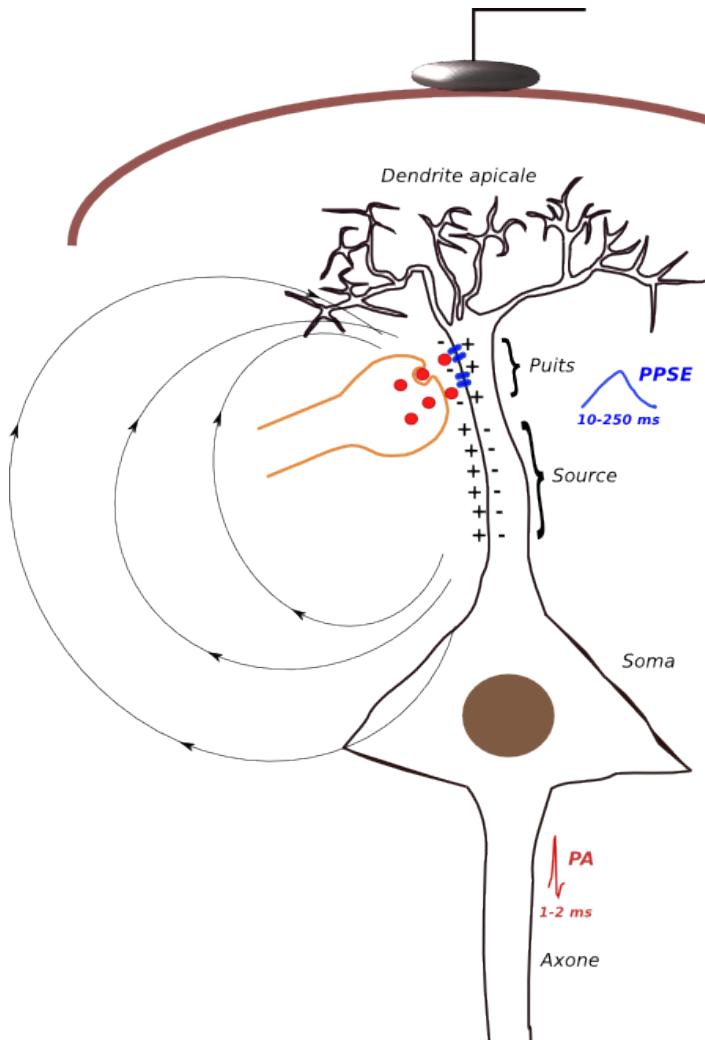
Electrode de terre

Papier d'enregistrement EEG

Ecran de réglages des
paramètres de stimulation et
d'enregistrement EEG

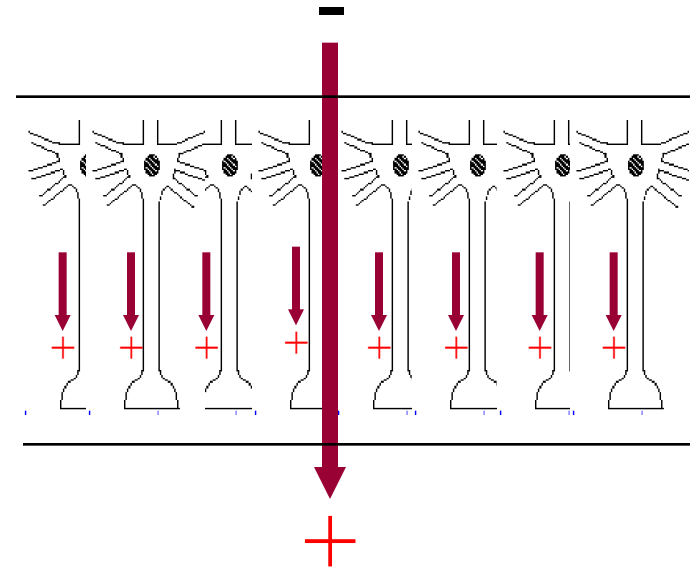
Qu'enregistre un signal EEG ?

Qu'enregistre l'EEG ?

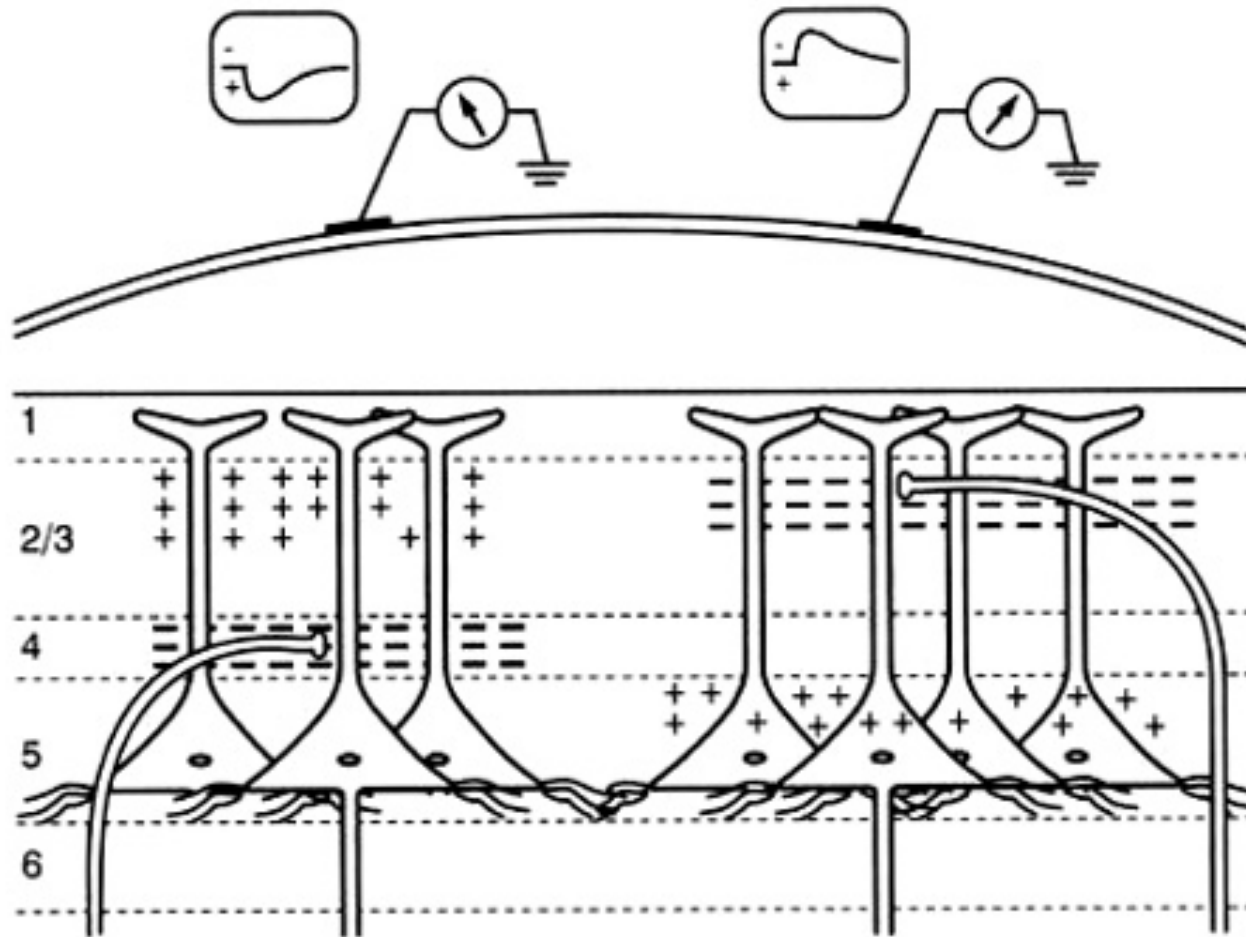


Organisation // des neurones dans le cortex

Dipole résultant
= Σ vectorielle des dipôles unitaires

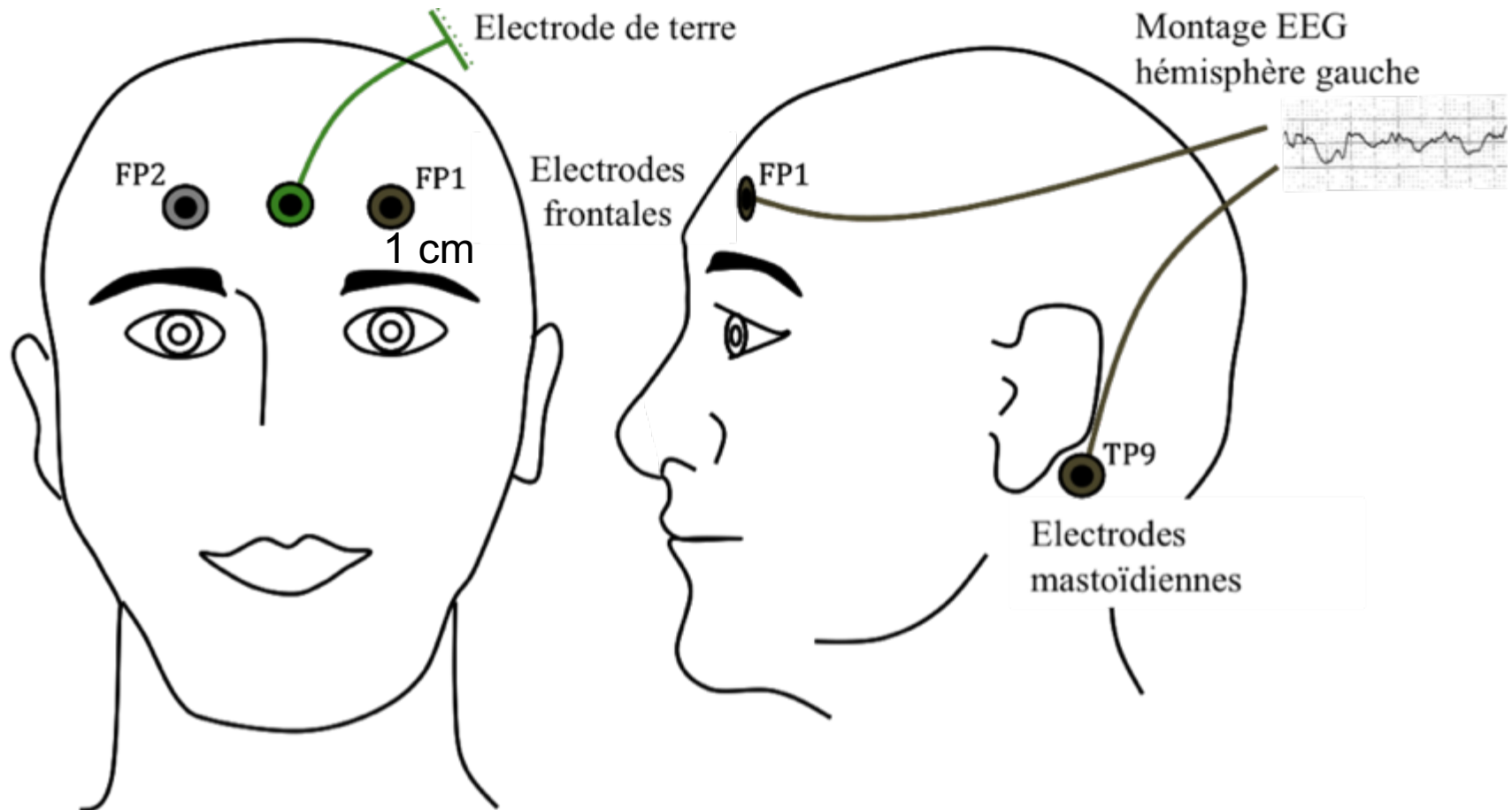


Qu'enregistre l'EEG ?



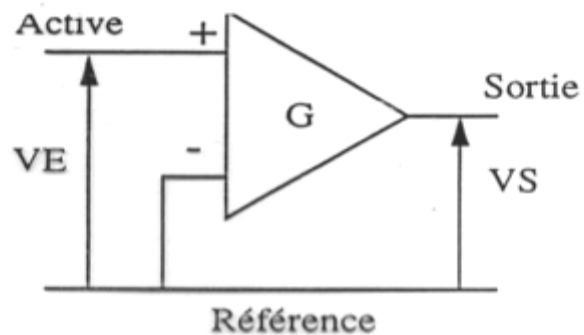
Comment enregistrer un signal EEG ?

Placement des électrodes



Amplification

Amplificateur simple



$$V_S = G \times V_E$$

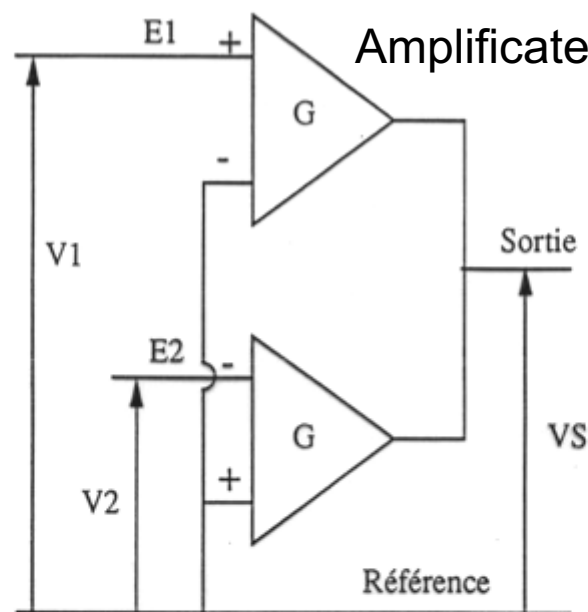
$$\text{Gain } G = V_S / V_E$$

Entrée

Sortie
amplifiée



Amplificateur différentiel



$$V_S = G \times (V_1 - V_2)$$

V1

V2

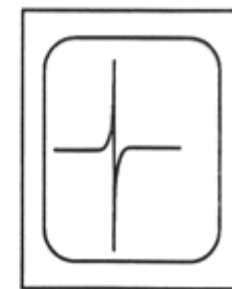
$G \times (V_1 - V_2)$



-



=

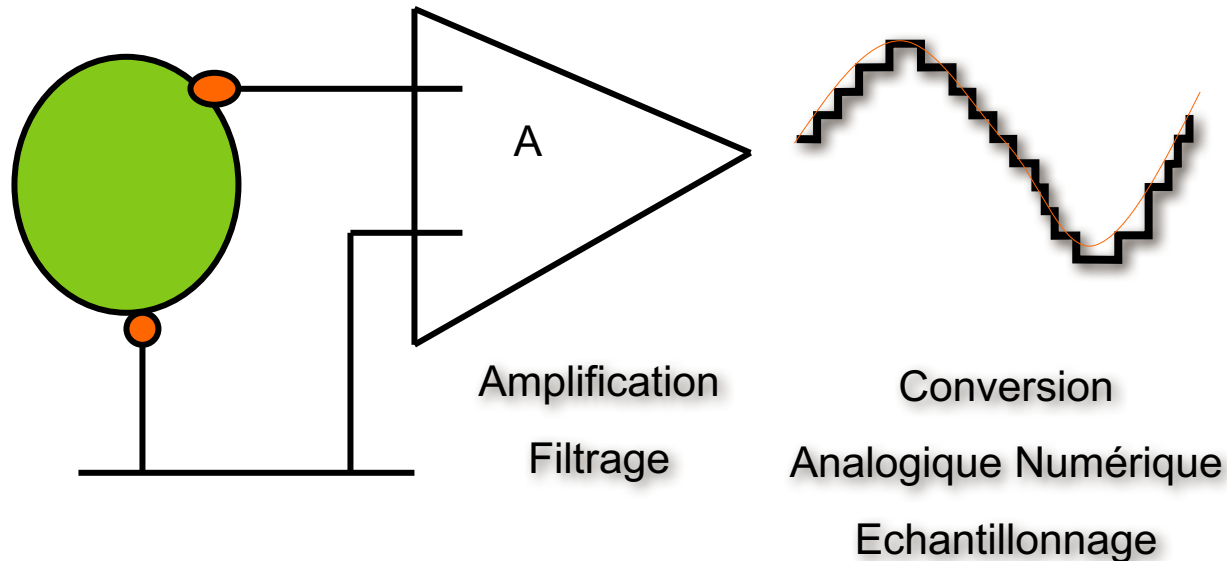


Rejection en mode commun

La chaine d'acquisition

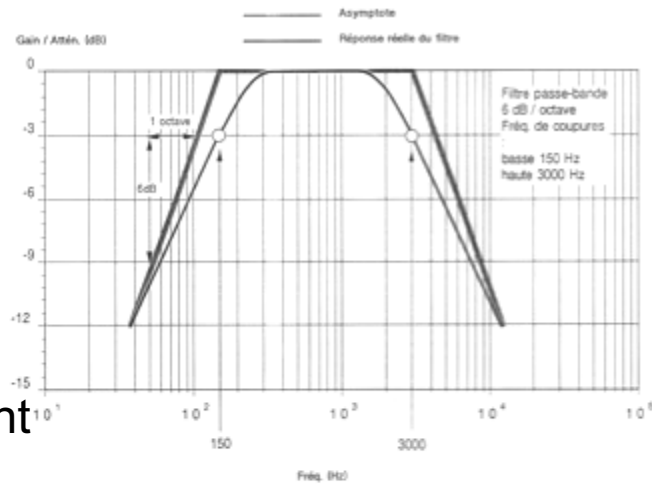
Potentiel d'électrode et impédance EEG : 100 μ V, 0,16 - 30 Hz Résolution
EMG : 10 m V, 20 - 10 000 Hz Précision
Gain d'amplification
Sensibilité (saturation)

Gain à l'affichage
Filtre à l'affichage



Visualisation du signal
Traitement du signal

Filtres



Mouvement
Sudation

EMG
Bruits
électriques

Filtre passe haut Filtre passe bas

L'EEG traditionnel

L'EEG conventionnel

Type de montage
présenté
Echelle de temps (vitesse
déroulement du papier)
Sensibilité en $\mu V/mm$
Valeur du filtre passe haut
constante de temps
Valeur du filtre passe-bas

Paramètres de l'enregistrement EEG

Montage
longitudinal

Dérivations
droites

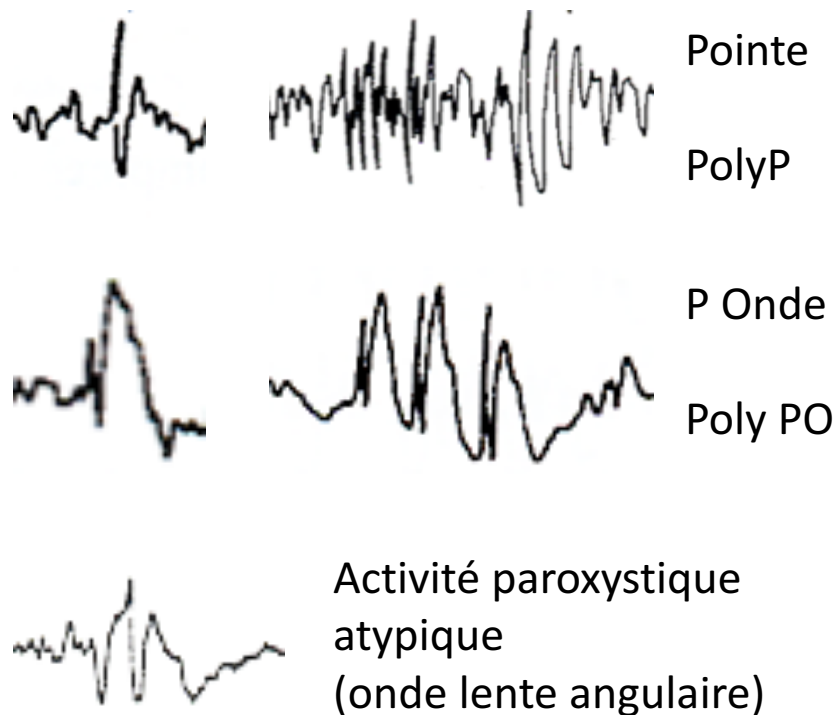
Dérivations
gauches

ECG
Ligne sur laquelle sera
indiqués les éclairs de
Informations temporel

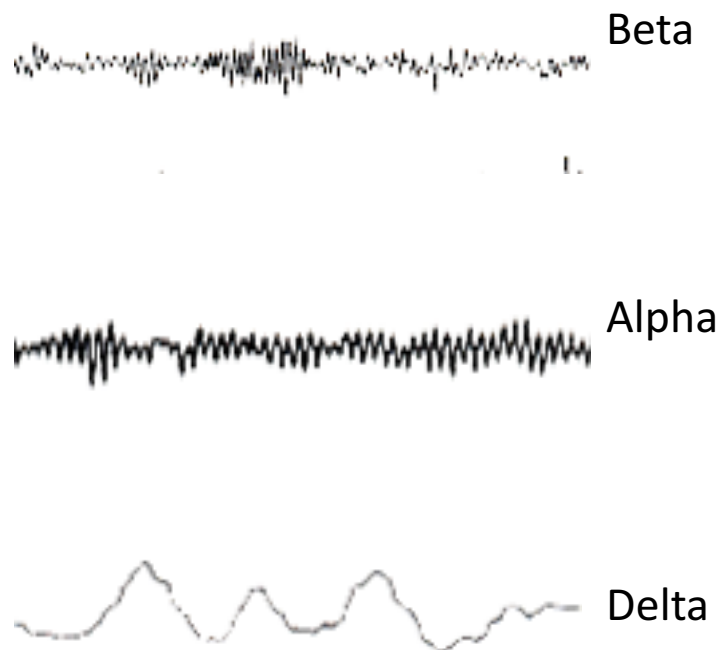


Le vocabulaire de description d'un EEG

Activités paroxystiques « épileptiformes »

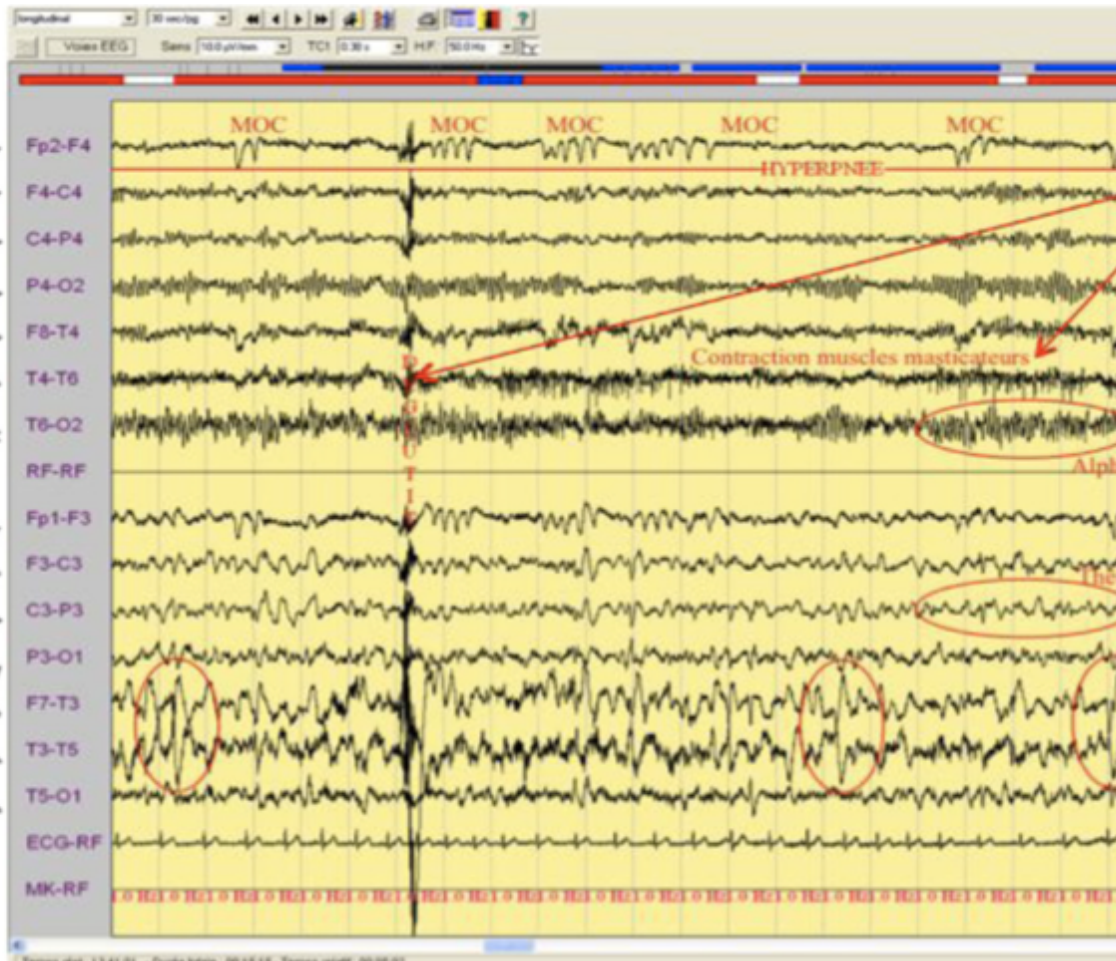
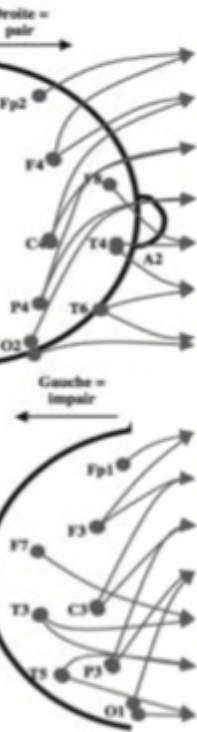


