



Minerva Access is the Institutional Repository of The University of Melbourne

Author/s:

Fisher, RS;Acevedo, C;Arzimanoglou, A;Bogacz, A;Cross, JH;Elger, CE;Engel, J;Forsgren, L;French, JA;Hesdorffer, DC;Lee, BI;Mathern, GW;Moshe, SL;Perucca, E;Scheffer, IE;Tomson, T;Watanabe, M;Wiebe, S

Title:

How long for epilepsy remission in the ILAE definition?

Date:

2017-08-01

Citation:

Fisher, R. S., Acevedo, C., Arzimanoglou, A., Bogacz, A., Cross, J. H., Elger, C. E., Engel, J., Forsgren, L., French, J. A., Hesdorffer, D. C., Lee, B. I., Mathern, G. W., Moshe, S. L., Perucca, E., Scheffer, I. E., Tomson, T., Watanabe, M. & Wiebe, S. (2017). How long for epilepsy remission in the ILAE definition?. *Epilepsia*, 58 (8), pp.1486-1487. <https://doi.org/10.1111/epi.13821>.

Persistent Link:

<https://hdl.handle.net/11343/293263>

Article type : Letter

How long for epilepsy remission in the ILAE definition?

by

*Robert S. Fisher, †Carlos Acevedo, ‡Alexis Arzimanoglou, §Alicia Bogacz, ¶J. Helen Cross, #Christian E. Elger, **Jerome Engel Jr, ††Lars Forsgren, ‡‡Jacqueline A. French, ¶¶Dale C. Hesdorffer, ## Byung-In Lee, ***Gary W. Mathern, †††Solomon L. Moshe, ‡‡‡Emilio Perucca, §§§Ingrid E. Scheffer, ¶¶¶Torbjørn Tomson, ###Masako Watanabe, and ****Samuel Wiebe

* (Fisher) Department of Neurology & Neurological Sciences, Stanford University School of Medicine, Stanford, California, U.S.A.;

† (Acevedo) Past President Chilean League Against Epilepsy, Santiago, Chile;

‡ (Arzimanoglou) University Hospitals of Lyon (HCL), Department of Clinical Epileptology, Sleep Disorders and Functional Neurology in Children, Lyon, France;

§ (Bogacz) Neurological Institute of Clinical Hospital, Universidad Mayor de la Republica. Montevideo, Uruguay;

¶ (Cross) UCL-Institute of Child Health, Great Ormond Street Hospital for Children, London & Young Epilepsy, Lingfield, United Kingdom;

This is the author manuscript accepted for publication and has undergone full peer review but has not been through the copyediting, typesetting, pagination and proofreading process, which may lead to differences between this version and the [Version of Record](#). Please cite this article as [doi: 10.1111/epi.13821](https://doi.org/10.1111/epi.13821)

This article is protected by copyright. All rights reserved

(Elger) Department of Epileptology, University of Bonn Medical Centre, Bonn, Germany;

** (Engel) Neurology, Neurobiology, and Psychiatry and Biobehavioral Sciences, UCLA Seizure Disorder Center, David Geffen School of Medicine at UCLA, Los Angeles, California, U.S.A.;

†† (Forsgren) Department of Pharmacology and Clinical Neuroscience/Neurology, Umea University, Umea, Sweden;

‡‡ (French) Department of Neurology, NYU School of Medicine, New York, New York, U.S.A.;

¶¶ (Hesdorffer) GH Sergievsky Center and Department of Epidemiology, Columbia University, New York, New York, U.S.A.;

(Lee) Yonsei Epilepsy Research Institute, Yonsei University College of Medicine, Seoul, Korea;

*** (Mathern) Departments of Neurosurgery and Psychiatry & BioBehavioral Medicine, Mattel Children's Hospital, David Geffen School of Medicine, University of California, Los Angeles, California, U.S.A.;

††† (Moshe) Saul R. Korey Department of Neurology, Dominick P. Purpura Department of Neuroscience and Department of Pediatrics, Laboratory of Developmental Epilepsy, Montefiore/Einstein Epilepsy Management Center, Albert Einstein College of Medicine and Montefiore Medical Center, Bronx, New York, U.S.A.;

‡‡‡ (Perucca) Department of Internal Medicine and Therapeutics University of Pavia and C. Mondino National Neurological Institute, Pavia, Italy;

§§§ (Scheffer) Departments of Medicine and Paediatrics, Florey Institute, Austin Health and Royal Children's Hospital, The University of Melbourne, Melbourne, Victoria, Australia;

¶¶¶ (Tomson) Department of Clinical Neuroscience, Karolinska Institutet, Stockholm, Sweden;

(Watanabe) Department of Psychiatry, National Center of Neurology and Psychiatry, Tokyo, Japan; and

**** (Wiebe) University of Calgary, Calgary, Alberta, Canada

Address correspondence to

Robert S. Fisher

Department of Neurology and Neurological Sciences

Stanford University School of Medicine

213 Quarry Road, Palo Alto, CA 94304-5979, U.S.A.

E-mail: robert.fisher@stanford.edu

Revision of the operational definition of epilepsy¹ has served in part as motivation to gather welcome new data on the time required for epilepsy to be considered in remission. In a recent study, Sillanpää² and colleagues followed 148 patients with childhood onset epilepsy. For certain subgroups, the 5-year remission was as predictive of a long-term remission as was a 10-year remission. However, for other subgroups, the 10-year remission was more predictive of long-term remission. For example, as estimated from their figure 3, both for all patients and for patients with generalized epilepsy, 5-year remissions with the last two years off anti-seizure medicines were associated with subsequent relapses in about 15%, with the 95% confidence

intervals extending above 20%. In comparison, all patients and patients with generalized seizures who were 10-years seizure-free with the terminal 5 years off anti-seizure medicines had a relapse rate less than 5% and an upper limit of confidence intervals around 15%.

The traditional epilepsy definition of two unprovoked seizures more than 24 hours apart was modified to make it possible to “outgrow” epilepsy. What seizure-free period of time would designate a likely condition of no future seizures? The Task Force recognized that relapse rates for people with epilepsy probably always remains above baseline general population levels. For that reason, the Task Force avoided the word “cure.” We employed “resolved” as a descriptor, rather than “remission,” which some people associate with cancer. In the absence of solid data, the Task Force chose the conservative time period of 10 years seizure-free and 5 years off anti-seizure medicines.

The study by Sillanpää ² is one investigation with a relatively small sample size and subjects having limited etiologies of epilepsy, in that all had childhood-onset epilepsy. Relapse rates might vary considerably among different populations with varied epilepsy etiologies. Self-limited age-dependent syndromes and other special etiologic categories might merit separate consideration. For these reasons, the Task Force and the ILAE consider it premature to alter the published definition of epilepsy ¹ but we can envision doing so in the future after further accumulation of broadly applicable data and larger cohort sizes from which to extrapolate findings.

None of the co-authors have a conflict of interest related to the basic definition of epilepsy.

‘We confirm that we have read the Journal’s position on issues involved in ethical publication and affirm that this report is consistent with those guidelines.

1. Fisher RS, Acevedo C, Arzimanoglou A, et al. ILAE official report: a practical clinical definition of epilepsy. *Epilepsia* 2014;55:475-482.
2. Sillanpää M, Schmidt D, Saarinen MM, et al. Remission in epilepsy: How long is enough? *Epilepsia* 2017;58:901-906.

Author Manuscript