Aetiologies and characteristics of Refractory Status Epilepticus cases in different areas of the world: results from a Global Audit.

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Key points

- We collected information about 776 cases of RSE in 50 countries around the world

- Good outcome was associated with younger age and a prior history of epilepsy; aetiology strongly influenced the outcome.

- Patients from Asia were younger, more frequently presented with convulsive SE, and were more frequently affected by infectious aetiologies

- Important differences exist among patients with RSE from different regions of the world, but these do not seem to influence patient outcomes

Figure legend:

Fig. 1: Map of involved countries

Fig. 2: Origin of patients

Fig. 3: Outcome of SE episodes

Fig. 4: Neurological outcome of patients at the end of anaesthesia

Abstract

In order to describe the demographics, aetiologies, types of status epilepticus (SE) and outcomes in people with refractory and super-refractory status epilepticus from around the world, we collected prospectively cases of refractory status epilepticus (RSE) requiring continuous intravenous anaesthetic drugs (CIVADs) in an intensive care unit setting, through online questionnaires using "active surveillance". We collected information about 776 cases of RSE in 50 countries over 4 years. Control of SE was achieved in 74% of the cases. Neurologic outcomes were poor in 41% of patients and 24% died. Good outcome was associated with younger age and a prior history of epilepsy. Aetiology strongly influenced the outcome. Patients from Asia were younger, more frequently presented with convulsive SE, and were more frequently affected by infectious aetiologies when compared with patients from Europe and the Americas. Despite these differences, outcomes were similar in all countries. Demographics of patients with RSE in a global audit are similar to those in prior single centre series providing evidence of generalizability of those studies. Important differences exist among patients with RSE from different regions of the world, but these do not seem to significantly influence patient outcomes.

Introduction

Refractory status epilepticus (RSE) is a dangerous condition, with a mortality rate of 24-38% in recent series, ^{1, 2} higher in prolonged episodes.³A generally accepted definition of RSE is a seizure that persists after 2 antiseizure drugs, typically including a benzodiazepine. At this stage, the standard treatment is with continuous intravenous anaesthetic drugs (CIVADs), in order to promptly stop seizure activity, prevent long-term neuronal damage, further refractoriness ^{4, 5} and severe acute systemic consequences, especially in convulsive SE. The current evidence base guiding optimal management of RSE is mostly based on small series given the rarity of the condition. ⁶ For these reasons, there has been an increasing interest in multinational registries.^{7, 8} The underlying aetiology of SE is considered the most important prognostic factor determining outcome.⁹ Apart from the treatment, aetiology itself significantly differs in developing countries as compared to the western world, with acute symptomatic aetiologies being more frequent in developing countries.^{10,11} In this study, we prospectively collected information about cases of RSE in different regions of the world.

Methods

Details about the audit procedures have been published previously.¹² Briefly, this was an anonymized online registry, collecting information prospectively from neurologists and intensivists caring for patients with RSE requiring admission to an intensive care unit (ICU), , defined as "SE not responding to first-line therapy and requiring general anaesthesia in an ICU", through online questionnaires. The "active surveillance" method, which utilized monthly reminders sent to all participating physicians, ensured maximal reporting. A modified Rankin scale (mRS) of 0-3 was considered a good outcome.¹³

All data were analysed using statistical software (IBM SPSS Statistics, version 20). When comparing continuous variables, Student's *t*-test and Mann-Whitney test were used. The analysis of categorical variables was performed using Chi-square and Fisher exact and analysis between groups with Anova and Kruskal Wallis.

Results

The data collection started on the 1st of March 2013 and was terminated after 4 years. In total, 776 cases were collected from 166 different physicians (see list of all contributors in appendix 1). A map of the 50 countries involved is shown in Figure 1, and the number of cases contributed per country in Figure 2. Patients were from Europe (n=408, 56%), Asia (n=169, 23%), the Americas (n=131, 18%), Australia and New Zealand (n=17, 2%) and Africa (n=9, 1%). The clinical characteristics of patients are summarized in Table 1. The majority of patients (n=x, 63%) had no history of epilepsy and the most common single etiology was cryptogenic (n=200, 26.1%). Among those with cryptogenic RSE, 78 (39%) had a positive history of epilepsy was uncertain.