Rythme de fond



Ce que l'on peut voir sur un EEG

Montage longitudinal



Descriptions EEG

Signifi EE

Mouvements oculaires

Mouvement de déglutition

Activités musculaires

Activité de fond postérieure dans la bande alpha, au niveau de l'hémisphère droit

Rythme de fond hémisphérique gauche ralenti dans la bande thêta

Paroxysmes de type ondes lentes angulaires et pointes ondes dégradés focalisés (en opposition de phase) sur T3

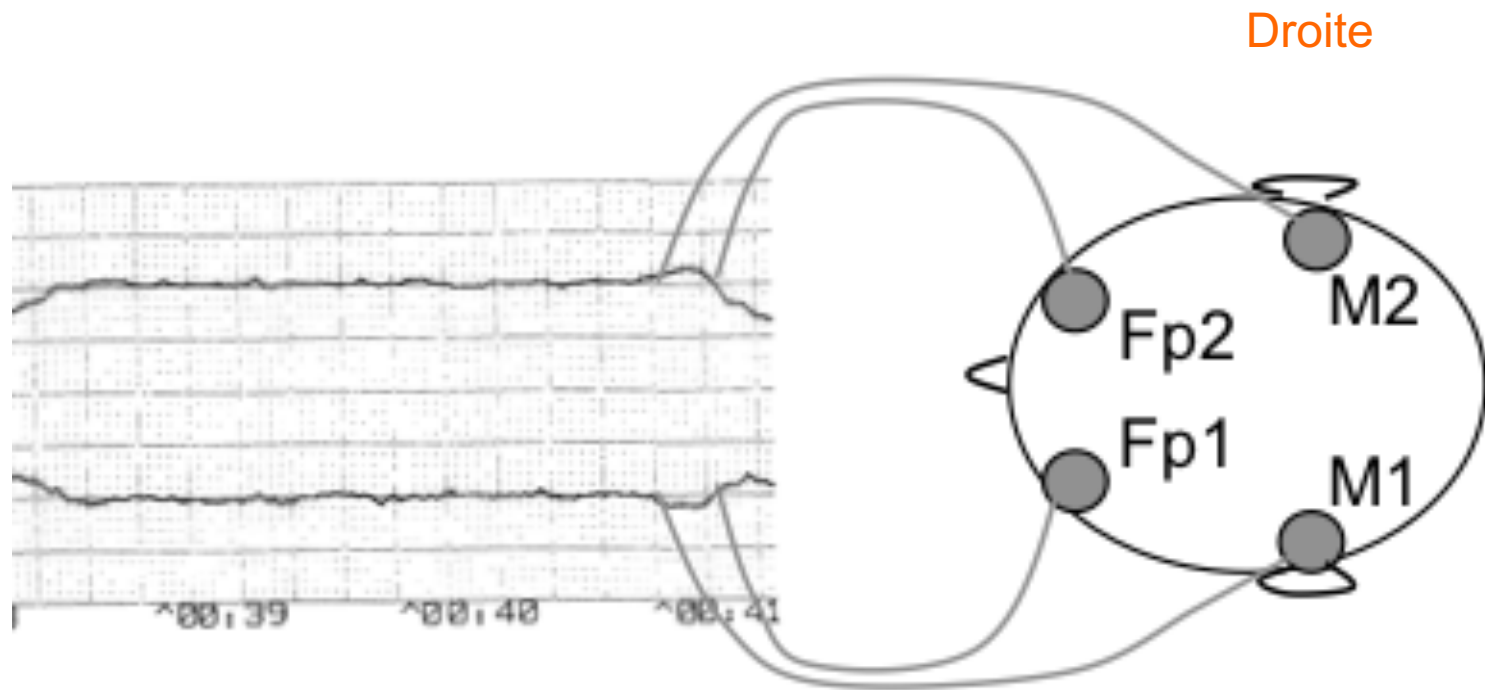
Artéf
biolog

Activ
physi
droite

Rythr
patho
gauch

Parox
inter-

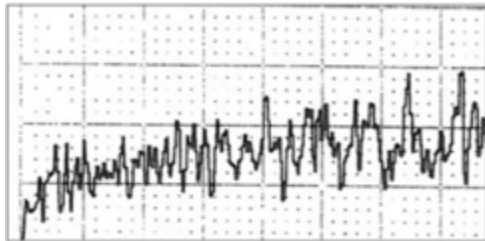
L'EEG des ECT



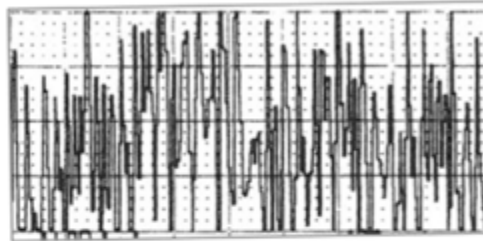
Gauche

Reconnaître les phases EEG de la crise

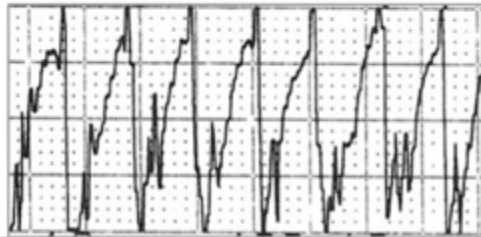
Phase critique 1:
Recrutement



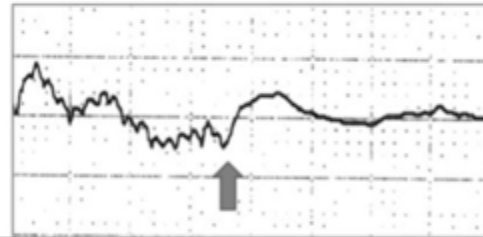
Phase critique 2:
Tonique (pointes)



Phase critique 3:
Clonique (pointes ondes)

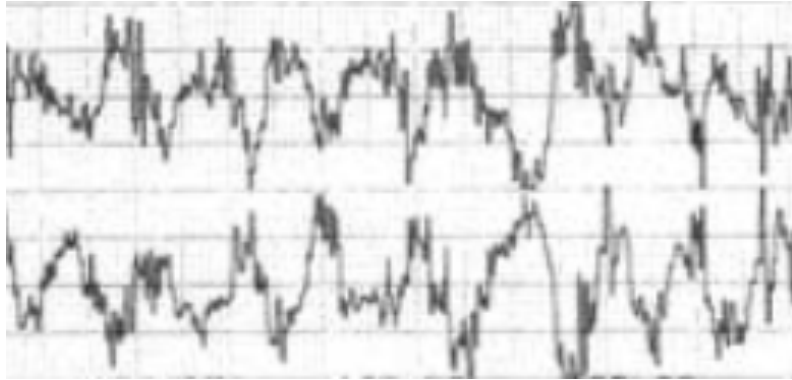


Phase critique 4 et
Suppression post-critique

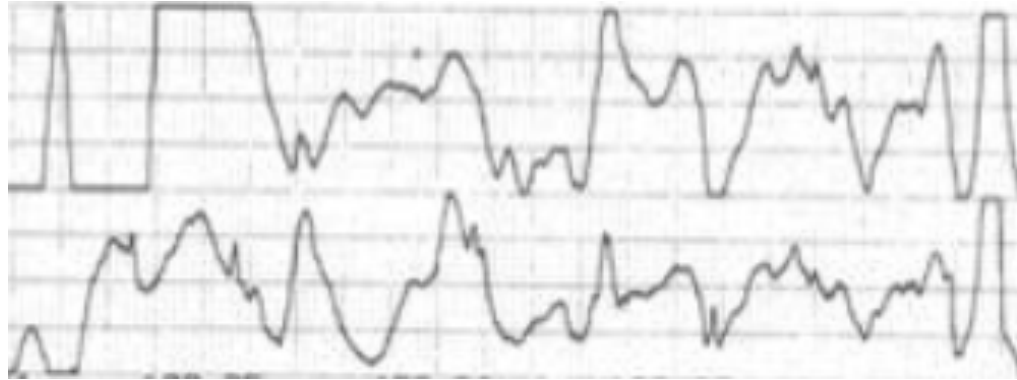


Reconnaître les artéfacts biologiques et techniques

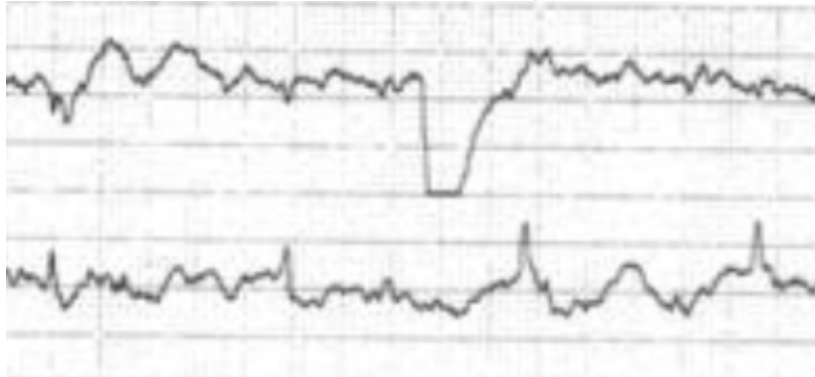
Artéfacts musculaires
(surchargeant des ondes lentes angulaires)



Artéfacts de mouvements
(manipulation de la tête du patient, ventilation au masque, etc.)



Artéfacts d'électrocardiogramme



Artéfacts d'électrode



L'EEG et les machines ECT

MECTA (spectrum) vs THYMATRON



Différence de principe de choix de dose

Spectrum



Thymatron



Titration

Sackheim
Duke université

Age / dose

Abrams

Table de titration

Exemple de table avec des *pulses* brefs sur MECTA spECTrum® pour comprendre le principe de la titration et l'augmentation par palier des charges électriques délivrées.

	Durée du <i>pulse</i>	Fréquence de stimulation	Durée de la stimulation	Intensité	Charge électrique (mC)	Quantité d'énergie (pour une résistance de 300 Ω)
1	1	40	0,5	0,8	32	7,6
2	1	40	0,75	0,8	48	11,5
3	1	40	1,25	0,8	80	19,2
4	1	40	2	0,8	128	30,7
5	1	60	2	0,8	192	46
6	1	60	3	0,8	288	69,1
7	1	90	3	0,8	432	103
8	1,8	90	2,5	0,8	648	155
9	1,8	90	3,5	0,8	907,2	248
10	2	90	4	0,8	1 152	275

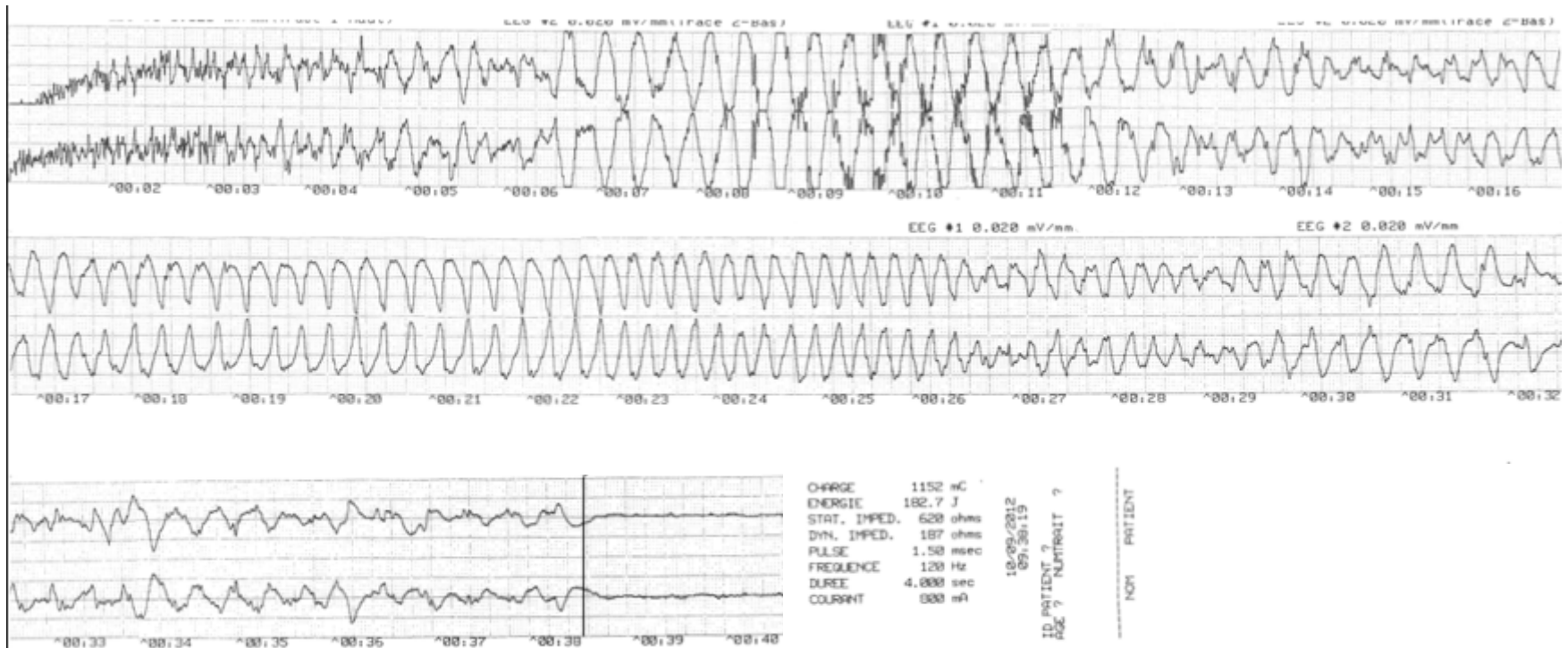
Il existe différents types de tables de titration qui peuvent dépendre du type de machine (MECTA spECTrum® et Somatics Thymatron®) et du type de *pulse* (ultra-bref ou bref), mais le principe reste le même.

Age / dose

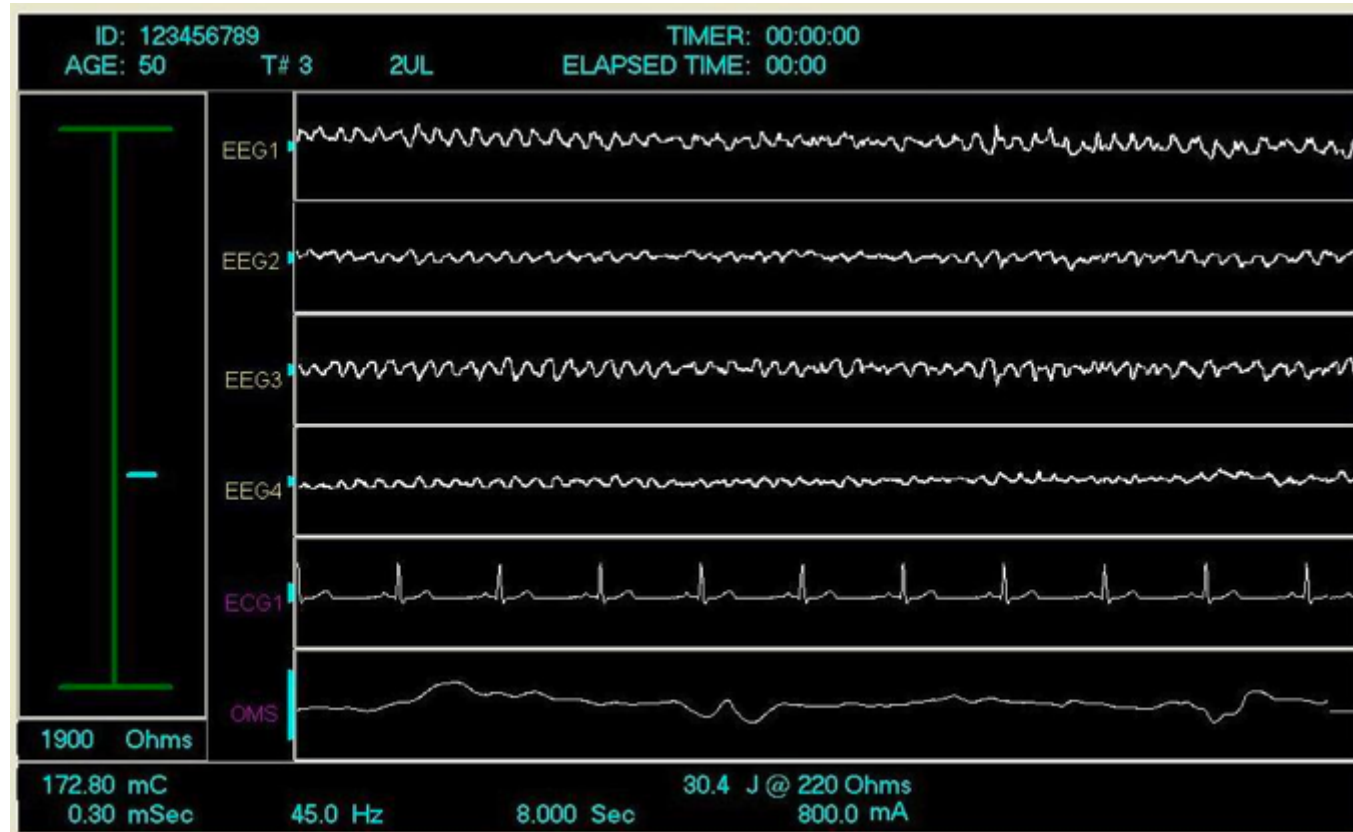
Exemple de table âge/dose pour comprendre le principe de la détermination de la charge électrique à délivrer en fonction de l'âge.

Âge	Fréquence de stimulation	Durée de la stimulation	Charge électrique (mC)
5	30	0,47	25,2
10	30	0,93	50,4
15	340	1,4	75,6
20	30	1,87	100,8
25	30	2,33	126
30	50	1,68	151,2
35	50	1,96	176,4
40	50	2,24	201,6
45	50	2,52	226,8
50	50	2,8	252
55	70	2,2	277,2
60	70	2,4	302,4
65	70	2,6	327,6
70	70	2,8	352,8
75	70	3	378
80	70	3,2	403,2
85	70	3,4	428,4
90	70	3,6	453,6
95	70	3,8	478,8
100	70	4	504

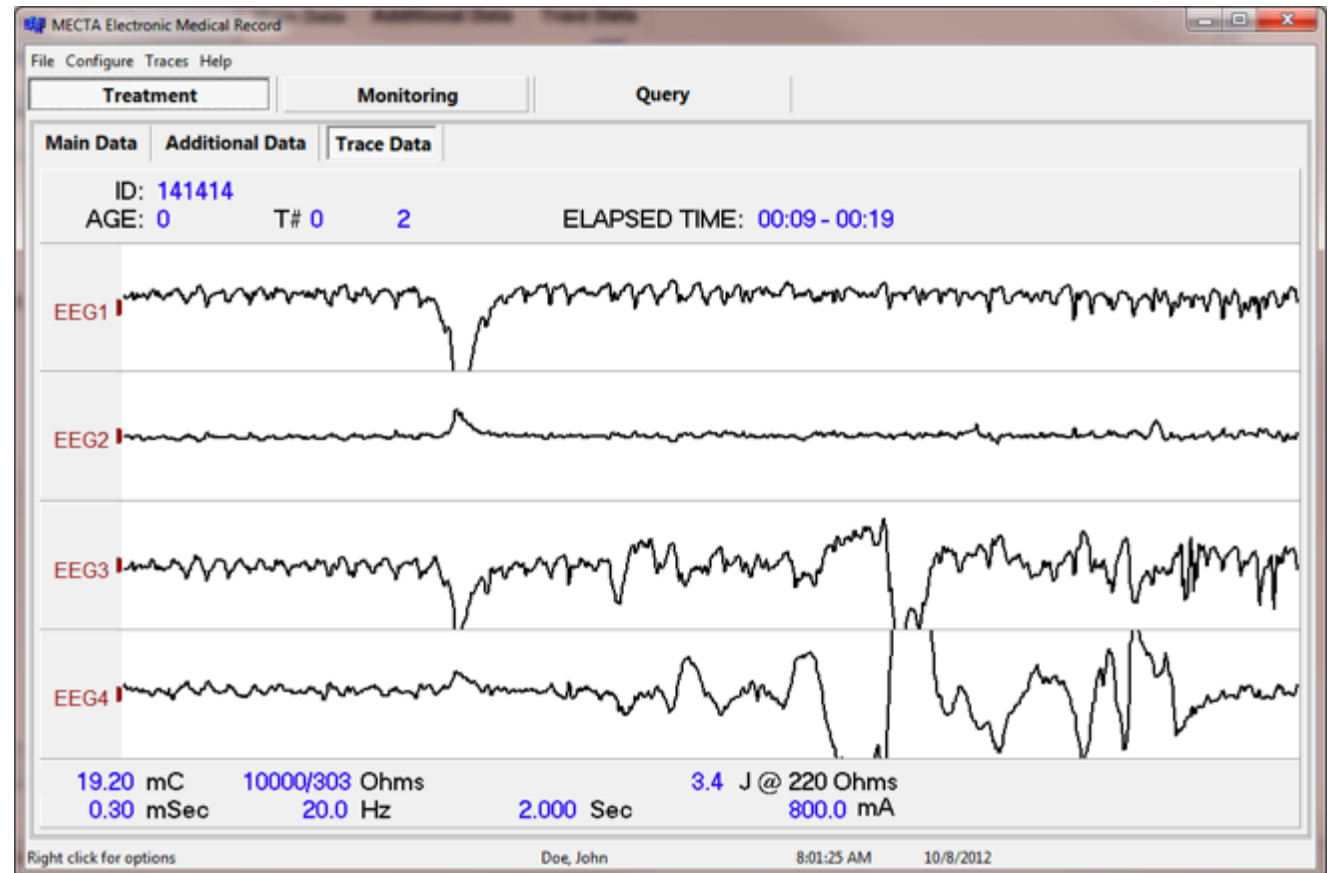
MECTA, EEG papier



MECTA, EEG numérisisé



MECTA, EEG numérisé



MECTA, EEG numérisisé



Charle perrens

STIMULUS and MONITORING REPORT

Patient
Patient ID
Sex

MRN
Encounter

Date	24/03/2017	Time	08:22:33	Course	1	Session	1	Treatment Number	1		
PW	,5	Freq	70	Dur	7,5	Current	900	Charge	472,5	Energy	69,2
Start	0:00									End	0:09
EEG1											
EEG2											
Start	0:09									End	0:18
EEG1											
EEG2											
Start	0:18									End	0:27
EEG1											
EEG2											
Start	0:27									End	0:37
EEG1											
EEG2											



MECTA, EEG indice de « seizure adequacy »



Analyse des données EEG : l'index « Seizure Adequacy » a été démontré comme pertinent pour les cliniciens. Basé sur 10 années de recherche et d'analyse partagée au sein de la Duke University, il s'agit d'un index efficace breveté, sous la licence exclusive de Mecta.

Krystal AD, Weiner RD.
ECT seizure therapeutic adequacy.
Convuls Ther. 1994 Jun;10(2):153-64.

MECTA, EEG indice de « seizure adequacy »



US005626627A

United States Patent [19]
Krystal et al.

[11] **Patent Number:** 5,626,627
[45] **Date of Patent:** May 6, 1997

[54] **ELECTROCONVULSIVE THERAPY METHOD USING ICTAL EEG DATA AS AN INDICATOR OF ECT SEIZURE ADEQUACY**

[75] Inventors: **Andrew D. Krystal; Richard D. Weiner**, both of Durham, N.C.

[73] Assignee: **Duke University**, Durham, N.C.

[21] Appl. No.: **508,062**

[22] Filed: **Jul. 27, 1995**

[51] Int. Cl.⁶ **A61N 1/32**

[52] U.S. Cl. **607/45; 128/731**

[58] Field of Search **128/731; 607/45**

[56] References Cited

U.S. PATENT DOCUMENTS

4,777,952	10/1988	Pavel	128/419
4,870,969	10/1989	Swartz	128/419
4,873,981	10/1989	Abrams et al.	128/419
4,878,498	11/1989	Abrams et al.	128/419
5,269,302	12/1993	Swartz et al.	128/419

OTHER PUBLICATIONS

Andrew D. Krystal et al., "ECT Seizure Therapy Adequacy" *Convulsive Therapy*, vol. 10, No. 2, pp. 153-164 (1994).
Conrad Melton Swartz, "Beyond Seizure Duration as a Measure of Treatment Quality" *Convulsive Therapy*, vol. 9, No. 1, pp. 1-7 (1993).
Nobler et al., "EEG Manifestations during ECT: Effects of Electrode Placement and Stimulus Intensity". *Biol. Psychiatry*, vol. 34, pp. 321-330 (1993).
Andrew D. Krystal et al., "The Largest Lyapunov Exponent of the EEG in EOT Seizures". *Proceedings of the Conference on Measuring Chaos in the Human Brain*, World Scientific Publishing Co. pp. 113-127 (1991).

Krystal et al. "EEG Evidence of More Intense Seizure Activity with Bilateral ECT". *Biol. Psychiatry*, vol. 31, pp. 617-621 (1992).

Krystal et al. "The Effects of ECT Stimulus Dose and Electrode Placement on the ictal Electroencephalogram: An Intraindividual Crossover Study" *Biol Psychiatry*, vol. 34, pp. 759-767 (1993).

Weiner et al., "The Monitoring and Management of Electrically Induced Seizures", *Psychiatric Clinics of North America*, vol. 14, No. 4, (Dec. 1991).

Weiner et al. "EEG Monitoring of ECT Seizures". *The Clinical Science of Electroconvulsive Therapy*, Washington, D.C., American Psychiatric Press, Inc. pp. 93-109, (1993).

Conrad Melton Swartz, "Low-Frequency Ictal EEG Activity and ECT Therapeutic Impact". *Convulsive Therapy*, vol. 9, No. 3, pp. 220-224 (1993).

Primary Examiner—William E. Kamm

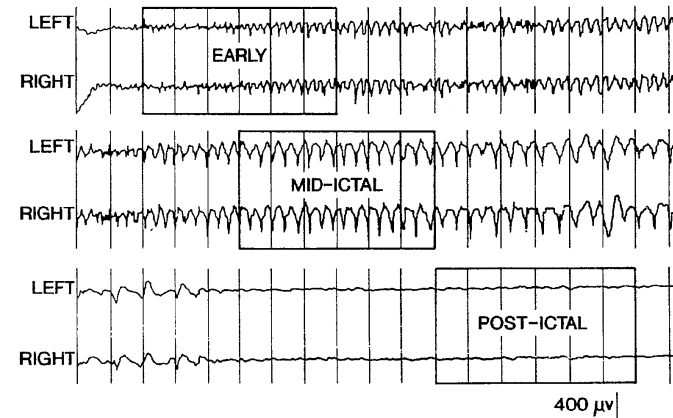
Attorney, Agent, or Firm—Richard E. Jenkins, P.A.

[57]

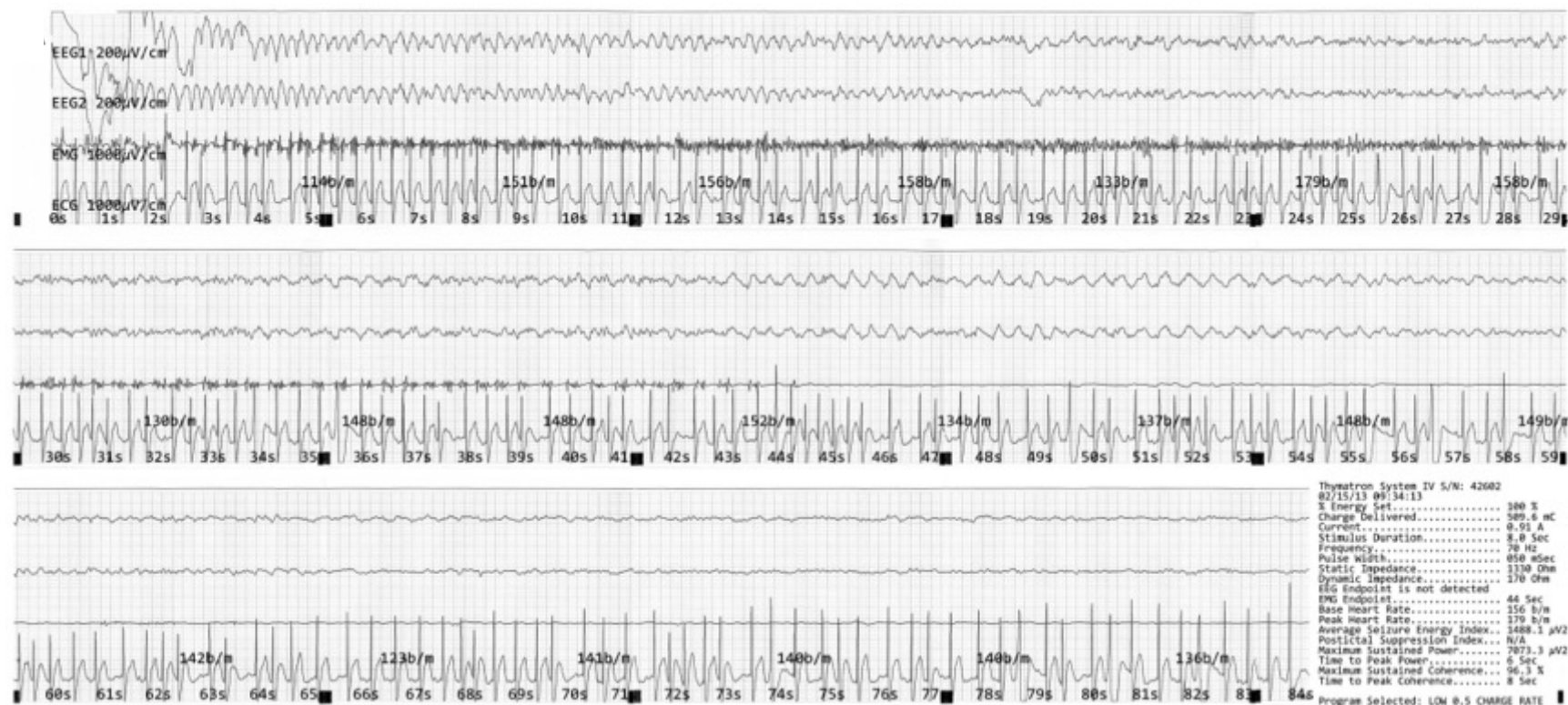
ABSTRACT

A method in electroconvulsive therapy (ECT) to use ictal EEG data for clinical determination of the adequacy of an induced seizure in a patient. The method includes employing an ECT device to apply electricity to the patient in an ECT session to induce seizure activity. The electroencephalographic (EEG) data is detected during the seizure and selected EEG data parameters are derived therefrom. Next, the likely adequacy of the induced seizure is computed by comparing the selected EEG data parameters of the patient to ictal EEG data parameters wherein the adequacy of the corresponding seizure or seizures is known, and the computed likely therapeutic adequacy of the induced seizure is displayed.

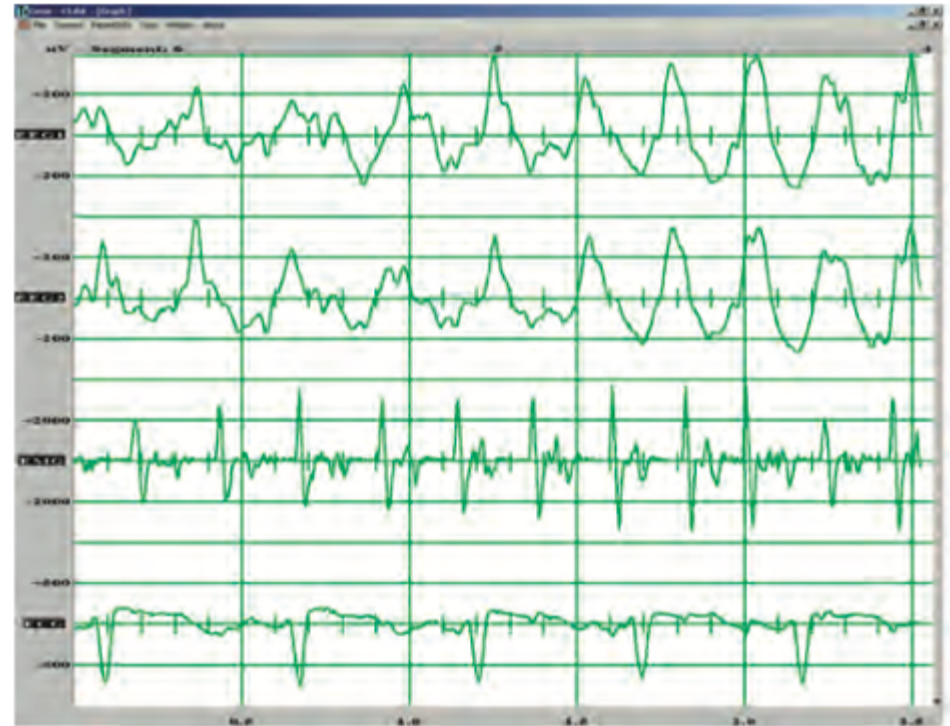
23 Claims, 1 Drawing Sheet



THYMATRON, EEG papier



THYMATRON, EEG numérisisé



GENIE™ IV

EEG Analysis Software for the Thymatron® System IV

THYMATRON, EEG numérisé



THYMATRON, EEG numérisé



SEIZURE MONITORING

The Thymatron® System IV allows the physician to monitor the physiological variables of EEG, ECG, and EMG. The paper tracing provides the wave forms and beats per minute for the ECG. The EEG and EMG also appear on the tracings, with additional information provided.

EEG SEIZURE MONITORING

- 1) The Audible EEG™ seizure monitor
- 2) The EEG paper recording
- 3) The Ictal Line™ seizure indicator
- 4) The EEG endpoint and indices determined values

Audible EEG™ Seizure Monitor

This feature operates automatically when the "TREAT" button is pressed and released. The knob marked "VOLUME" on the back panel controls the volume of the tone. To inactivate this feature, turn the volume control knob all the way counterclockwise.

The pitch of the Audible EEG™ signal varies with the amplitude of the EEG. It will waver and warble intensely and rapidly during the initial tonic phase. It becomes increasingly irregular, with superimposed staccato bursts, during the clonic phase, and tends to correspond to each muscular contraction. Seizure termination is marked by a change to a nearly steady tone with little modulation or variability. Each Thymatron® System IV is supplied with a cassette tape guide for the interpretation of the Audible EEG™ seizure monitor.

EEG paper recording

- a) Paper EEG recording prior to the treatment stimulus can be initiated (after the EEG electrodes have been properly applied) by pressing the "START/STOP" button on the front panel. This will provide a paper record of the patient's baseline recording. The printer stops during treatment stimulus administration.
- b) Automatic paper EEG recording begins or resumes when the treatment stimulus ends. The EEG recording continues through ictal and postictal periods, until the "START/STOP" button is pressed, which generates the end-of-treatment report.

NOTE: Obtaining a paper baseline EEG record does not replace the baseline EEG collection procedures described above.

Ictal Line™ Seizure Indicator

After baseline EEG collection is completed by the Thymatron® System IV and the "READY" message light appears, a thin black line is printed along the top of the paper recording strip when the EEG amplitude exceeds a specified baseline value. An unbroken, solid black line reflects continuous seizure activity. A broken or intermittent line reflects waxing and waning, or intermittent seizure activity. Complete cessation of the black line reflects EEG seizure termination, as determined by the Thymatron® System IV. Wait several seconds before pressing the "START/STOP" button to terminate recording because the computer takes that long to process and report the seizure endpoints and indices.

Endpoints and Indices

A unique feature of the Thymatron® System IV, (U.S. patents: 4873981, 4878498, 5269302 and 5871517), provides two computer-determined estimates of the duration of the induced seizure, derived from the EEG and EMG data.

Automatic EEG Seizure Endpoint Determination

The Thymatron® System IV continuously monitors the EEG for the endpoint of seizure activity and prints the EEG seizure duration, in seconds, on the end-of-treatment report, provided the baseline EEG collection procedures have been properly followed and the "READY" message has appeared. (If the treatment stimulus is administered before the "READY" message appears, automatic EEG analysis will not occur and the end-of-treatment report will state the message "Baseline not available.")

In about 10-20% of ECT treatments, the EEG endpoint is not readily determined (Abrams, 1997). This typically occurs when paroxysmal activity decreases too gradually to provide a clear visual endpoint, or when the immediate post-seizure EEG contains high amplitude activity. In these circumstances, inability to detect a precise EEG endpoint is expected with any method of examination. The Ictal Line™ might show an on-again-off-again broken line pattern, and the end-of-treatment report might state: "EEG Endpoint is not detected".

Automatic EMG (Motor) Seizure Endpoint Determination

The Thymatron® System IV is shipped with the EMG monitor enabled in channel A. When EMG monitoring electrodes have been properly applied, the lead-wires and monitoring cable connected, then EMG tracing automatically appears on the paper record after the treatment stimulus ends.

The Thymatron® System IV continuously monitors the EMG for motor seizure activity and prints the EMG endpoint seizure duration in seconds, on the end-of-treatment report. Baseline EMG collection is neither required, nor possible, in obtaining this measure.

CAUTION: The computer-derived endpoint seizure duration measures, including the Ictal Line™ seizure indicator, are derived solely by calculation and are provided to aid, not replace, the physician's judgment. It is possible for seizure activity to continue in the brain after any or all of the computer reports indicate seizure termination. It is also possible for artifact to be interpreted by the computer programs as seizure activity.

THYMATRON, EEG indices



- **EEG COHERENCE MEASURES** of maximum sustained coherence, and time to peak coherence, interhemispheric cross-correlation measures reported to reflect seizure quality and clinical impact (Krystal & Weiner, 1994; Krystal et al, 1995; Krystal, 1998).
- **EEG AMPLITUDE** measures of maximum sustained EEG power, and average seizure energy, with separate values for early, mid- and postictal seizure phases, found by the Duke University group to be important correlates of seizure quality and efficacy (Krystal & Weiner, 1994; Krystal et al, 1995; Krystal, 1998).
- The **POSTICTAL SUPPRESSION INDEX** reports the degree of EEG flattening immediately following the seizure, which correlates with clinical efficacy (Nobler et al, 1993; Krystal & Weiner, 1994; Krystal et al, 1995; Krystal, 1998; Nobler et al, 2000). A recent study of the Thymatron's Postictal Suppression Index found that it significantly differentiated ECT remitters from non-remitters (Petrides et al, 2000). The authors concluded: "higher PSI values (more abrupt ending of ictal EEG) are correlated with better clinical outcome of ECT in depression".

Krystal AD, Weiner RD (1994): ECT seizure therapeutic adequacy. *Convul. Ther.* 10:153-164.

Krystal et al (1995): The ictal EEG as a marker of adequate stimulus intensity with unilateral ECT. *J. Neuropsych.* 7:295-303.

Krystal AD (1998): The clinical utility of ictal EEG seizure adequacy models. *Psych. Ann.* 28:30-35.

Nobler MS et al (1993): EEG manifestations during ECT: Effects of electrode placement and stimulus intensity. *Biol. Psych.* 34:321-330.

Nobler MS et al (2000): Quantitative EEG during seizures induced by electroconvulsive therapy: relations to treatment modality and clinical features. I. Global analyses. *J ECT* 16:211-28.

Petrides G, Kellner C et al (2000): Can Ictal EEG Indices predict response to ECT? [presentation] Jan. 2000 NCDEU meeting.

THYMATRON, EEG numérisisé



SEIZURE QUALITY MEASURES

The Thymatron® System IV provides 7 Seizure Quality Measures under the “INDEXES” function level that can be individually enabled/disabled. EEG monitoring must be enabled to obtain these measures. Their names and FlexDial™ designations are as follows:

<i>Average Seizure Energy Index</i>	ASEI ON/OFF
<i>Postictal Suppression Index</i>	PSI ON/OFF
<i>Maximum Sustained Power and Time to Peak Power</i>	MSP ON/OFF
<i>Maximum Sustained Coherence and Time to Peak Coherence</i>	COH ON/OFF
<i>Duke University Amplitude Measures</i>	DUKE ON/OFF

The AVERAGE SEIZURE ENERGY INDEX (ASEI) integrates the total ictal EEG power across the entire seizure and divides the result by the total seizure duration.

The POSTICTAL SUPPRESSION INDEX (PSI) measures the percentage decrease in ictal EEG amplitude immediately following seizure termination.

The MAXIMUM SUSTAINED POWER (MSP) measure reports the mean value of the 10-second EEG segment with the highest average power recorded during the seizure.

TIME TO PEAK POWER is the time elapsed from stimulus termination to the point of maximum EEG power.

The MAXIMUM SUSTAINED COHERENCE (COH) measure reports the mean value of the 5-second EEG segment with the highest average coherence recorded

TIME TO PEAK COHERENCE is the time elapsed from stimulus termination to the point of maximum EEG coherence.

DUKE UNIVERSITY AMPLITUDE MEASURES display the amplitudes of the 3 seizure segments (early ictal, mid-ictal, and post-ictal) reported by Duke University investigators to correlate with ECT treatment response.

BASELINE RETENTION

Baseline Retention, BLV, found under ENDPOINTS, is the length of time the EEG baseline will be kept in memory after the treatment. It can be set from 0 - 5 minutes. After this time a new EEG baseline must be acquired. This feature is useful for re-stimulation without acquiring a new EEG baseline.

Approche clinique

La première utilité de l'EEG pendant les ECT :

Evaluer la durée de la crise

Pourquoi ?

Crise prolongé



A court terme

ETAT DE MAL EPILEPTIQUE



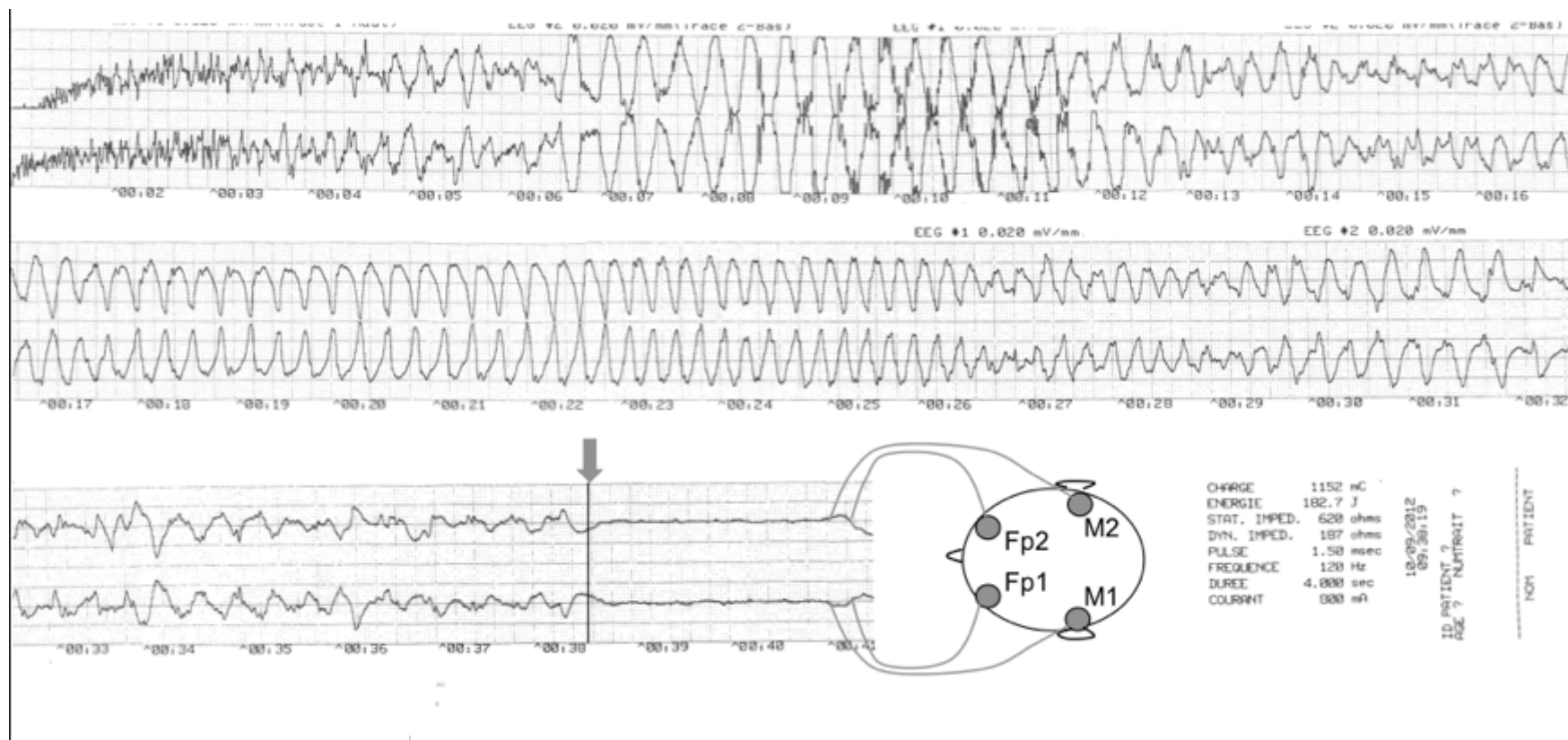
A moyen terme

MAUVAISE TOLERANCE COGNITIVE



Meilleure efficacité

Fixer la durée de la crise induite



Arrêt des signes cliniques fins avant les signes EEG

La surveillance EEG
des séances d'ECT
permet de repérer une
crise épileptique prolongée (>3min, définition APA)

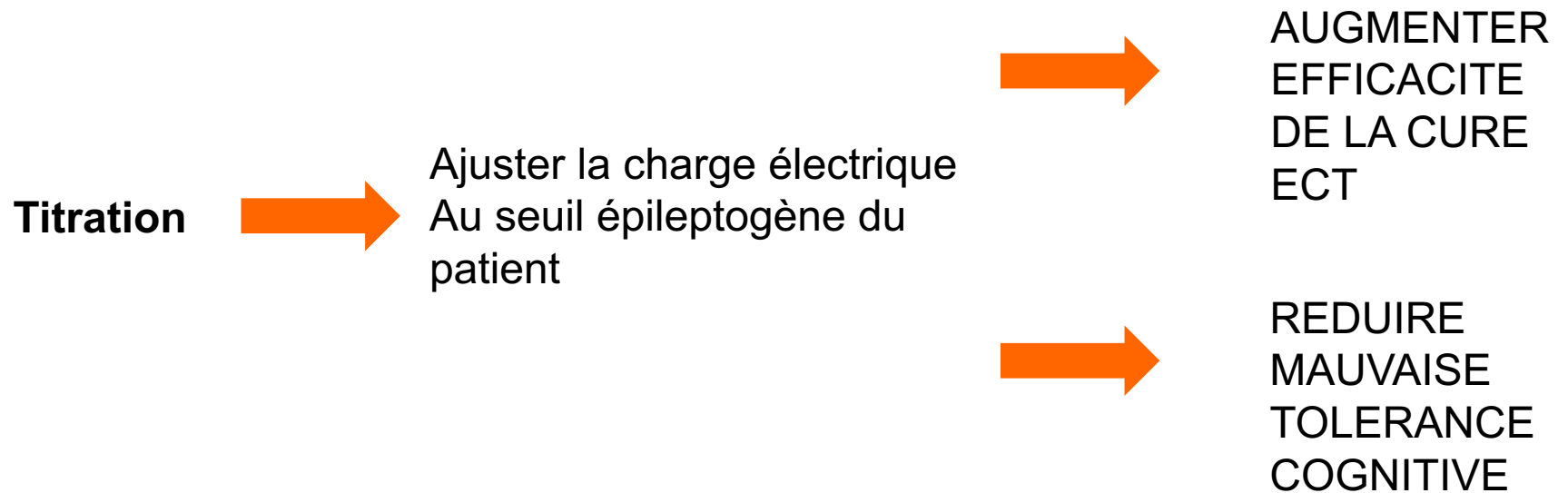
Difficultés pour déterminer la durée de la crise

1. Diminution progressive de l'activité de la phase critique et continuité avec la phase post critique
2. Suppression post critique incomplète
3. Fin de crise artefactée



1. Laisser défiler le papier 10 secondes
(*when in doubt, play it out*)
2. Période de *hand-off*
3. Relire le tracé en partant de la fin

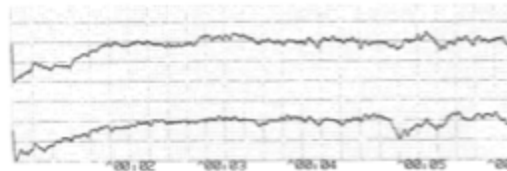
Pourquoi ?