SE was convulsive in 55% of cases, non-convulsive in 19%, convulsive evolving to nonconvulsive in 21%, of other semiology (epilepsia partialis continua, absence status, other) in 4%. Mean duration of ICU stay was 18.41 ± 22.8 days.

Regional differences

There were too few patients from Africa and Oceania to justify subgroup analyses. Patients from Asia were significantly younger than those from Europe and the Americas (Mean age 22.4, 48.2, and 40.5 years, p<0.001) and more frequently presented with convulsive SE compared with non-convulsive forms (71%, 53% and 44%, p<0.001). The ICU duration was longer in Asia (mean 22.8 ± 24.1 days) than in Europe (16.3 \pm 18.9 days, p<0.05) or in the Americas (19.31 \pm 26.3 days).

There were some notable differences regarding aetiologies of SE (Table 2). In Asia, the most frequently reported aetiology was infectious (n=59, 30.1%), compared with while this represented only (n=23, 15.4%) of cases in the Americas and (n=56, 12.3%) in Europe. In particular, the percentage of cases with acute encephalitis was significantly higher in Asia (n=41,

20.9%), than in Europe (n=26, 5.7%) or the Americas (n=7, 4.7%), p<0.01). Vascular aetiologies were more frequent in Europe (n=75, 16.6%) than in Asia (n=11, 5.6%), p<0.01. There was a non significant trend towards a higher incidence of traumatic etiologies in Europe (n=28, 6.2%) and the Americas (n=7, 4.7%) than in Asia (n=1, 0.5%).

Outcomes

In 686 cases information about the outcome was provided: 510 patients (74%) recovered, 148 patients (22%) died during treatment, and 28 patients (4%) had therapy actively withdrawn. The neurological status of the patients at the end of anaesthesia was good in 35% of patients, poor in 41%, and 24% of patients died. Outcomes in patients with a long-term outcome data provided (n=208) are shown in Table 3. There was a higher proportion of patients with a better outcome at six-month follow up.

No differences were found in outcome with respect to gender or type of SE. Prior history of epilepsy and younger age was positively associated with recovery from SE (p<0.001). As expected, aetiology strongly influenced outcome (Table 4). Patients with post-anoxic SE had the worst outcomes, as did those with metabolic aetiologies or acute encephalitis when compared with other etiologies. Patients with antiseizure drug withdrawal as the aetiology of SE had the best outcomes, as did those with a genetic or chromosomal aetiology.

In the analysis between geographical regions, we did not find any significant differences in rate of success in controlling SE, or in the neurological outcome of patients (fig. 3 and 4).

Discussion

In studies on RSE, the setting (ICU, academically-driven, general hospital, rural hospital) and the geographical region may have an important impact on the results.^{10, 14, 15} Observational studies are almost the only ones available in RSE, and the validity of such studies depends on the range of participation and the quality of their data. This is to our knowledge the largest and most widely collected series of RSE although we acknowledge limitations in drawing conclusions about associations given the non-systematic method of collection. The characteristics of patients with RSE in our global audit are similar to those in prior single centre series, providing evidence of generalizability of those studies.

To ensure accuracy and completeness of the data, we modified the format of the questionnaires several times, but as increasing complexity reduced the number of cases reported and we had to compromise on a limited data set. We also did not make any complex statistical analysis of the data as a registry carries intrinsically constituted limitations. ¹⁶ Despite these limitations, this registry adds important information to our knowledge of the demographics, types and aetiologies of RSE around the world.

As expected, aetiology of SE and characteristics of patients can significantly differ in Asian countries as compared to the western world.^{10, 14} In this study, patients from Asia were younger: this could simply reflect the lower mean age of the Asian population. We found less prevalence of non-convulsive status in Asian countries, presumably because of lower availability of continuous EEG monitoring. Globally, cryptogenic SE was the most frequent cause of RSE around the world. As most of these cases had no prior history of epilepsy, future research must focus on identifying causes of cryptogenic RSE and in particular on autoimmune aetiologies, which probably account for a significant number of these cases. As expected, anoxic SE has the worst outcome; acute encephalitis and metabolic aetiologies are also associated with poor outcomes.