Fixer le seuil épileptogène pour un patient

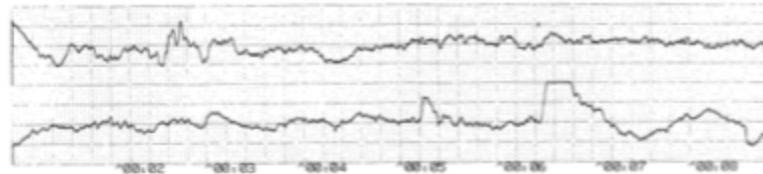
Enregistrer 15-20 secondes
Car possible **crise**
retardée

CHARGE 72 mC
ENERGIE 12.7 J
at 220 ohms
IMPEDANCE 740 ohms
PULSE 1.00 msec
FREQUENCE 30 Hz
DUREE 1.500 sec
COURANT 600 mA



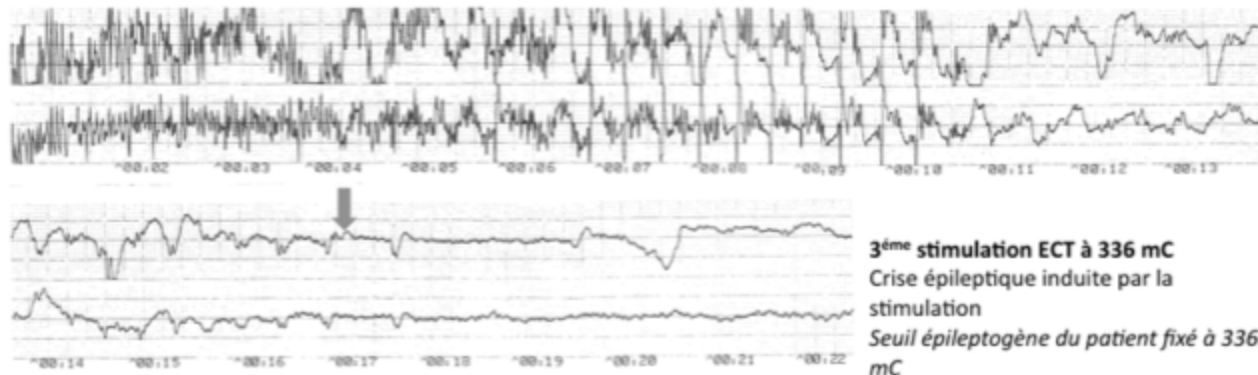
1^{er} stimulation ECT à 72 mC
Au dessous du seuil épileptogène

CHARGE 120 mC
ENERGIE 22.5 J
at 220 ohms
IMPEDANCE 500 ohms
PULSE 1.00 msec
FREQUENCE 40 Hz
DUREE 2.000 sec
COURANT 600 mA



2^{ème} stimulation ECT à 120 mC
Au dessous du seuil épileptogène

CHARGE 336 mC
ENERGIE 59.1 J
at 220 ohms
IMPEDANCE 440 ohms
PULSE 1.00 msec
FREQUENCE 70 Hz
DUREE 3.000 sec
COURANT 600 mA



3^{ème} stimulation ECT à 336 mC
Crise épileptique induite par la
stimulation
Seuil épileptogène du patient fixé à 336
mC

La surveillance EEG
des séances d'ECT
permet de réaliser la titration de la charge ECT

La deuxième utilité de l'EEG pendant les ECT :

Evaluer la qualité de la crise

Pourquoi ?

**Crise de durée
adéquate**



Crise optimale



AUGMENTER
EFFICACITE
DE LA CURE
ECT

Pourquoi ?

**Au fur et à mesure
des séances :
le seuil
épileptogène des
patients
augmentent**



**Diminution de
la qualité de
crise**



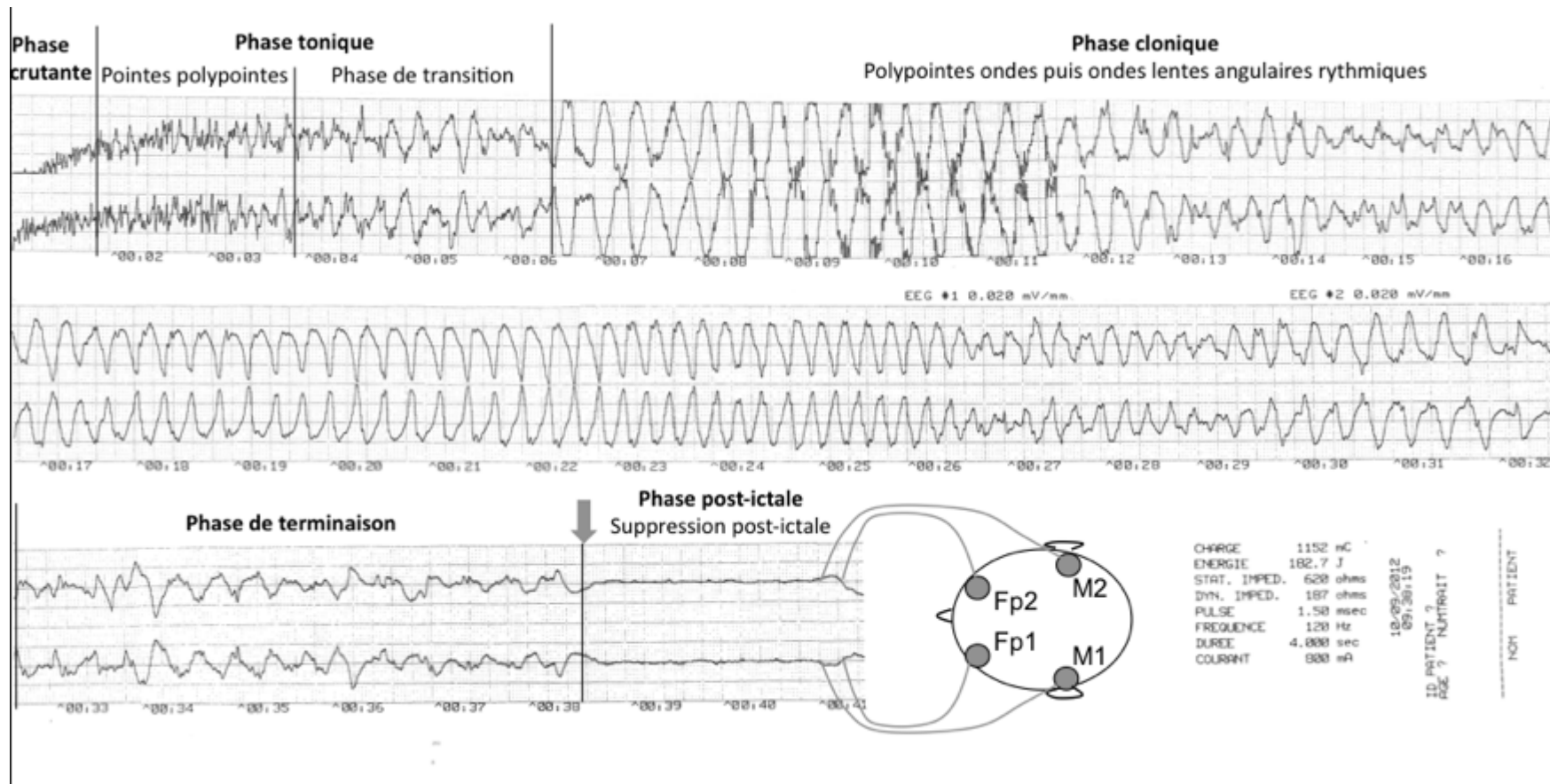
**DIMINUTION
EFFICACITE
DE LA CURE
ECT**

**Ajustement des paramètres de stimulation
au cours des séances ECT
(et après la séance de titration)**

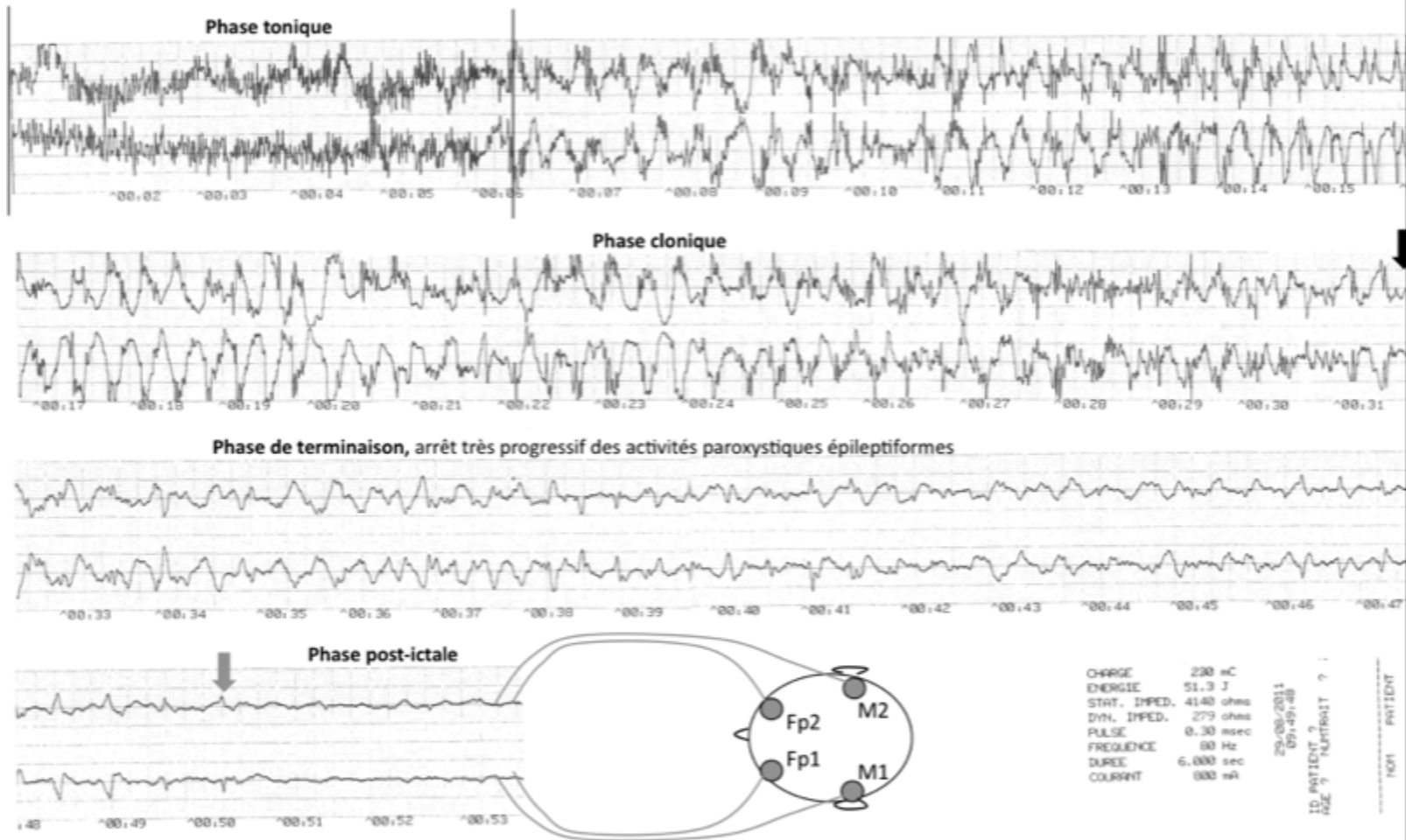
La surveillance EEG
des séances d'ECT
permet de repérer une
crise épileptique avortée (<15-20 s)
crise épileptique adéquate (>20s & <3min)
crise épileptique optimale

Permet d'adapter les paramètres de stimulation pour
optimiser l'efficacité des ECT

Savoir lire la trace EEG



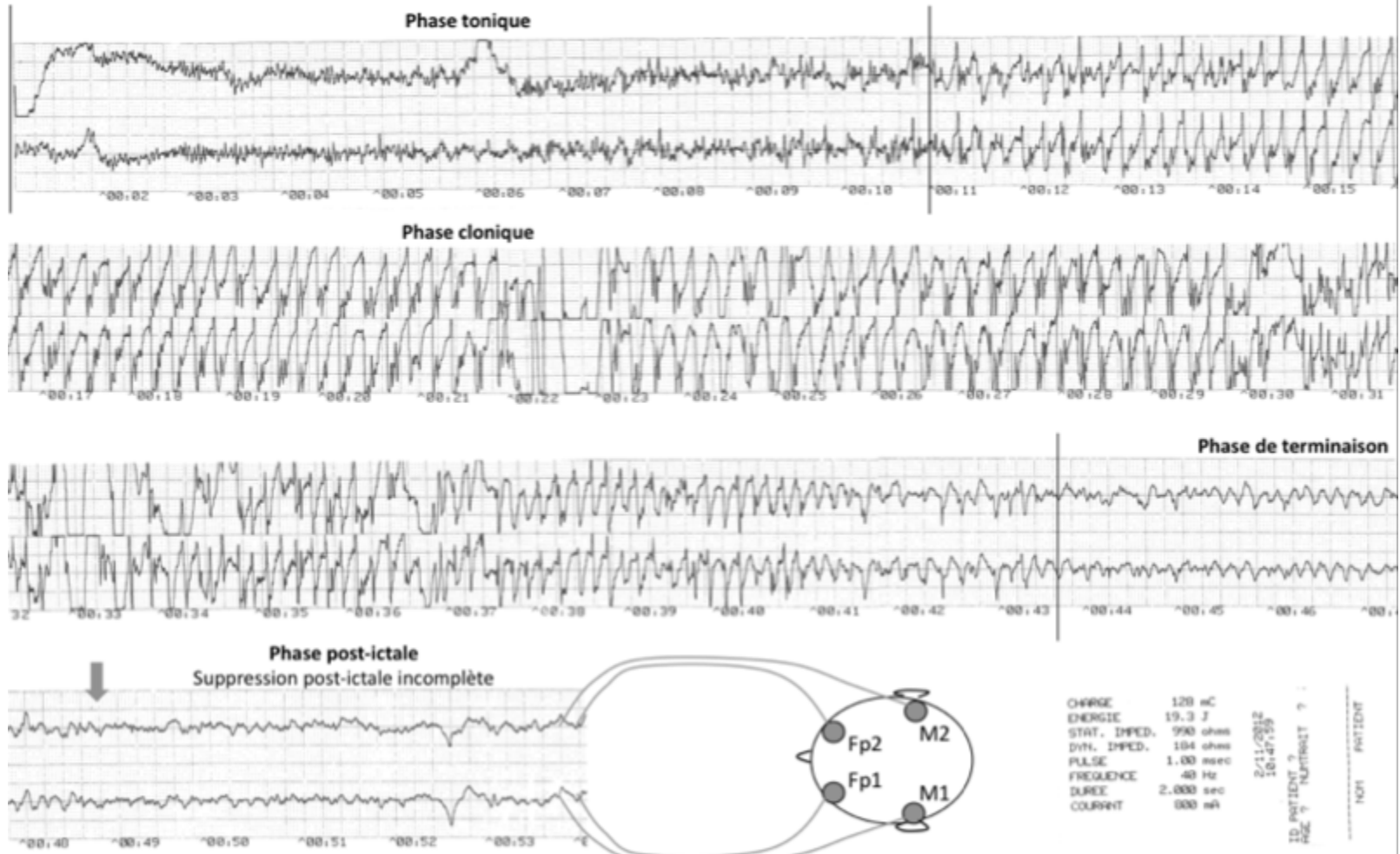
Savoir lire la trace EEG



« When in doubt, play it out »

Poursuivre l'EEG 10 secondes après la fin de la crise
(pour s'assurer de l'absence de reprise de crise)

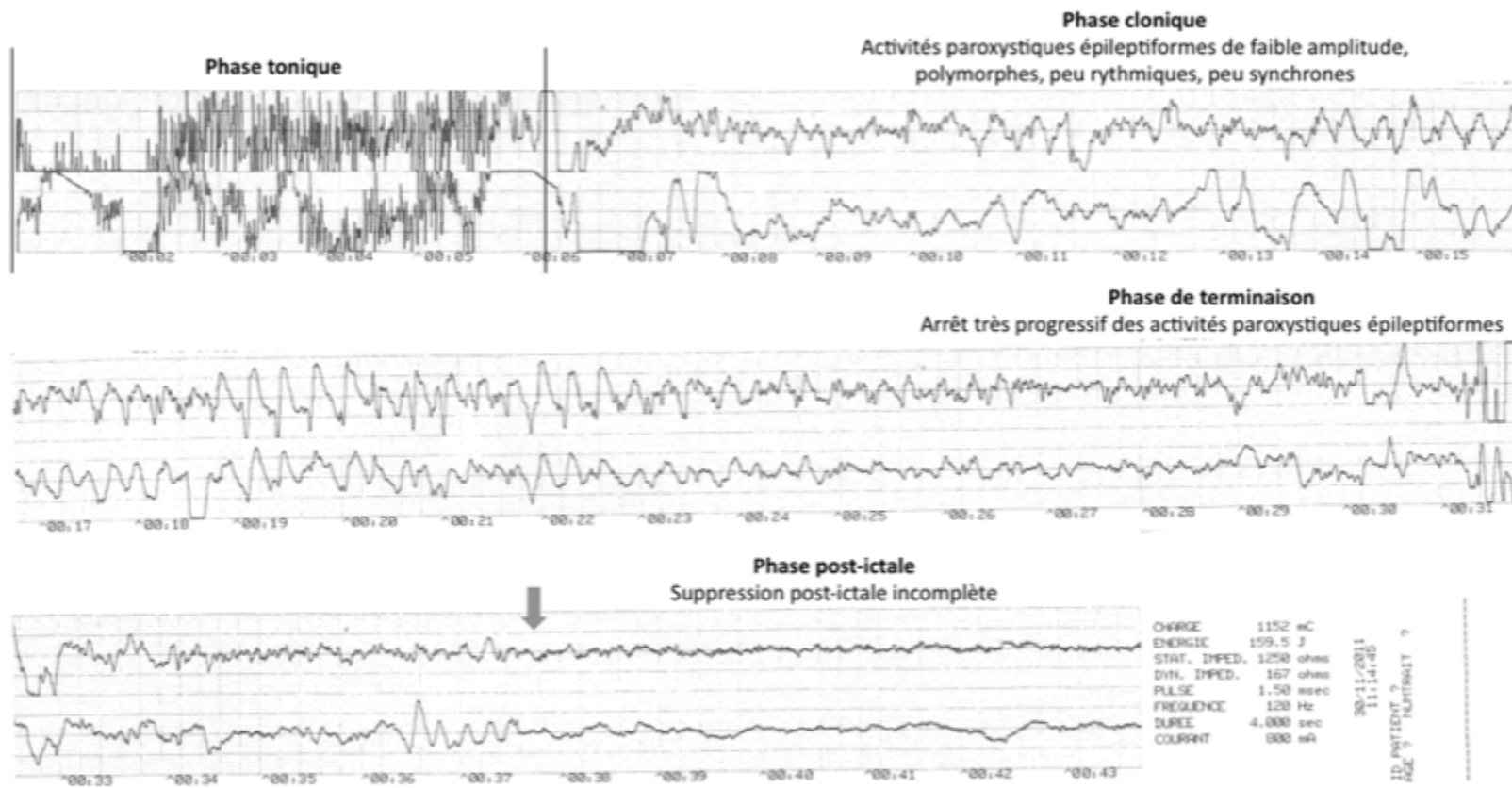
Savoir lire la trace EEG



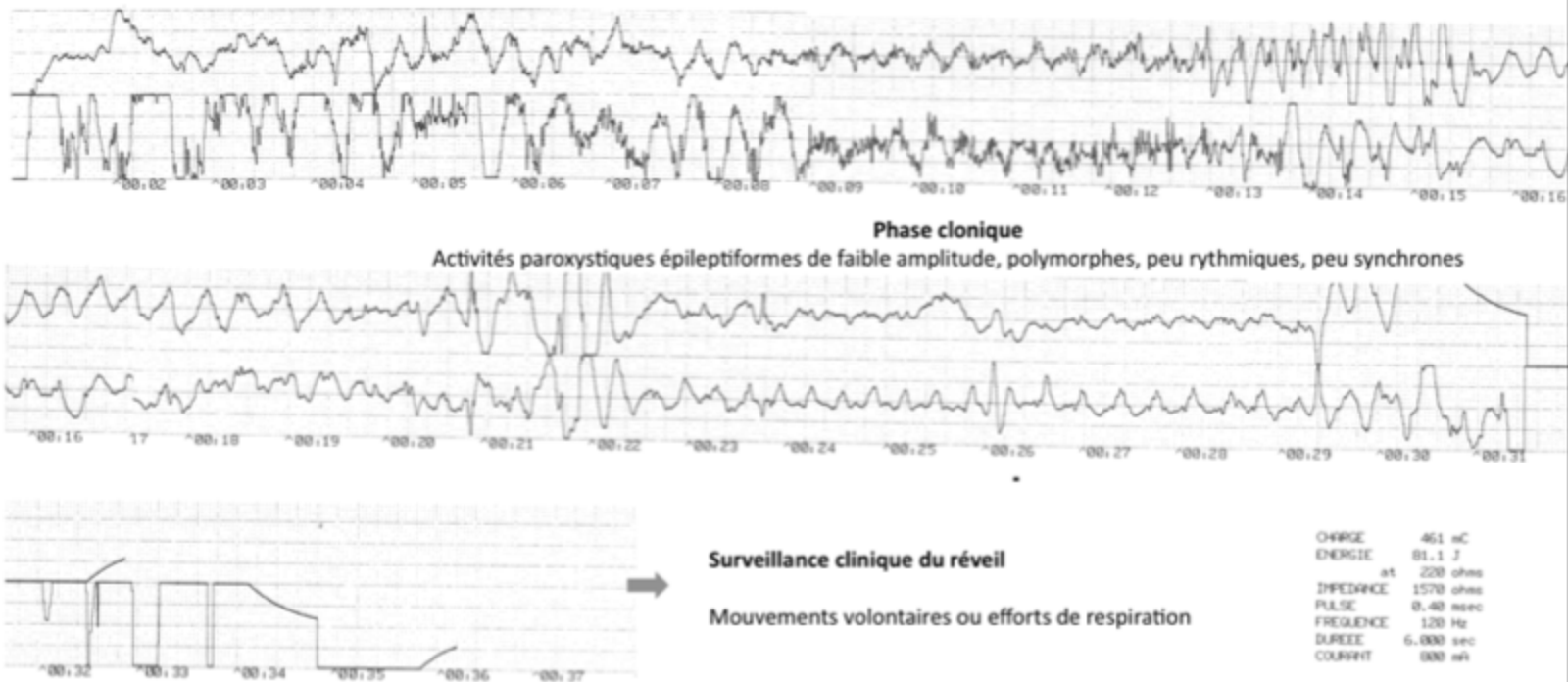
« When in doubt, play it out »

Période de hand off et s'assurer de la reprise de mouvements volontaires et d'efforts respiratoires

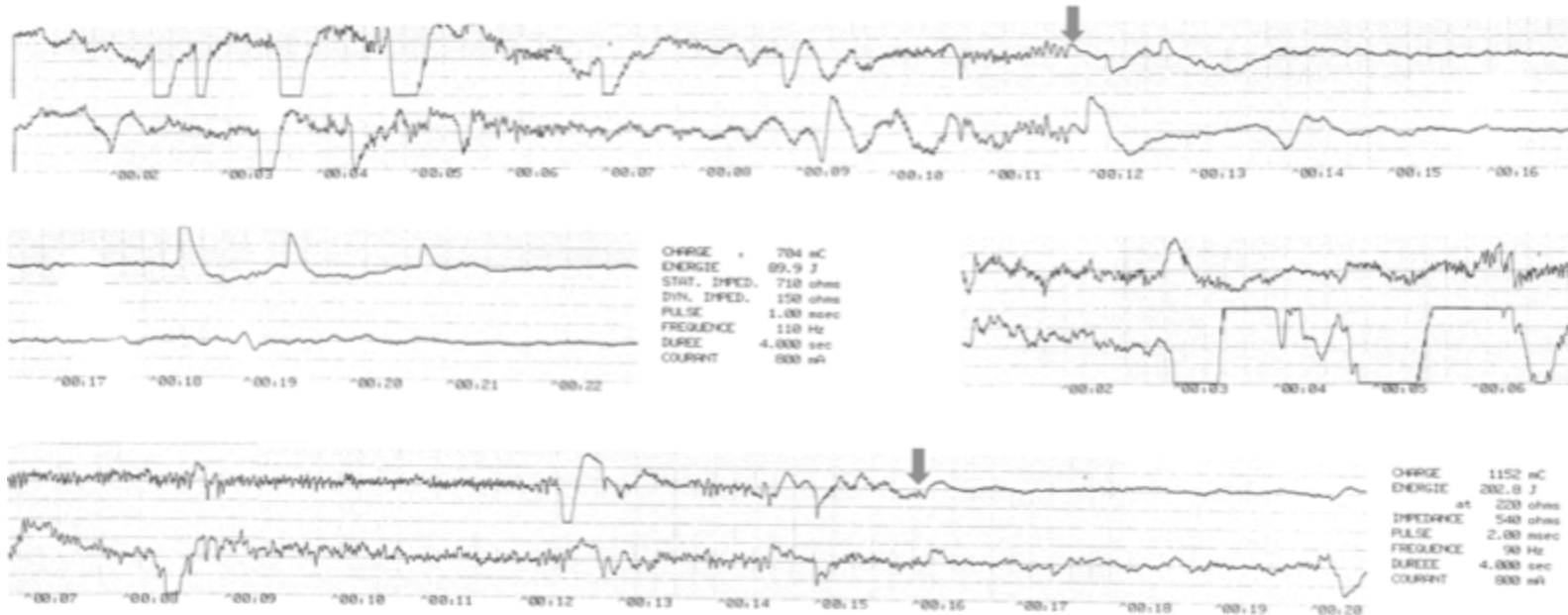
Savoir lire la trace EEG



Savoir lire la trace EEG



Savoir lire la trace EEG

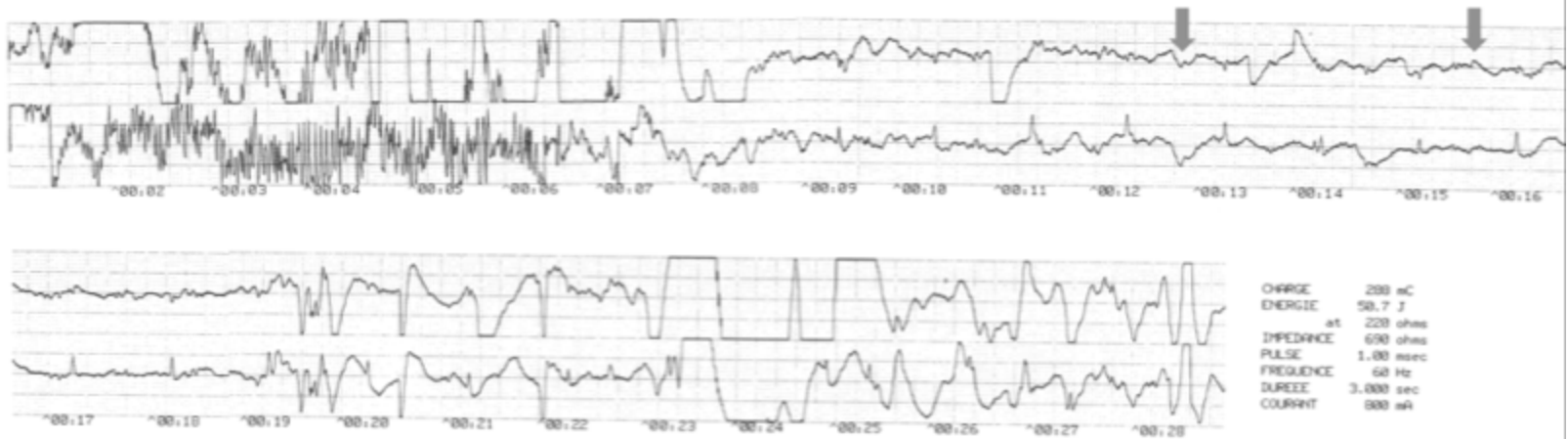


Rechercher des facteurs favorisant

Antiépileptiques?

Préférer l'étomidate (hypnomidate®) au propofol (diprivan®)

Savoir lire la trace EEG





L'échelle

université
de **BORDEAUX**

A la quête de l'échelle électrophysiologique

Brain Stimulation 9 (2016) 72–77



Contents lists available at ScienceDirect

Brain Stimulation

journal homepage: www.brainstimjrnل.com



The Anaesthetic-ECT Time Interval in Electroconvulsive Therapy Practice – Is It Time to Time?

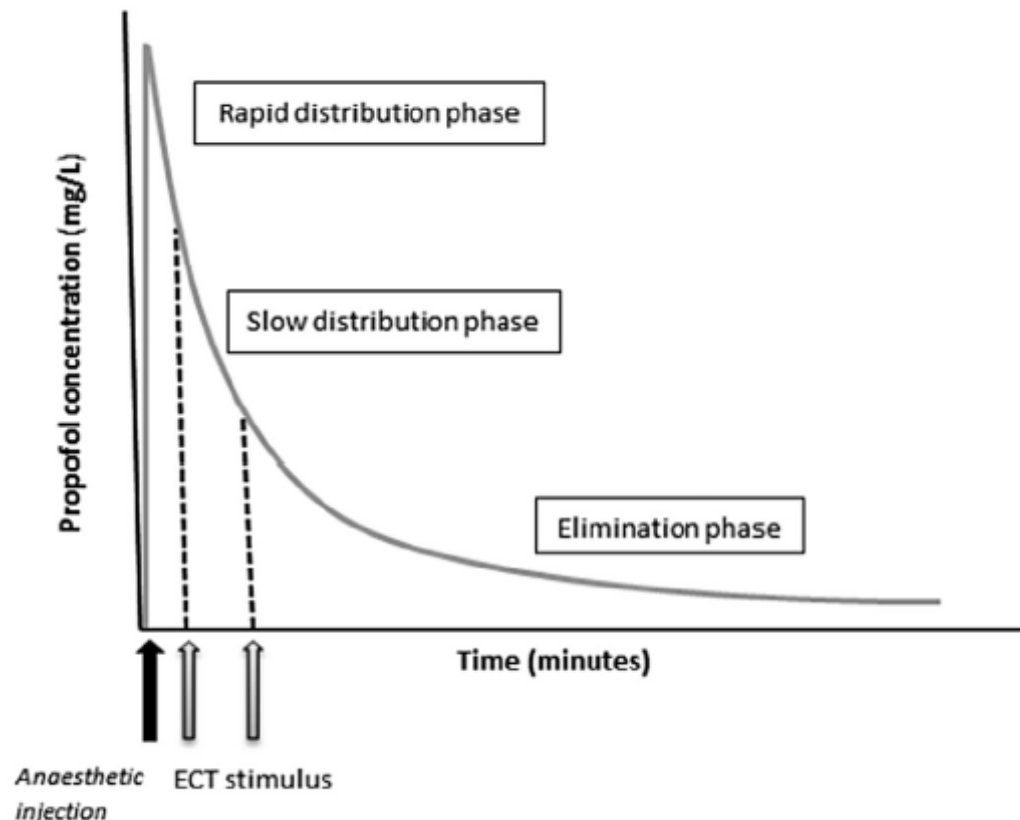


Verònica Gálvez ^{a,b,c}, Dusan Hadzi-Pavlovic ^{a,b}, Harry Wark ^{a,c,d,e}, Simon Harper ^{c,f}, John Leyden ^{c,g}, Colleen K. Loo ^{a,b,c,h,*}

A la quête de l'échelle électrophysiologique

Results: Longer anaesthetic-ECT time intervals lead to significantly higher quality seizures ($p < 0.001$ for amplitude, regularity, stereotypy and post-ictal suppression).

Conclusions: These results suggest that the anaesthetic-ECT time interval is an important factor to consider in ECT practice. This time interval should be extended to as long as practically possible to facilitate the production of better quality seizures. Close collaboration between the anaesthetist and the psychiatrist is essential.



A la quête de l'échelle électrophysiologique

Seizure quality EEG indices [TSLOW (time to onset of seizure activity ≤ 5 Hz, seconds); peak mid-ictal amplitude (mm); regularity (e.g. intensity or morphology of the seizure (0–6)); stereotypy (or global seizure patterning, 0–3) and post-ictal suppression (0–3)], as well as seizure duration (in seconds) were rated through a qualitative–quantitative structured rating scale [33], based on the one developed by Nobler [17,34]. Ratings were performed by a single



Brief report

Low dose lignocaine added to propofol does not attenuate the response to electroconvulsive therapy

Ross D. MacPherson^{a,*}, Jessica Lawford^b, Brett Simpson^c, Michelle Mahon^d,
Debra Scott^e, Colleen Loo^f

A la quête de l'échelle électrophysiologique

EEG traces were examined for EEG seizure duration and ictal quality. The latter was rated manually according to five criteria previously found to be related to the efficacy of ECT seizures (Krystal et al., 1993, 1998; McCall et al., 1996; Nobler et al., 1993; Weiner and Krystal, 1993). A manual system of rating these criteria was developed, based on descriptions provided in the above studies, with further specific guidelines as required (see [Appendix A](#)).



Appendix A. EEG Seizure Quality Rating Sheet

Parameter	Score/value
Time to onset of slowing (TSLOW) • Time from end of ECT stimulus to slowing of frequency to ≤ 5 Hz. • Score as 0 if frequency does not initially exceed 5 Hz.	[secs]
Peak mid-ictal amplitude • Calculate the mean value of 3 consecutive spike and wave complexes which represent the largest of the slow wave/spike-and-wave phase (ignore transition phases).	Wave 1: [mm] Wave 2: [mm] Wave 3: [mm] Mean:
Seizure regularity (0-6) • Score the predominant pattern during slow wave/spike-and-wave phase according to the scale (ignore transition phases). See scale by Krystal and Weiner, (1993).	
Seizure stereotypy (0-3; 0.5 pt increments) • Score based on three characteristics. Score 0, 0.5 or 1 for each and sum the total. 1. Transition from recruitment polyspike phase to the slow wave/spike-and-wave phase (0=no recruitment phase, 1=recruitment phase present and transition to slow wave is clear). Consider transition in terms of both morphology and increase in amplitude. 2. Quality of the spike-and-wave morphology and reappearance of chaotic polyspikes during the slow wave/spike-and-wave phase (0=no clear spike-and-wave and/or chaotic polyspikes are present during the slow wave/spike-and-wave phase; 1=a stereotypic slow wave/spike-and-wave phase is present, and no reappearance of chaotic polyspikes). 3. Variability in amplitude of the slow wave/spike-and-wave phase (0=marked variability, the amplitude is inconsistent for >50% of the slow wave phase; 1=the amplitude is consistent for >80% of the slow wave phase) NT: this is excluding transition phases.	
Post-ictal suppression (0-3; 0.5 pt increments) • Score based on three characteristics. Score 0, 0.5 or 1 for each and sum the total. 1. Seizure end-point (0=cannot determine end-point; 1=clear end-point). 2. Transition to seizure termination (0=gradual, the transition phase consists of >1/3 of the trace; 1=abrupt). 3. Flatness of trace after seizure end-point (0=poor; 1=very flat).	

A la quête de l'échelle

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A la quête de l'échelle

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Krystal, A., Weiner, R., McCall, V., Shelp, F., Arias, R., Smith, P., 1993. The effect of ECT stimulus dose and electrode placement on the ictal electroencephalogram: an intraindividual crossover study. *Biol. Psychiatry* 34, 759–767.

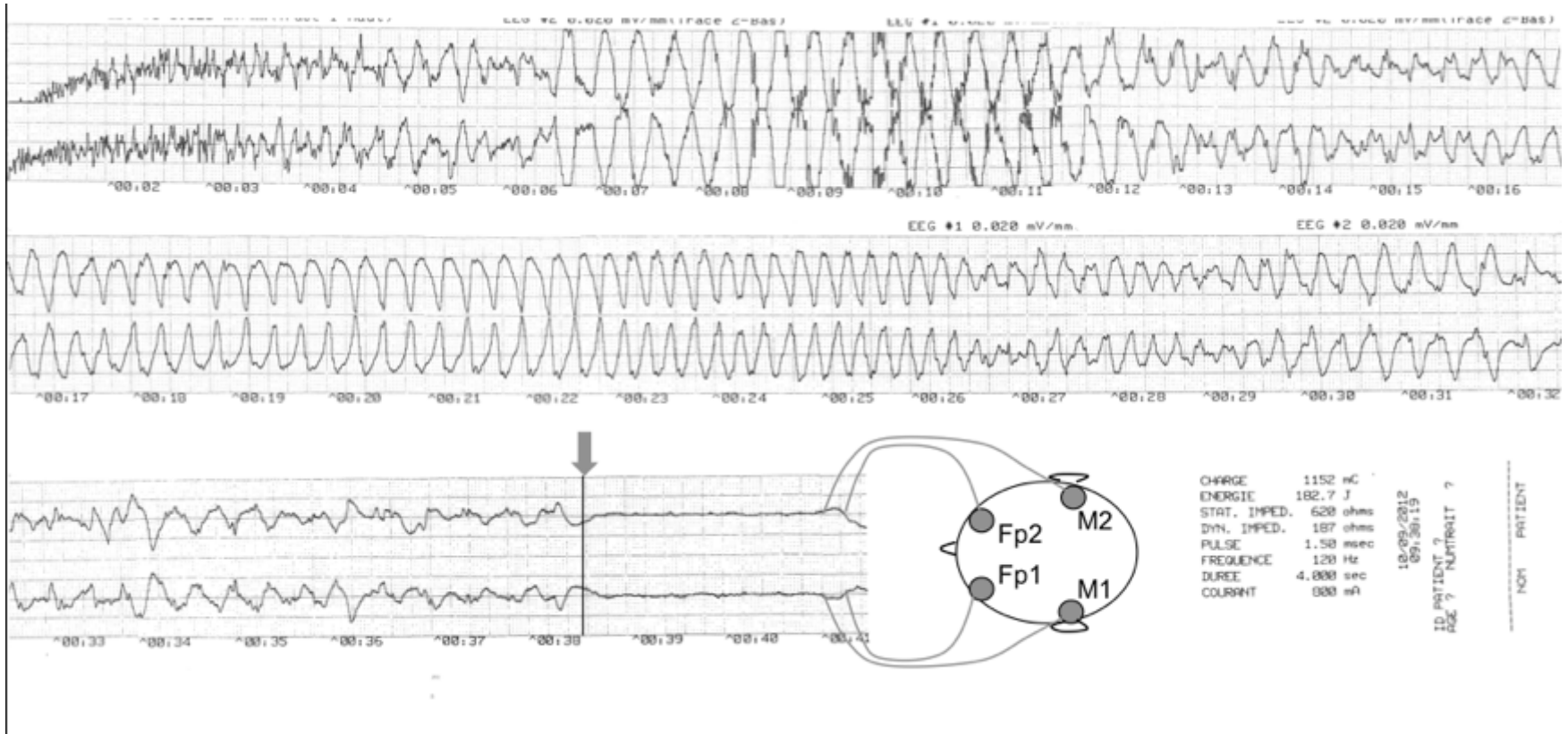
La compréhension de l'échelle

Appendix A. EEG Seizure Quality Rating Sheet

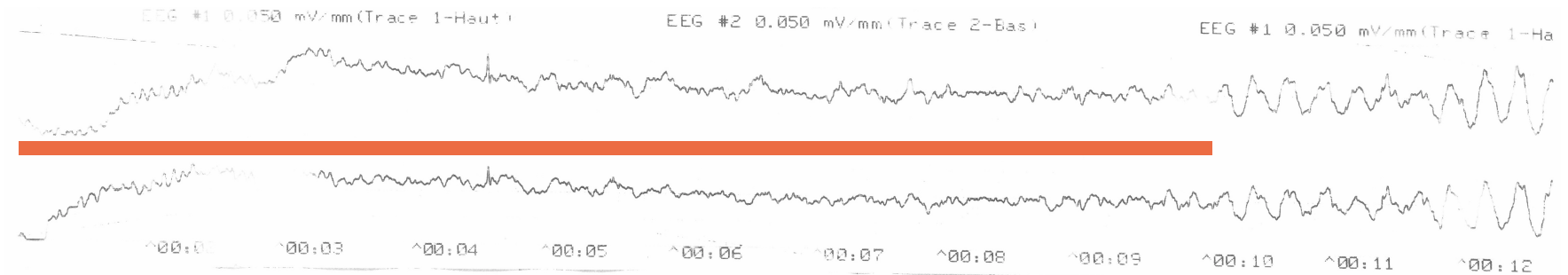
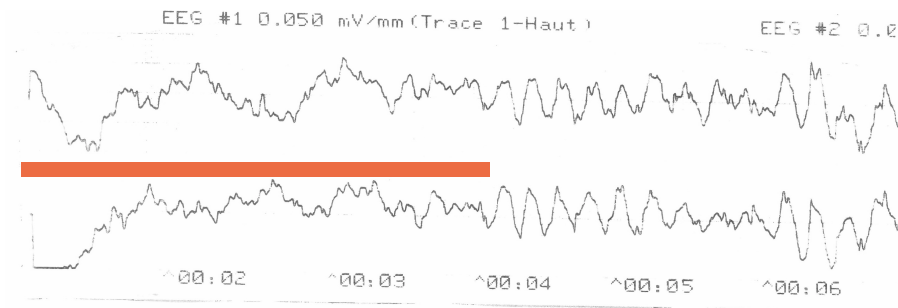
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Peak mid-ictal amplitude Calculate the mean value of 3 consecutive spike and wave complexes which represent the largest of the <i>slow wave/spike-and-wave phase</i> (ignore transition phases).	Wave 1: [mm] Wave 2: [mm] Wave 3: [mm] Mean:
Seizure regularity (0–6) Score the predominant pattern during <i>slow wave/spike-and-wave phase</i> according to the scale (ignore transition phases). See scale by Krystal and Weiner, (1993).	
Seizure stereotypy (0–3; 0.5 pt increments) - Score based on three characteristics. Score 0, 0.5 or 1 for each and sum the total. - Transition from recruitment <i>polyspike</i> phase to the <i>slow wave/spike-and-wave phase</i> (0=no recruitment phase, 1=recruitment phase present and transition to slow wave is clear). Consider transition in terms of both morphology and increase in amplitude. - Quality of the <i>spike-and-wave</i> morphology and reappearance of <i>chaotic polyspikes</i> during the <i>slow wave/spike-and-wave phase</i> (0=no clear spike-and-wave and/or chaotic polyspikes are present during the slow wave/spike-and-wave phase; 1=a stereotypic slow wave/spike-and-wave phase is present, and no reappearance of chaotic polyspikes). - Variability in amplitude of the <i>slow wave/spike-and-wave phase</i> (0=marked variability, the amplitude is inconsistent for >50% of the slow wave phase; 1=the amplitude is consistent for >80% of the slow wave phase) NT: this is excluding transition phases.	
Post-ictal suppression (0–3; 0.5 pt increments) - Score based on three characteristics. Score 0, 0.5 or 1 for each and sum the total. - Seizure end-point (0=cannot determine end-point; 1=clear end-point). - Transition to seizure termination (0=gradual, the transition phase consists of >1/3 of the trace; 1=abrupt). - Flatness of trace after seizure end-point (0=poor; 1=very flat).	