Aetiologies differ remarkably among continents. In Asian countries infectious aetiologies were the most commonly reported, with acute encephalitis occurring significantly more frequently than in other regions of the world. Acute encephalitis has been associated with refractoriness to treatment and higher mortalities in prior studies.^{17, 18, 19} A fascinating finding of this audit is that, despite such great differences in patients' and SE characteristics, outcomes around the world were largely similar. It is possible that such great differences in aetiologies and SE characteristics are somehow compensated by other factors, like younger age, and the nature of this study does not allow for a full exploration of these relationships. Data from this audit have been reviewed with all the participant doctors at the 6th London-Innsbruck Colloquium on Status Epilepticus and Acute Seizures, where further analyses have

been planned and future research discussed.

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Disclosure of conflicts of interest

None of the authors has any conflict of interest to disclose.

Ethical Publication Statement

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.

References

1. Delaj L, Novy J, Ryvlin P, et al. Refractory and super-refractory status epilepticus in adults: a 9-year cohort study. Acta Neurol Scand 2017;135:92-99.

2. Sutter R, Marsch S, Fuhr P, et al. Mortality and recovery from refractory status epilepticus in the intensive care unit: a 7-year observational study. Epilepsia 2013;54:502-11.

3. Cooper AD, Britton JW, Rabinstein AA. Functional and cognitive outcome in prolonged refractory status epilepticus. Arch Neurol 2009;66:1505–1509.

4. Hillman J, Lehtimäki K, Peltola J, et al. Clinical significance of treatment delay in status epilepticus. Int J Emerg Med. 2013 Feb 27;6:6.

5. Kapur J. Rapid seizure-induced reduction of benzodiazepine and Zn2 sensitivity of hippocampal dentate granule cell GABAA receptors. J Neurosci. 1997;17:7532.

6. Shorvon S, Ferlisi M. The treatment of super-refractory status epilepticus: a critical review of available therapies and a clinical treatment protocol. Brain 2011;134:2802-18.

7. Ferlisi M, Hocker S. What can we learn from status epilepticus registries? Epilepsia. 2013;54 Suppl 6:72-3.

8. Kellinghaus C, Lang N, Rossetti AO, et al. Making SENSE--Sustained Effort Network for treatment of Status Epilepticus as a multicenter prospective registry. BMC Neurol 2015;15:230.

9. Neligan A, Shorvon SD. The frequency and prognosis of convulsive status epilepticus of different causes: a systematic review. *Arch Neurol* 2010; 67: 931-40.

10. Hassan H, Rajiv KR, Menon R, et al. An audit of the predictors of outcome in status epilepticus from a resource-poor country: a comparison with developed countries. Epileptic Disord. 2016 Jun 1;18:163-72.

11. Murthy JMK, Jayalaxmi SS, Kanikannan MA. Convulsive status epilepticus: clinical profile in a developing country. *Epilepsia* 2007; 48: 2217-23

12. Ferlisi M, Hocker S, Grade M, et al. Preliminary results of the global audit of treatment of refractory status epilepticus. Epilepsy Behav. 2015;49:318-24.

13. Bamford JM, Sandercock PA, Warlow CP, et al. Interobserver agreement for the assessment of handicap in stroke patients. Stroke 1989;20:828.

14. Sinha S, Prashantha DK, Thennarasu K, et al. Refractory status epilepticus: a developing country perspective. *J Neurol Sci* 2010; 290: 60-5.

15. Alvarez V, Lee JW, Westover MB, et al. Therapeutic coma for status epilepticus: Differing practices in a prospective multicenter study. Neurology 2016;87:1650-1659.

16. Byar DP. Why data bases should not replace randomized clinical trials. Biometrics. 1980;36:337-42.

17. Holtkamp M, Othman J, Buchheim K, et al. Predictors and prognosis of refractory status epilepticus treated in a neurological intensive care unit. J Neu-rol Neurosurg Psychiatry. 2005;76:534-539.

18. Jayalakshmi S, Ruikar D, Vooturi S, et al. Determinants and predictors of outcome in super refractory status epilepticus-a developing country perspective. Epilepsy Res 2014;108:1609-17.

19. Sahin M, Menache CC, Holmes GL, et al. Outcome of severe refractory status epilepticus in children. Epilepsia 2001; 42: 1461-1467.