La compréhension de l'échelle

Temps avant l'activité lente (<5Hz)



Time to onset of slowing (TSLOW)

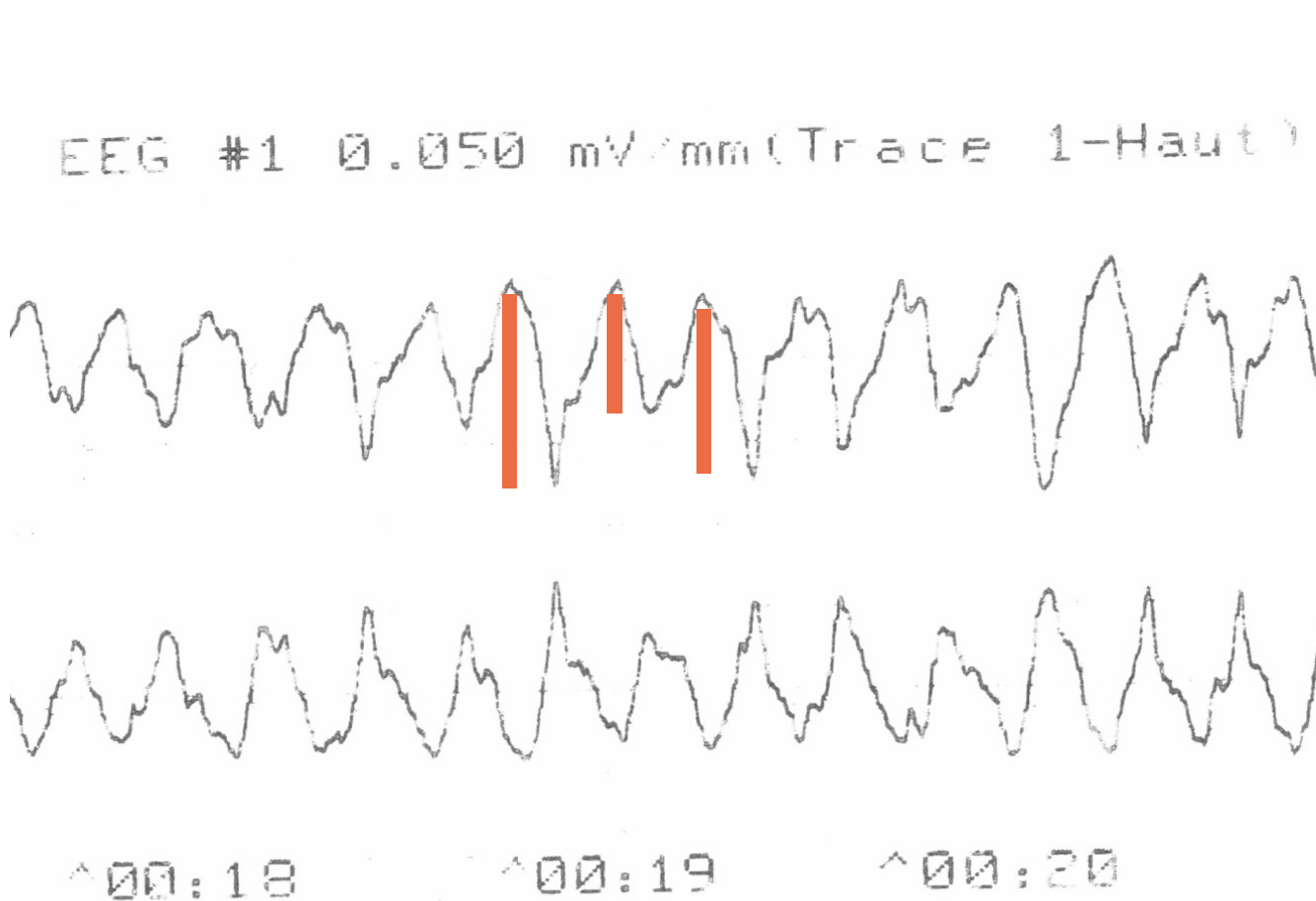


La compréhension de l'échelle

Appendix A. EEG Seizure Quality Rating Sheet

Parameter	Score/value
Time to onset of slowing (TSLOW) ▪ Time from end of ECT stimulus to slowing of frequency to ≤ 5 Hz. ▪ Score as 0 if frequency does not initially exceed 5 Hz.	[secs]
Peak mid-ictal amplitude ▪ Calculate the mean value of 3 consecutive spike and wave complexes which represent the largest of the <i>slow wave/spike-and-wave phase</i> (ignore transition phases).	Wave 1: [mm] Wave 2: [mm] Wave 3: [mm] Mean:
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Post-ictal suppression (0–3; 0.5 pt increments) ▪ Score based on three characteristics. Score 0, 0.5 or 1 for each and sum the total. 1. Seizure end-point (0=cannot determine end-point; 1=clear end-point). 2. Transition to seizure termination (0=gradual, the transition phase consists of >1/3 of the trace; 1=abrupt). 3. Flatness of trace after seizure end-point (0=poor; 1=very flat).	

Peak mid-ictal amplitude



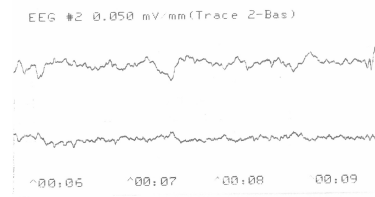
La compréhension de l'échelle

Appendix A. EEG Seizure Quality Rating Sheet

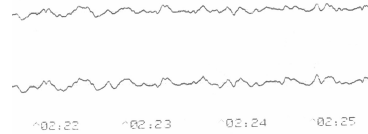
Parameter	Score/value
<i>Time to onset of slowing (TSLOW)</i> - Time from end of ECT stimulus to slowing of frequency to ≤ 5 Hz. - Score as 0 if frequency does not initially exceed 5 Hz.	[secs]
<i>Peak mid-ictal amplitude</i> Calculate the mean value of 3 consecutive spike and wave complexes which represent the largest of the <i>slow wave/spike-and-wave phase</i> (ignore transition phases).	Wave 1: [mm] Wave 2: [mm] Wave 3: [mm] Mean:
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Seizure regularity

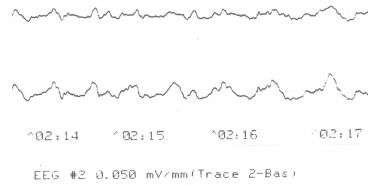
0



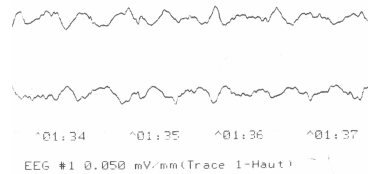
1



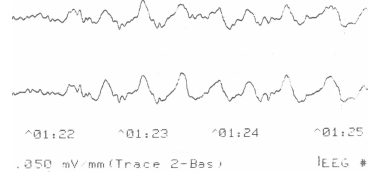
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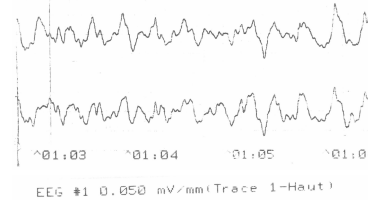
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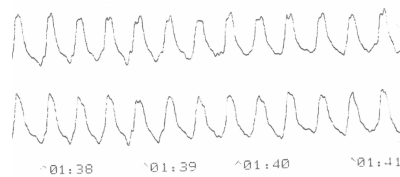
4



5



6



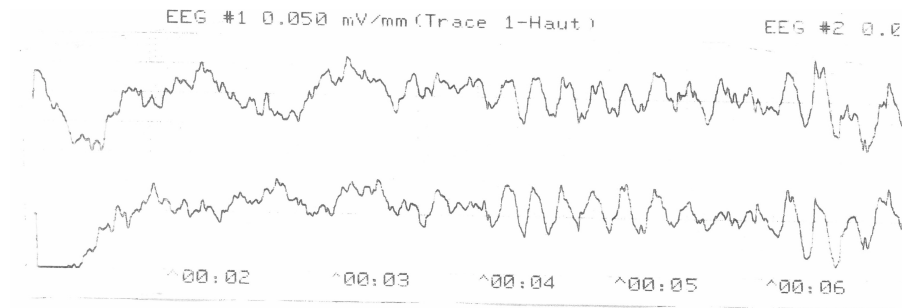
La compréhension de l'échelle

Appendix A. EEG Seizure Quality Rating Sheet

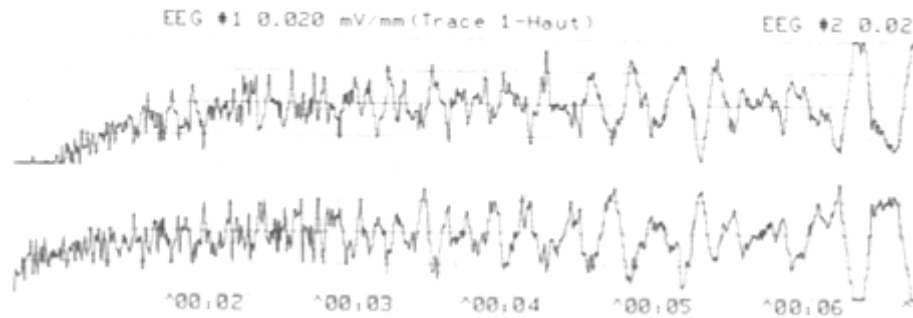
Parameter	Score/value
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<i>Post-ictal suppression (0–3; 0.5 pt increments)</i> - Score based on three characteristics. Score 0, 0.5 or 1 for each and sum the total. - Seizure end-point (0=cannot determine end-point; 1=clear end-point). - Transition to seizure termination (0=gradual, the transition phase consists of >1/3 of the trace; 1=abrupt). - Flatness of trace after seizure end-point (0=poor; 1=very flat).	

Seizure stereotypy : transition recrutement

0

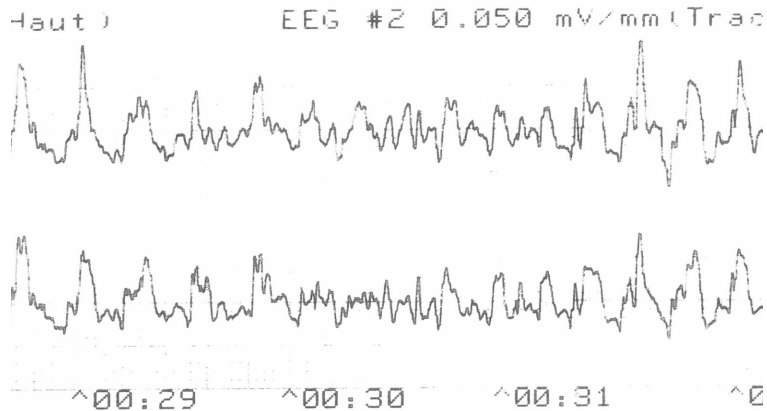


1

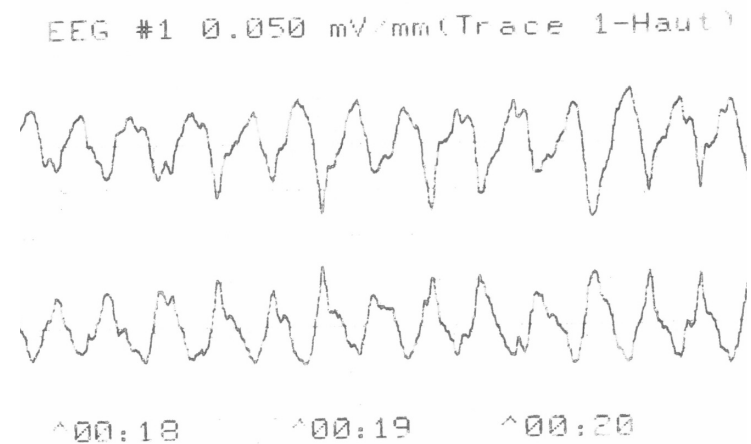


Seizure stereotypy : Chaotic polyspikes

0



1

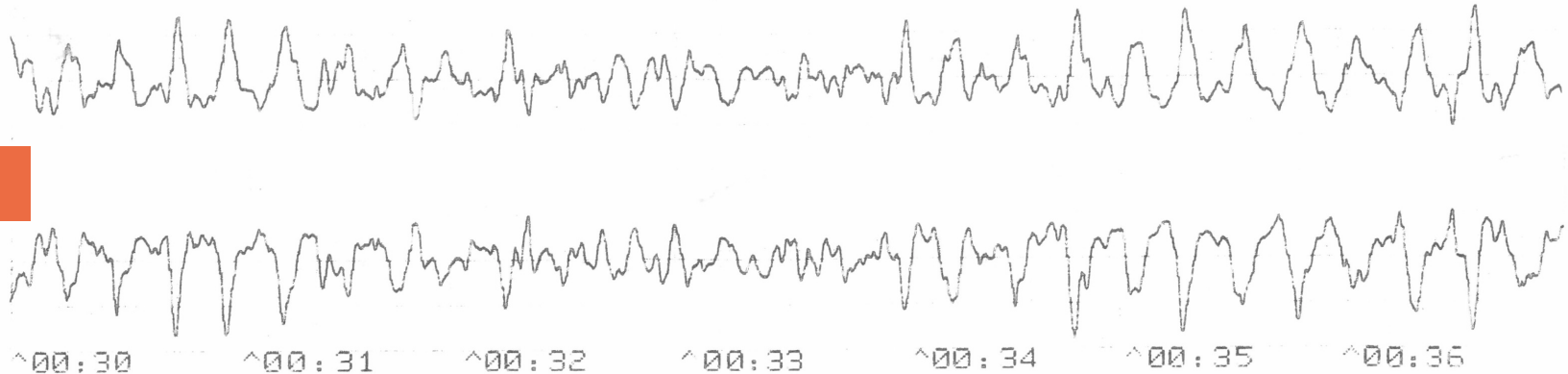


Seizure stereotypy : Variability amplitude

EEG #2 0.050 mV/mm (Trace 2-Bas)

EEG #1 0.050 mV/mm (Trace 1-Hau)

0



La compréhension de l'échelle

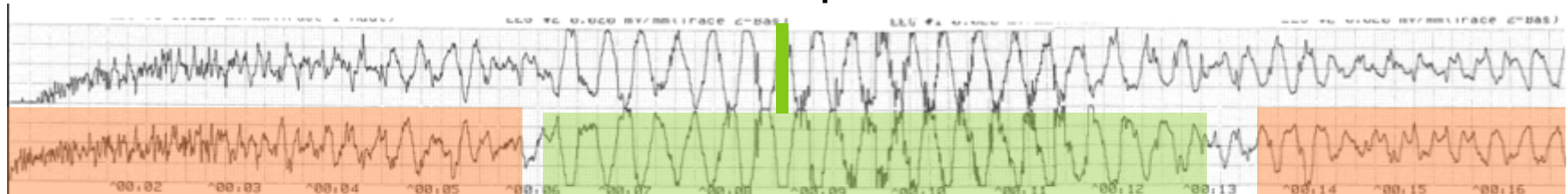
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La compréhension de l'échelle

Temps avant l'activité lente (<5Hz)

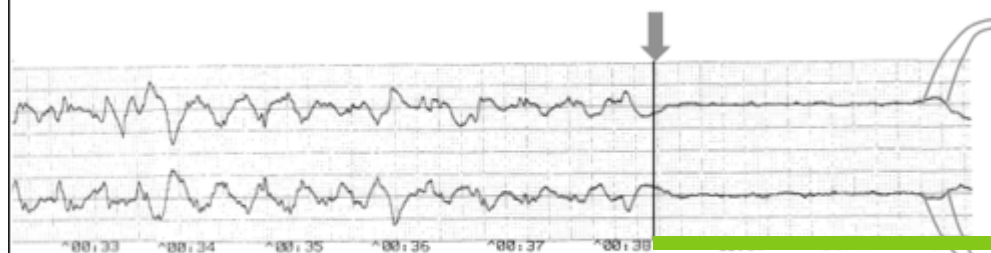
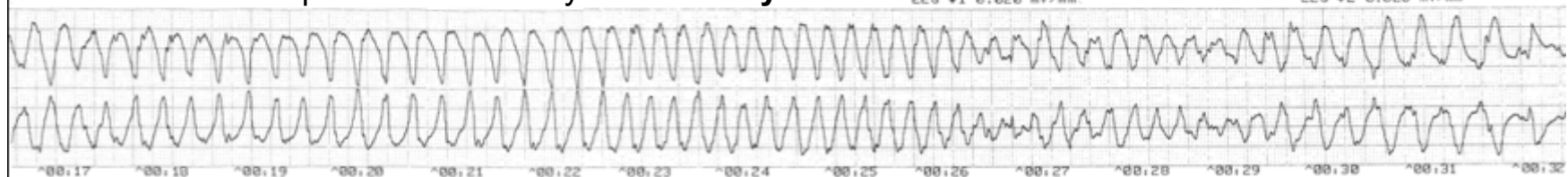
Amplitude des ondes lentes



Recrutement/tonique/transition

Rythmicité et **synchronie**

Variabilité



Suppression post ictale

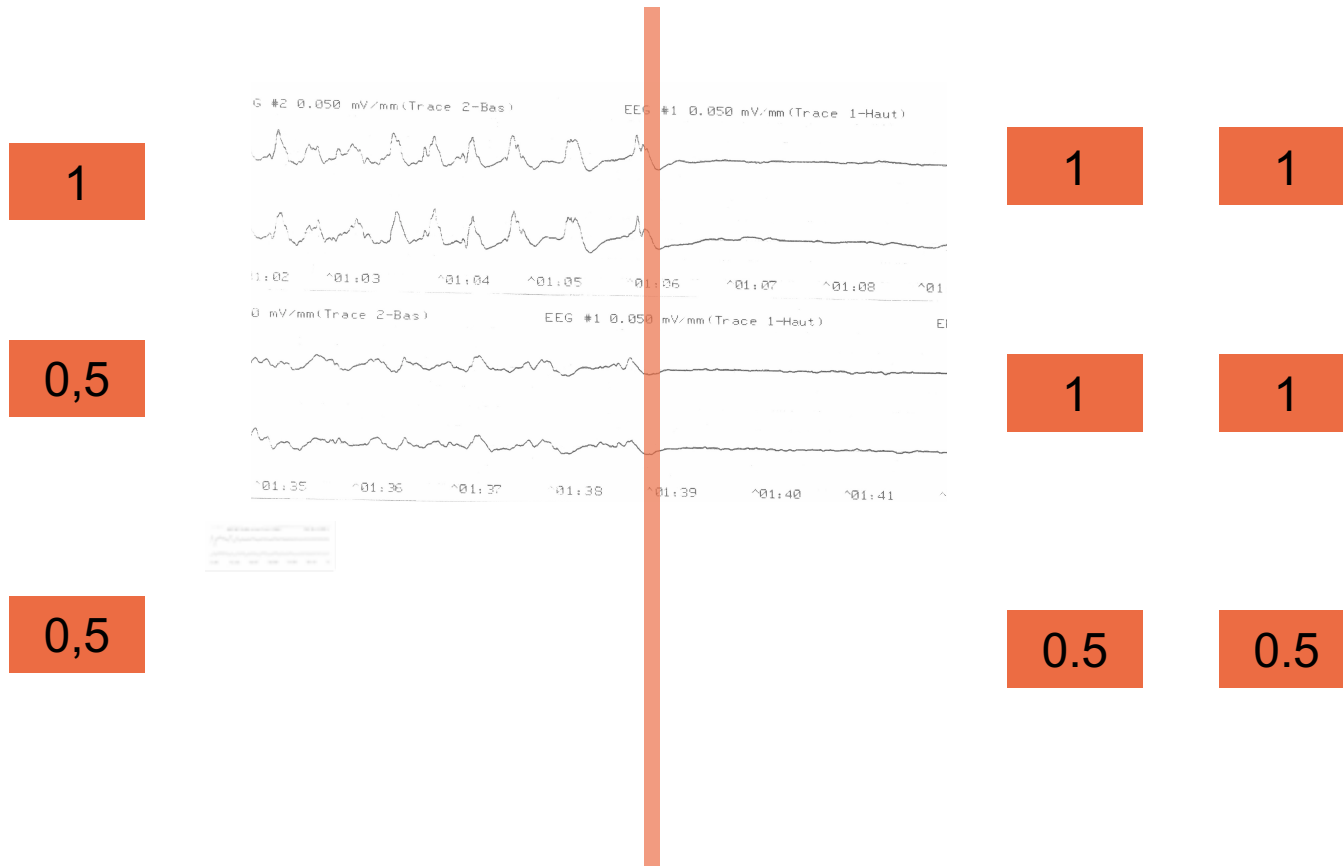
Stéréotypie

CHARGE 1152 mC
ENERGIE 182.7 J
STAT. IMPED. 628 ohms
DYN. IMPED. 187 ohms
PULSE 1.50 msec
FREQUENCE 120 Hz
DUREE 4.000 sec
COURANT 000 mA

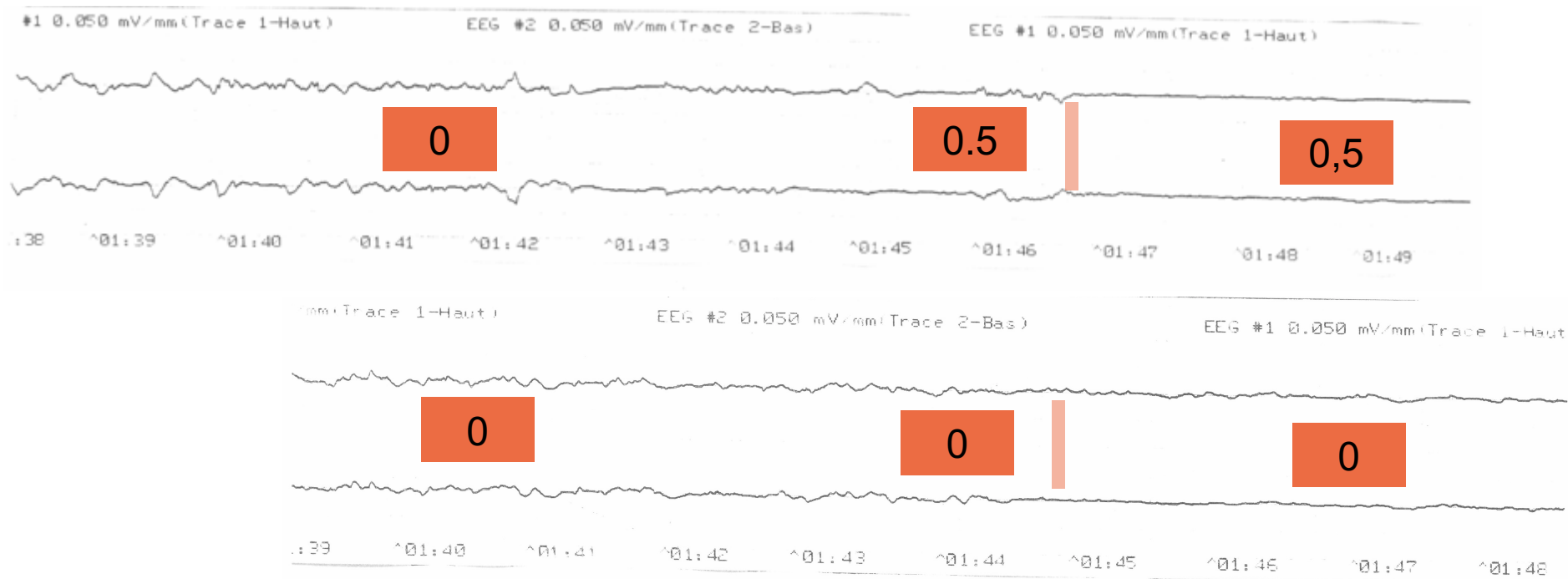
18/09/2012
09:30:15
ID PATIENT ?
AGE ? NUTRIT ?

NON PATIENT

Post ictal suppression

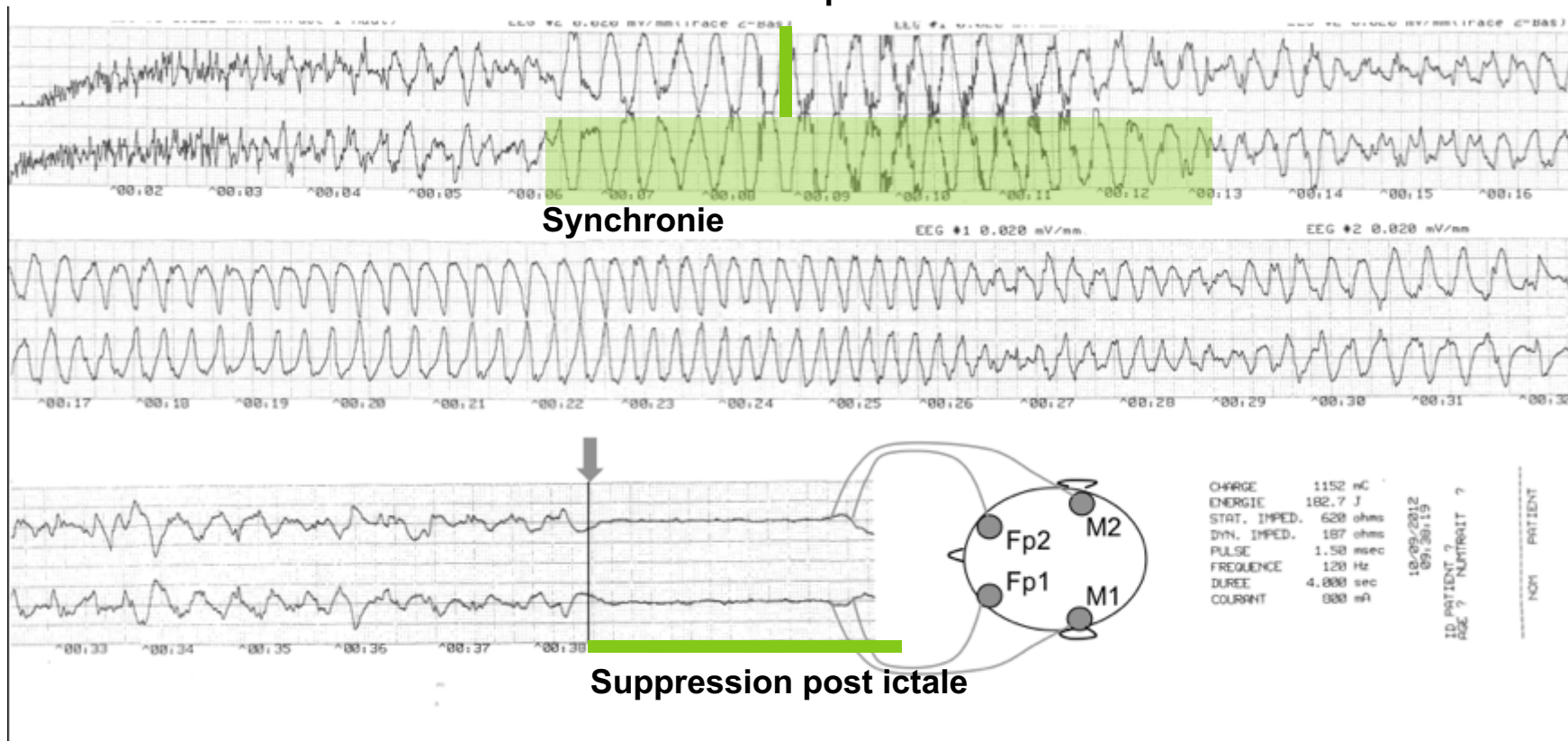


Post ictal supression

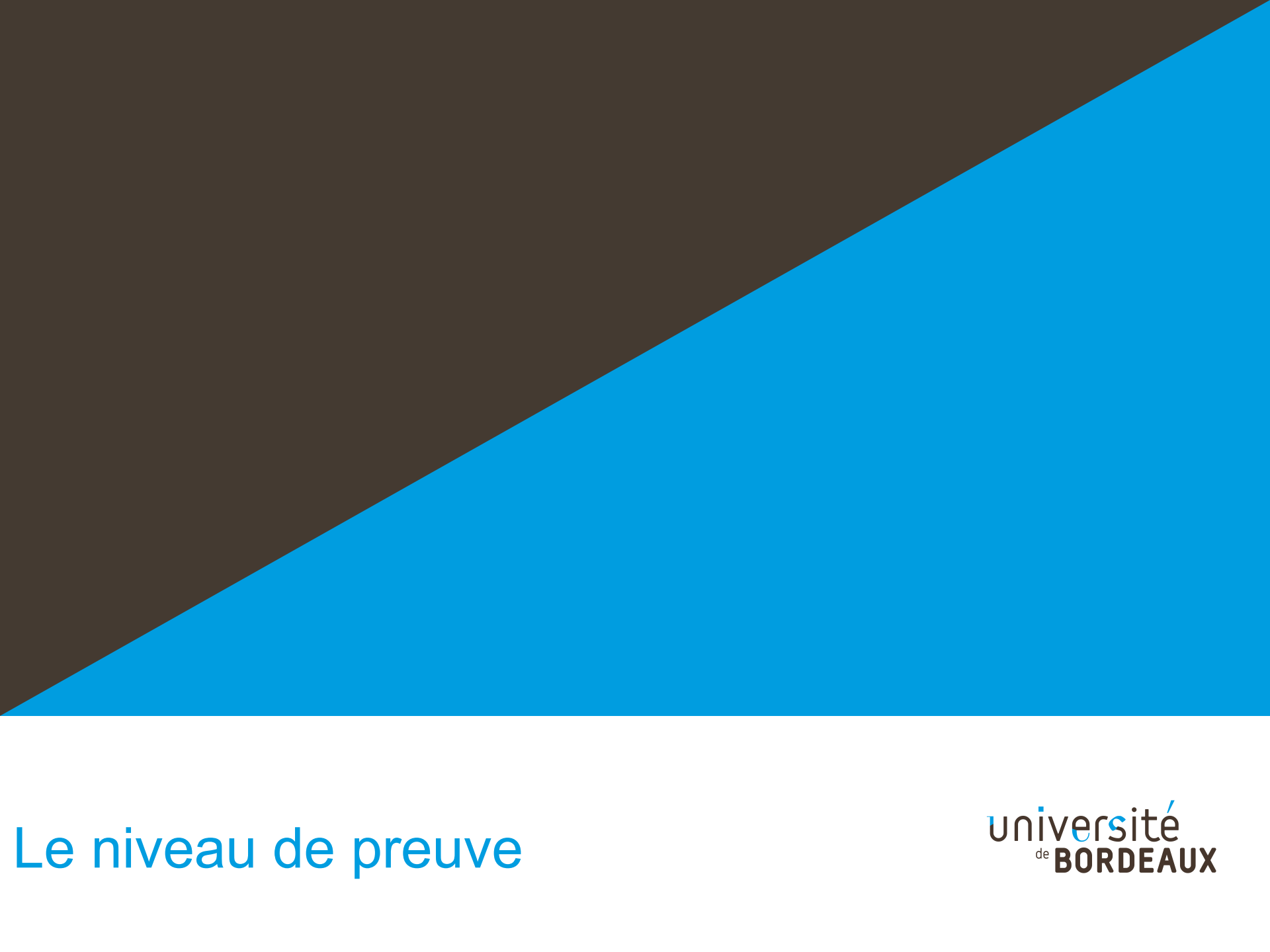


Critères quantifiés

Amplitude des ondes lentes



- 1- Indice de suppression post-critique (ou de l'activité biocorticale, SABC)
- 2- Amplitude maximale de l'EEG durant la crise
- 3- Indice de cohérence inter hémisphérique



Le niveau de preuve

Quel niveau de preuve pronostique pour la cure ECT ?

UNE LITTERATURE PLUTOT ANCIENNE ?

Les prémisses

Brit. J. Psychiat. (1982), **141**, 357-366

A Double-Blind Controlled Comparison of the Therapeutic Effects of Low and High Energy Electroconvulsive Therapies

ASHLEY ROBIN and SANATH DE TISSERA

Amplitudes de crise plus amples et suppression post-ictale reliée à l'efficacité des crises

MAIS

En courant sinusoïdal

EEG Manifestations during ECT: Effects of Electrode Placement and Stimulus Intensity

Mitchell S. Nobler, Harold A. Sackeim, Maria Solomou, Bruce Lubner, D.P. Devanand, and Joan Prudic

La durée de la crise n'influe pas

BIOL. PSYCHIATRY
1993;34:321-330

321

EEG Manifestations during ECT: Effects of Electrode Placement and Stimulus Intensity

Mitchell S. Nobler, Harold A. Sackeim, Maria Solomou, Bruce Luber, D.P. Devanand, and Joan Prudic

A variety of qualitative ratings were scored on 7-point Likert-like scales ranging from 0 to 3, in increments of 0.5. These included global seizure strength, global seizure patterning (stereotypy), two indices of the termination phase (extent of amplitude and frequency reduction), and degree of immediate postictal suppression (suppression of bioelectric activity). Seizures were rated as having greater strength if slow-wave activity of high amplitude predominated during the slow-wave phase. Seizures were rated as more stereotypic if there was a clear progression from low amplitude chaotic polyspike activity to high amplitude slow-wave activity without the reappearance of chaotic polyspike activity or marked variability in amplitude during phases. Ratings for some of these features for prototypic examples of ictal activity are provided in Figure 1. In addition, ratings of total seizure duration and measurement of peak slow-wave amplitude were also made for the left hemisphere channel. Symmetry between left and right channels was rated on a scale from -3.0 to +3.0, again with 0.5 increments, with -3.0 representing markedly greater seizure strength in the left hemisphere lead, and +3.0 indicating an opposite pattern of asymmetry (see Figure 1).



La durée de la crise n'influe pas

BIOL PSYCHIATRY 321
1993;34:321-330

EEG Manifestations during ECT: Effects of Electrode Placement and Stimulus Intensity

Mitchell S. Nobler, Harold A. Sackeim, Maria Solomou, Bruce Lubner, D.P. Devanand, and Joan Prudic

UL (29)		BL (31)	
Faible dose	Forte dose	Faible dose	Forte dose

La durée des crises ne diffèrent pas

MAIS : les critères qualitatifs diffèrent

En particulier la suppression post ictale qui prédit les répondeurs

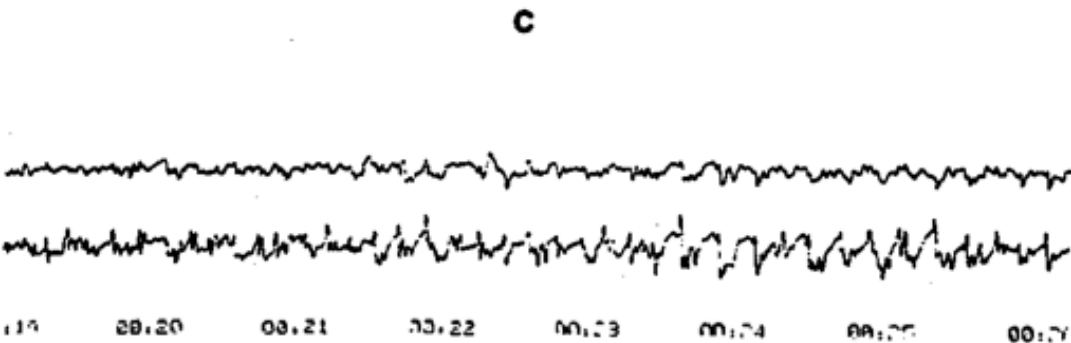
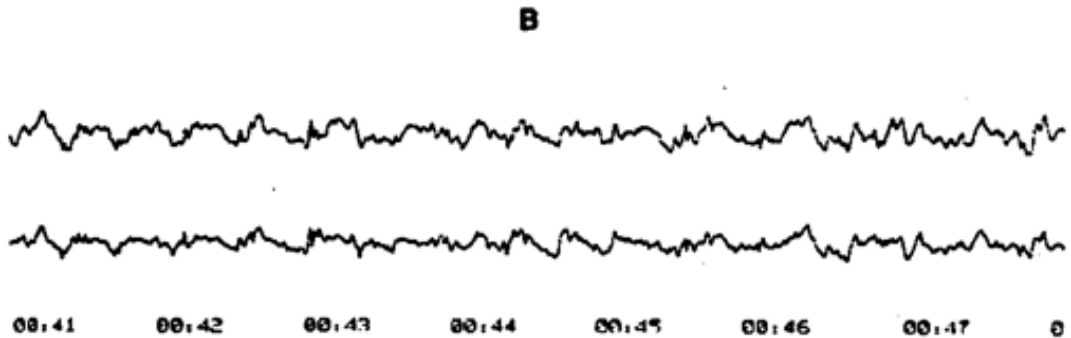
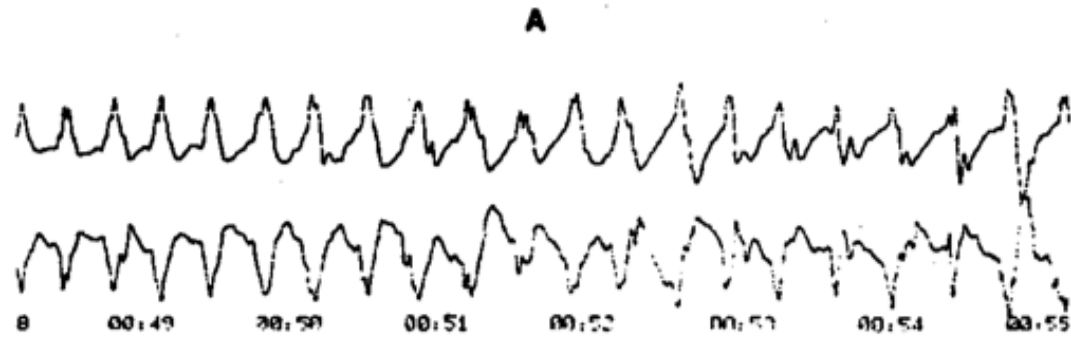
La durée de la crise n'influe pas

Table 3. Mean Values for Ictal EEG Ratings*

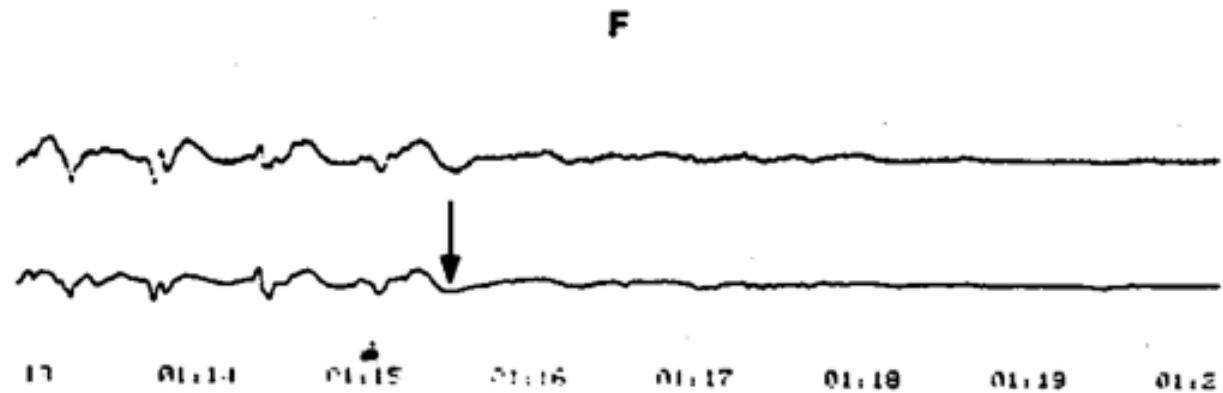
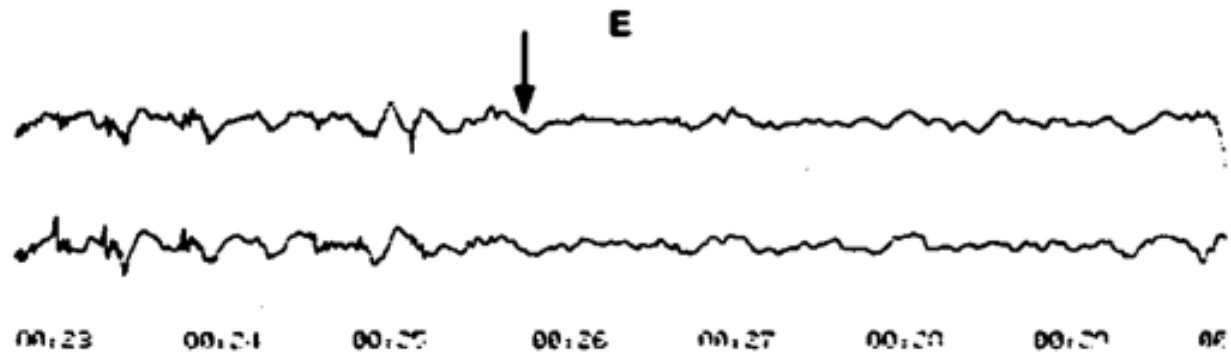
	Low dose unilateral (n = 15)		High dose unilateral (n = 14)		Low dose bilateral (n = 15)		High dose bilateral (n = 19)	
	Mean	SD	Mean	SD	Mean	SD	Mean	SD
<i>Right Hemisphere</i>								
Duration of polyspike phase (sec)	10.29 ^a	6.17	5.60 ^b	2.01	6.78 ^b	2.42	5.58 ^b	3.54
Duration of slow-wave phase (sec)	40.60 ^a	13.75	52.23 ^b	19.76	44.44 ^b	10.38	45.37 ^b	12.65
Total duration (sec)	50.89	13.89	57.83	19.06	51.22	11.48	50.95	12.32
Peak slow-wave amplitude (μv)	550.83 ^a	166.24	526.07 ^a	158.02	466.77 ^a	127.84	633.73 ^b	171.88
Overall strength	1.28 ^a	0.58	1.50 ^{a,b}	0.69	1.23 ^a	0.41	1.63 ^b	0.60
Overall pattern	1.38 ^a	0.67	1.78 ^b	0.62	1.51 ^{a,b}	0.49	1.78 ^b	0.50
Postictal suppression	1.09 ^a	0.67	1.64 ^b	0.93	1.95 ^b	0.43	1.78 ^b	0.77
<i>Left Hemisphere</i>								
Total duration (sec)	53.67	12.70	58.37	19.27	52.88	13.02	51.09	12.51
Peak slow-wave amplitude (μv)	515.33 ^a	209.92	548.91 ^a	168.96	457.34 ^a	157.11	598.95 ^b	201.99
Overall symmetry	+0.19	0.89	+0.29	0.90	+0.38	0.74	+0.70	1.17

*Groups with differing superscripts (e.g., *a* versus *b*) differed in pair-wise *t*-tests on least-squares adjusted means at trend or significant levels (all *p*'s < 0.1, two-tailed).

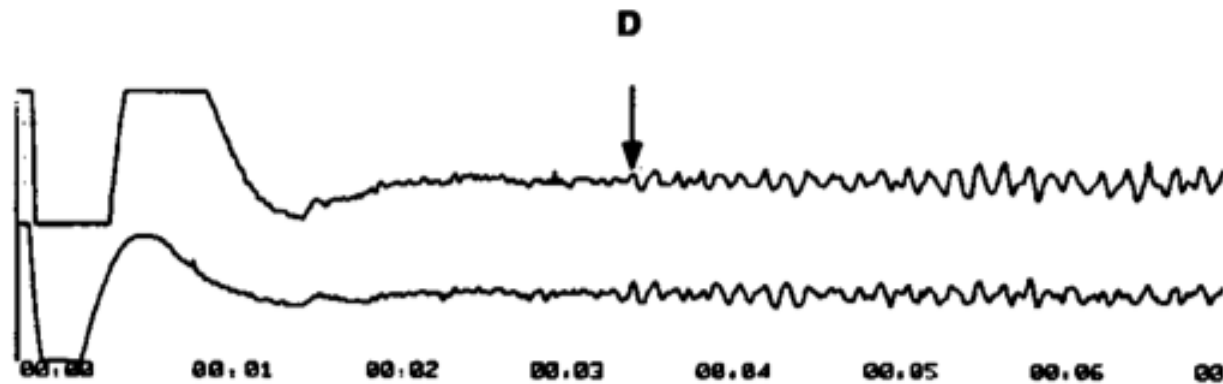
Différente amplitude



Différente terminaison



Apparition de la phase clonique



The Journal of ECT
16(3):211-228 © 2000 Lippincott Williams & Wilkins, Inc., Philadelphia

Quantitative EEG During Seizures Induced by Electroconvulsive Therapy: Relations to Treatment Modality and Clinical Features. I. Global Analyses

Mitchell S. Nobler, M.D., Bruce Luber, Ph.D., James R. Moeller, Ph.D.,
Gary P. Katzman, B.A., Joan Prudic, M.D., *D. P. Devanand, M.D.,
Gabriel S. Dichter, B.A., and †Harold A. Sackeim, Ph.D.

62 patients, UL & BL

Régularité et stérotypie : faible predicteur
Suppression post ictale: plus robuste

Les études de Krystal et al. Duke University

BIOL PSYCHIATRY
1993;34:759-767

759

The Effects of ECT Stimulus Dose and Electrode Placement on the Ictal Electroencephalogram: An Intraindividual Crossover Study

Andrew D. Krystal, Richard D. Weiner, W. Vaughn McCall, Frank E. Shelp, Rebecca Arias, and Pamela Smith

BIOL PSYCHIATRY
1992;31:617-621

617

EEG Evidence of More “Intense” Seizure Activity with Bilateral ECT

Andrew D. Krystal, Richard D. Weiner, C. Edward Coffey, Pamela Smith, Rebekka Arias, and Eric Moffett

Changes in Seizure Threshold Over the Course of Electroconvulsive Therapy Affect Therapeutic Response and Are Detected by Ictal EEG Ratings

Andrew D. Krystal, M.D., M.S.
C. Edward Coffey, M.D.
Richard D. Weiner, M.D., Ph.D.
Tracey Holsinger, M.D.

J Neuropsychiatry Clin Neurosci. 1995 Summer;7(3):295-303.

The ictal EEG as a marker of adequate stimulus intensity with unilateral ECT.

Krystal AD¹, Weiner RD, Coffey CE.

Les études de Krystal et al. Duke University

BIOL PSYCHIATRY 617
1992;31:617-621

EEG Evidence of More “Intense” Seizure Activity with Bilateral ECT

Andrew D. Krystal, Richard D. Weiner, C. Edward Coffey,
Pamela Smith, Rebekka Arias, and Eric Moffett

Manual Analysis. Manual ratings of EEGs recorded on paper during the ECT seizures were made by EM and included (1) *maximum peak-to-peak amplitude* of the seizure (mean values over 3 contiguous spike-and-wave complexes), (2) *immediate postictal amplitude* (mean value over first 5 sec following the end of the seizure), and qualitative estimates of both (3) *seizure symmetry* (7-point ordinal scale, -3 to +3, and 0 = symmetrical), and (4) *seizure regularity* (7-point ordinal scale, 0 to 7).

+ QUANTITATIVE ANALYSIS

Analyse spectrale : amplitude ondes lentes et suppression post ictale

Analyse de cohérence

La durée de la crise n'influe pas

BIOL PSYCHIATRY 617
1992;31:617-621

EEG Evidence of More “Intense” Seizure Activity with Bilateral ECT

Andrew D. Krystal, Richard D. Weiner, C. Edward Coffey,
Pamela Smith, Rebekka Arias, and Eric Moffett

UL (4)

BL (5)

La durée des crises ne diffèrent pas

MAIS : les critères qualitatifs et quantitatifs diffèrent

Les études de Krystal et al. Duke University

1993, November
1993, Vol. 19, No. 10

The Effects of ECT Stimulus Dose and Electrode Placement on the Ictal Electroencephalogram: An Intraindividual Crossover Study

Andrew D. Krystal, Richard D. Weiner, W. Vaughn McCall, Frank E. Shelp, Rebecca Arias, and Pamela Smith

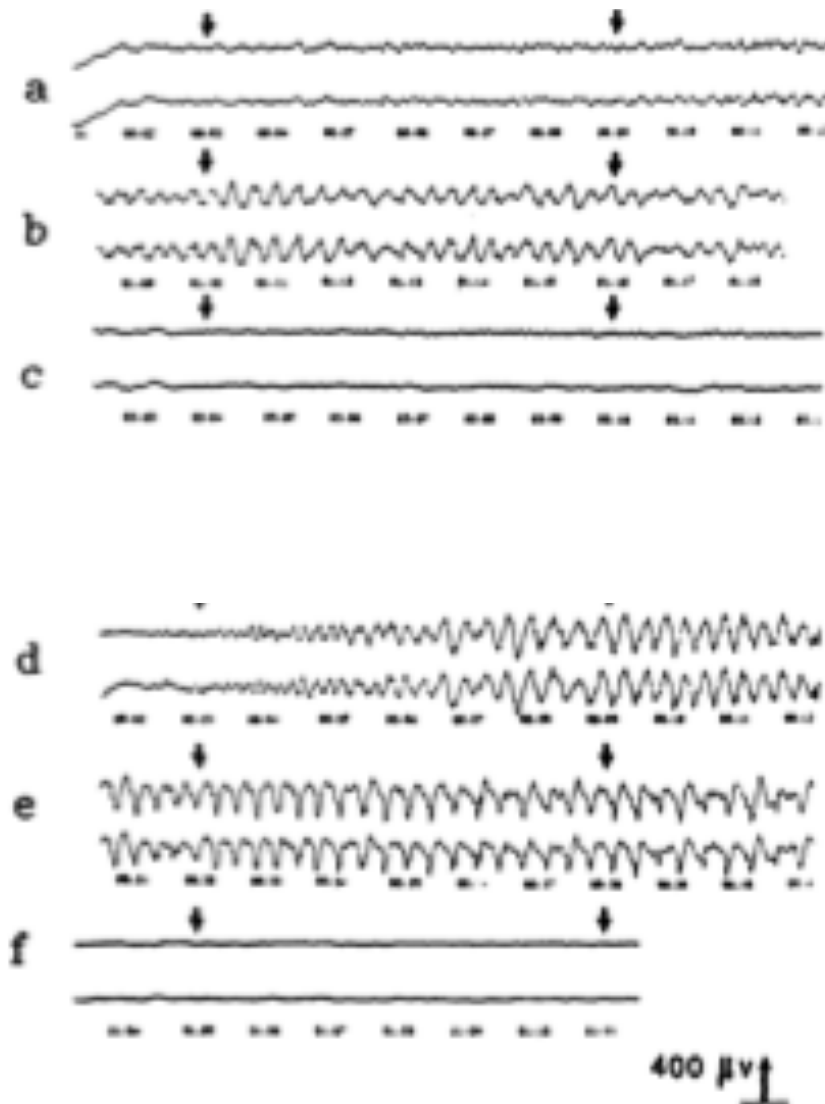
UL (9)		BL (10)	
Faible dose	Forte dose	Faible dose	Forte dose

La durée des crises ne diffèrent pas

MAIS : les critères qualitatifs et quantitatifs diffèrent

Attention : la durée de crise diminue en augmentant la dose en BL

Les études de Krystal et al. Duke University



Les études de Krystal et al. Duke University

Changes in Seizure Threshold
Over the Course of
Electroconvulsive Therapy
Affect Therapeutic Response
and Are Detected by Ictal EEG
Ratings

Andrew D. Krystal, M.D., M.S.
C. Edward Coffey, M.D.
Richard D. Weiner, M.D., Ph.D.
Tracey Holsinger, M.D.

UL (37)

BL (10)



Le seuil épileptogène augmente

En UL : les critères qualitatifs prédisent cette augmentation

Et prédisent la réponse à 6 séances

Les autres études



Journal of Affective Disorders 41 (1996) 55–58



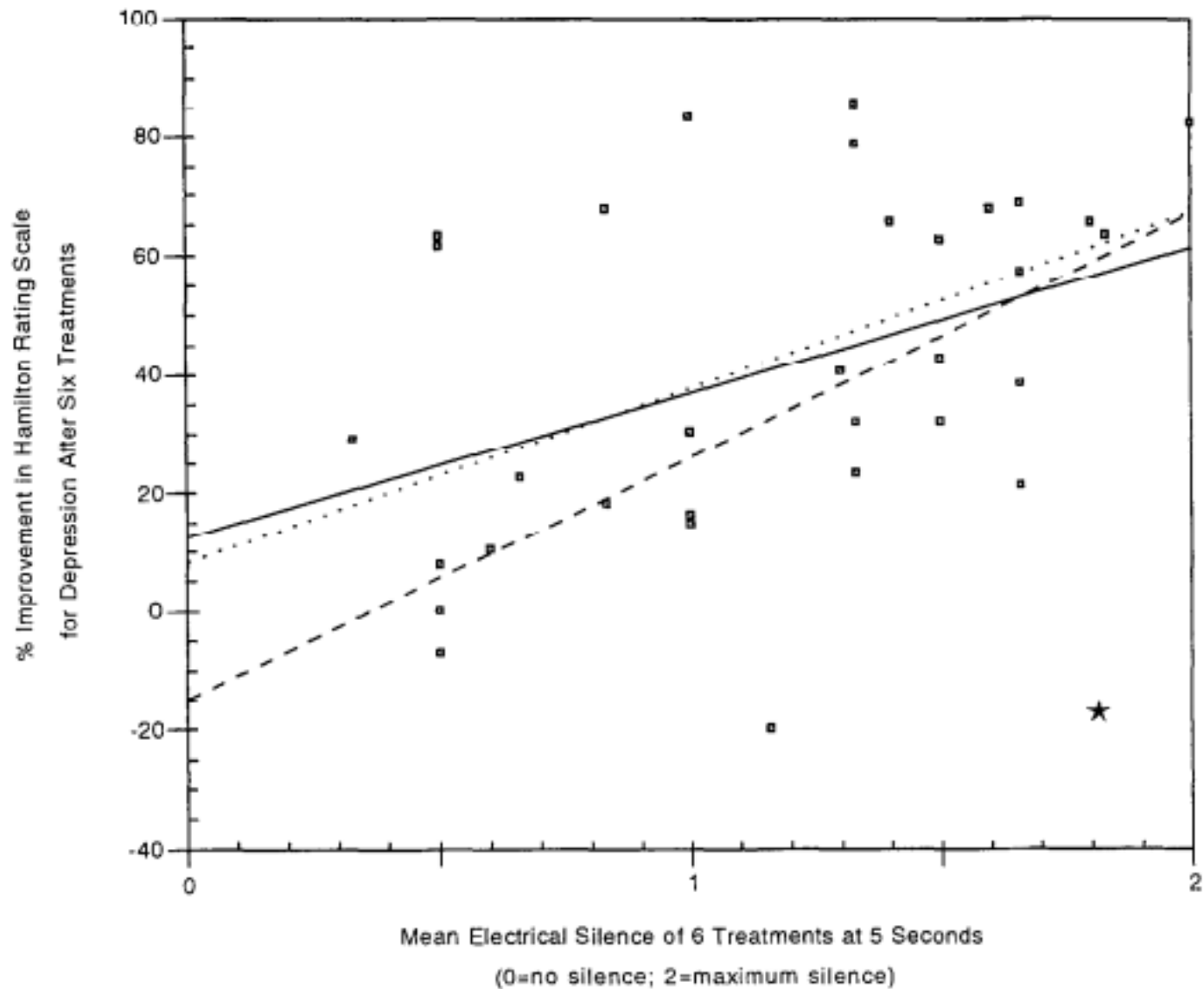
Research report

Is postictal electrical silence a predictor of response to electroconvulsive therapy?

Trisha Suppes ^{a,*}, Andrew Webb ^a, Thomas Carmody ^a, Eunice Gordon ^b,
Rolando Gutierrez-Esteinou ^c, James I. Hudson ^d, Harrison G. Pope Jr. ^d

33 patients, UL ou BL

Les autres études



ORIGINAL PAPER

Here Folkerts

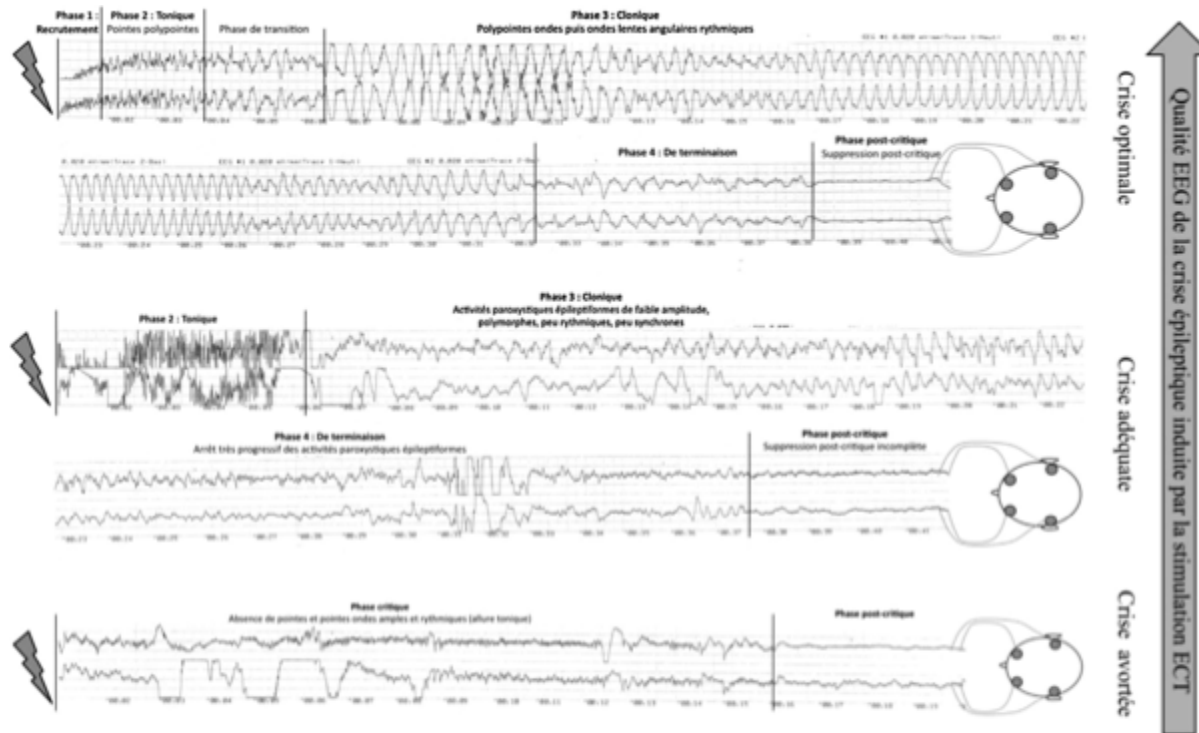
The ictal electroencephalogram as a marker for the efficacy of electroconvulsive therapy

40 patients, UL

Régularité et stérotypie

En résumé

Aide mémoire

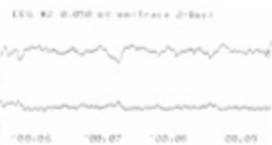




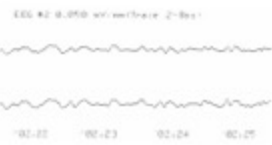
Critère 3 : Régularité de la crise
Seizure regularity
 (0 – 6)

Critère adapté de Weiner & Krystal, 1993

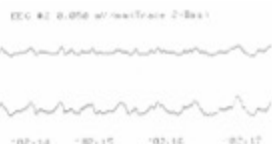
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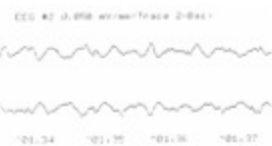
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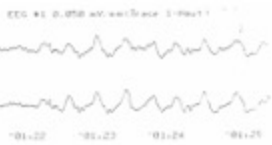
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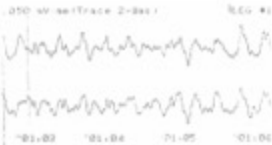
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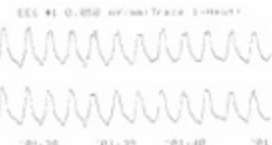
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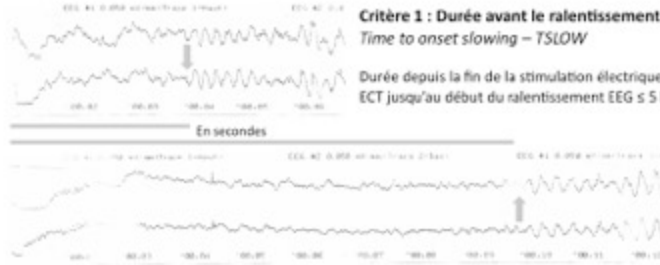


6



Critère 1 : Durée avant le ralentissement
Time to onset slowing – TSLOW

Durée depuis la fin de la stimulation électrique
 ECT jusqu'au début du ralentissement EEG ≤ 5 Hz



Critère 2 : Amplitude pic à pic en milieu de la phase clonique
Peak mid-ictal amplitude

Amplitudes de 3 ondes consécutives représentatives
 de la phase clonique la plus ample

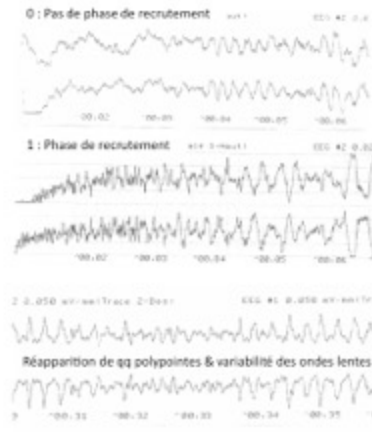


Critère 4 : Stéréotypie de la crise
Seizure stereotypy
 (0 – 3)

1 – Coder (0, 0,5, 1) la présence d'une
 phase de recrutement vers la phase
 clonique

2 – Coder (0, 0,5, 1) la qualité de la
 morphologie des ondes lentes /
 pointes-ondes et la réapparition de
 polypointes chaotiques

3 – Coder (0, 0,5, 1) la variabilité en
 amplitude des ondes lentes /
 pointes-ondes de la phase clonique

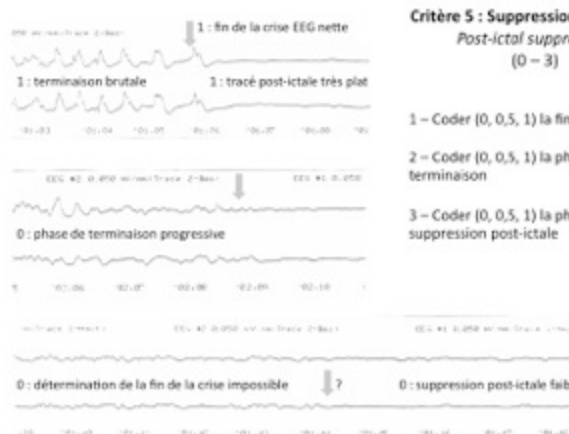


Critère 5 : Suppression post-ictale
Post-ictal suppression
 (0 – 3)

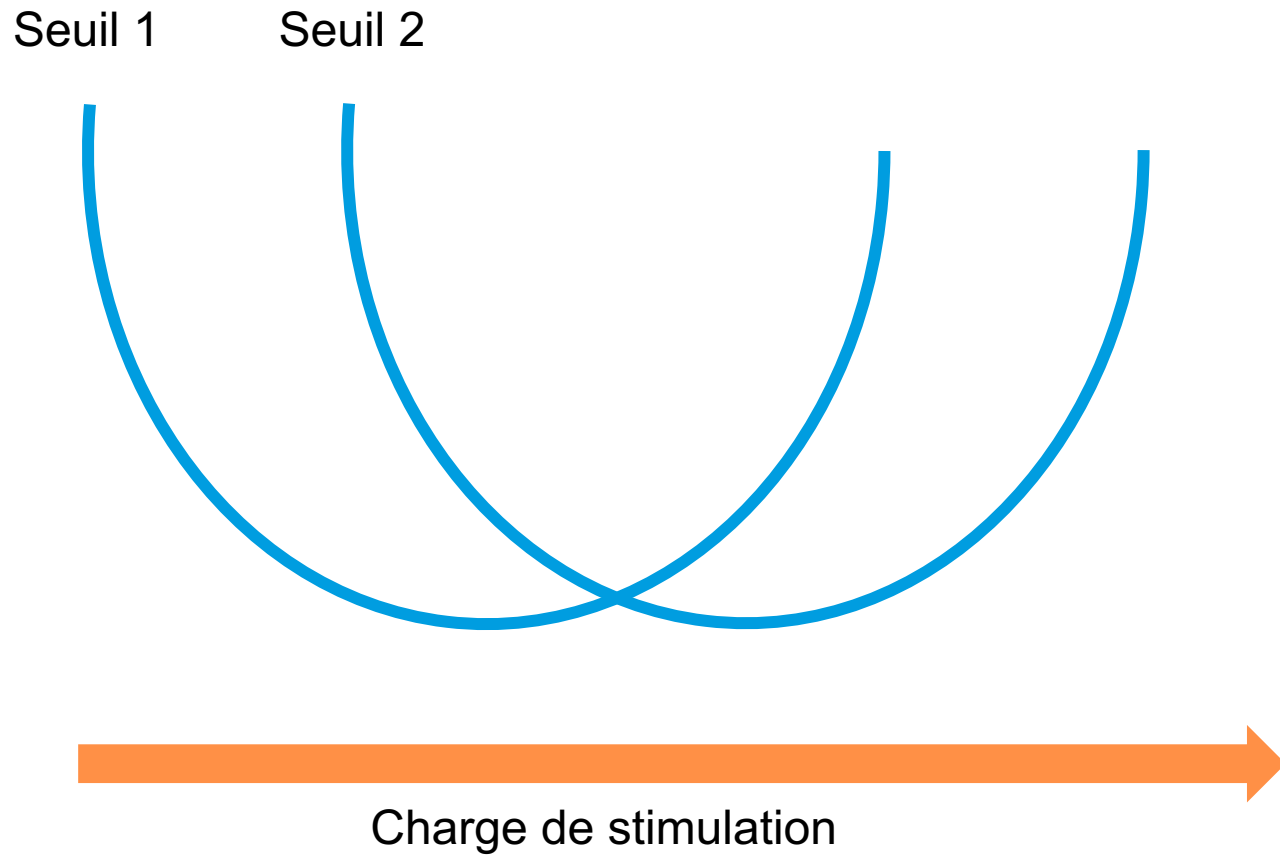
1 – Coder (0, 0,5, 1) la fin de la crise

2 – Coder (0, 0,5, 1) la phase de
 terminaison

3 – Coder (0, 0,5, 1) la phase de
 suppression post-ictale



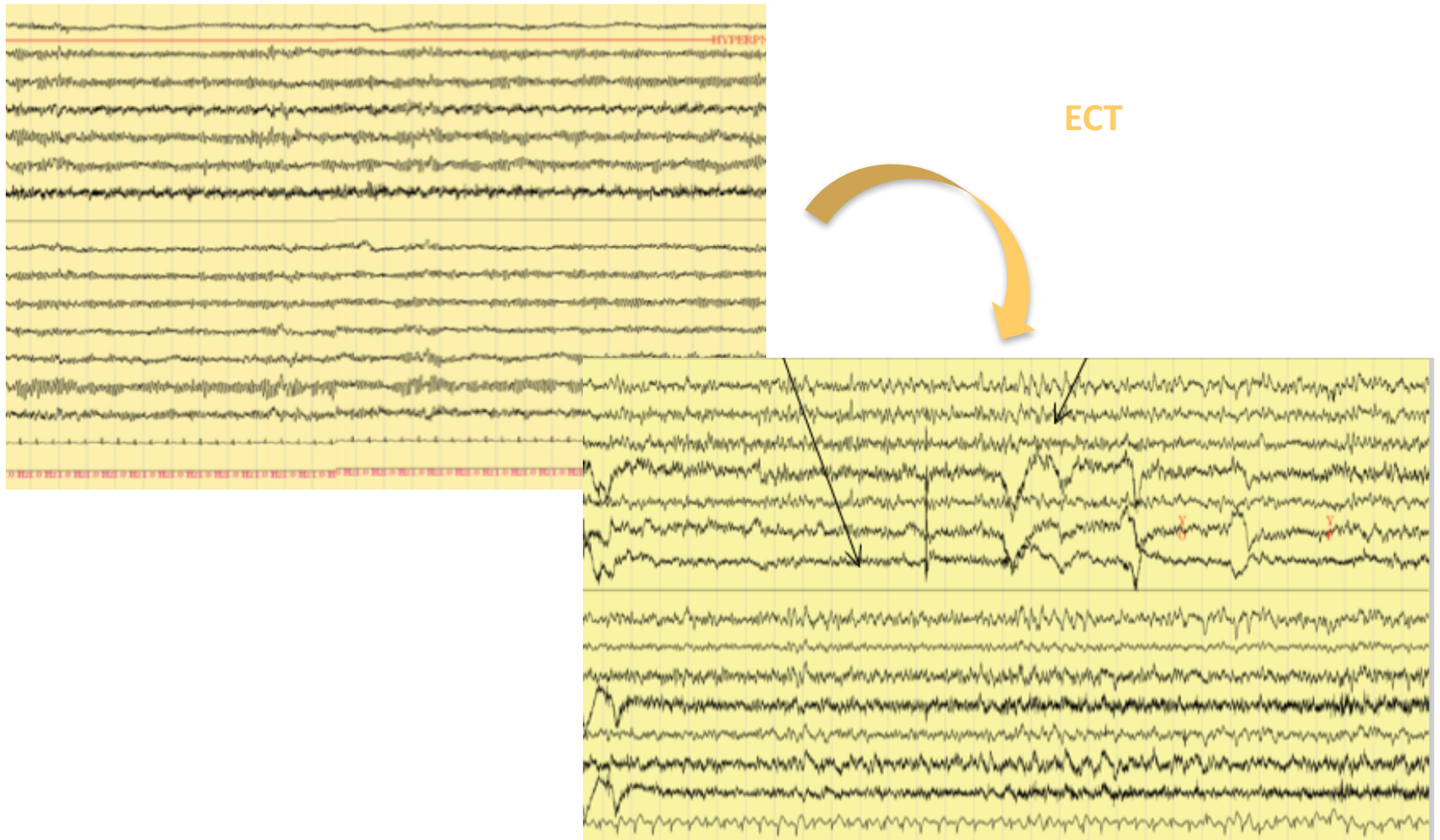
Take home message



En inter critique

EEG conventionnel et ECT

Modification EEG post ECT



Crise épileptique indésirable

Pas de surveillance EEG systématique

Les modifications EEG sont reliées à une meilleure réponse thérapeutique aux ECT

Crise épileptique indésirable / Etat de mal épileptique indésirable

Pas de rôle prédictif de l'EEG de base

Facteurs favorisant:

- ATCD de crises prolongées
- Première séance ECT
- ATCD de pathologies neurologiques (crise épileptique, maladie de parkinson)
- TRT abaissant seuil épileptogène, arrêt TRT AE

Risque de récurrence (discuter TRT AE)

EEG devant toute confusion post ECT pour éliminer état de mal épileptique indésirable

The logo of the University of Bordeaux is displayed against a background with a blue diagonal stripe in the top left and a dark grey diagonal stripe in the bottom right. The text 'université' is in a dark brown, lowercase, sans-serif font, with a blue accent on the 'u' and 's'. Below it, 'de' is in a smaller, dark brown, lowercase, sans-serif font, and 'BORDEAUX' is in a larger, dark brown, uppercase, sans-serif font.

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