

Anti-NMDAR encephalitis

A single-center, longitudinal study in China

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Abstract

Objective

To describe the detailed clinical characteristics, immunotherapy, and long-term outcomes of patients with anti-NMDA receptor (NMDAR) encephalitis in China.

Methods

A single-center, prospective study. Patients who met the diagnostic criteria were enrolled from 2011 to 2017 and followed up. The clinical features, treatment, and long-term outcomes were collected prospectively. Factors affecting the long-term prognosis were analyzed.

Results

The study included 220 patients. The most common clinical presentations were psychosis (82.7%) and seizures (80.9%). Of the patients, 19.5% had an underlying neoplasm; of which ovarian teratoma was 100% of tumors in females and only one male had lung cancer. Most patients (99.5%) received first-line therapy (glucocorticoids, IV immunoglobulin, or plasmapheresis alone or combined), and only 7.3% received second-line immunotherapy (rituximab, cyclophosphamide alone, or combined). Long-term immunotherapy (mycophenolate mofetil or azathioprine >1 year) was administered to 53.2% of patients. During the first 12 months, 207 (94.1%) patients experienced improvement, and 5 (2.3%) died, whereas 38 (17.3%) experienced relapses. At 12-month follow-up, 92.7% had favorable clinical outcomes (modified Rankin Scale score ≤ 2).

Conclusions

Patients in China present with psychosis and seizure frequently but have a low percentage of underlying neoplasms. Re-enforced first-line immunotherapy is effective in managing anti-NMDAR encephalitis in the acute phase. Although relapse is relatively common, with combined first-line and long-term immunotherapy, most patients reached favorable outcomes.

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Glossary

AE = autoimmune encephalitis; **AQP4** = aquaporin-4; **AZA** = azathioprine; **CTX** = cyclophosphamide; **HSV** = herpes simplex virus; **ICU** = intensive care unit; **IQR** = interquartile range; **IVIG** = IV immunoglobulin; **MMF** = mycophenolate mofetil; **MOG** = myelin oligodendrocyte glycoprotein; **mRS** = modified Rankin Scale; **MTX** = methotrexate; **NMDAR** = NMDA receptor; **PE** = plasmapheresis; **PUMCH** = Peking Union Medical College Hospital; **RTX** = rituximab.

Anti-NMDA receptor (NMDAR) encephalitis is the most common type of autoimmune encephalitis (AE), which is associated with autoantibodies against neurosurface or synaptic antigens.^{1–3} Since its first report in 2007,¹ with the proposed clinical approach,³ increasing numbers of anti-NMDAR encephalitis cases were identified.

Differences in clinical characteristics and treatment strategies of anti-NMDAR encephalitis were reported among races and countries.^{4–10} Main barriers to AE management in China consist of the availability of referral centers, the timeliness of correct diagnosis, and financial concerns.¹¹ In 2017, China proposed a domestic consensus on the management of AE, aiming to increase awareness of the disease and determine the optimal treatment for Chinese patients.¹² However, limited data of clinical characteristics and long-term prognosis of Chinese anti-NMDAR patients are available owing to few reports with small sample size.^{9,11,13,14}

Taking the advantage of Peking Union Medical College Hospital (PUMCH) as the national referral center for complicated disease, a prospective anti-NMDAR encephalitis disease cohort was established to describe the clinical characteristics, treatment regimen, and long-term outcomes of patients with anti-NMDAR encephalitis in China.

Methods

Study design and population

In this study, patients with anti-NMDAR encephalitis were enrolled consecutively at PUMCH between May 2011 and December 2017. The inclusion criteria were as follows: (1) acute onset of 1 or more of the 8 major groups of manifestations: psychosis, memory deficit, speech disturbance, seizures, movement disorder, loss of consciousness, autonomic dysfunction, and central hypoventilation; (2) CSF tested positive for NMDAR antibodies (cell-based assay [EUROIMMUN, Lübeck, Germany]); and (3) reasonable exclusion of other disorders.

To better understand the clinical characteristics, we also recorded the co-occurrence of fever, headache, arrhythmia, intensive care unit (ICU) admission, and other atypical symptoms. Because of limited resources of the hospital and financial concerns the patients, the absolute indications for ICU admission included severe anti-NMDAR encephalitis with ovarian teratoma requiring surgical operation, status epilepticus, mechanical ventilation requirement, and hemodynamic instability.

Demographic data and ancillary tests results were recorded, including age at onset, sex, disease course, CSF tests results, MRI, and EEG results. All patients were screened at least once for systemic tumors at onset. Patients with tumors underwent tumor removal. Immunotherapy included first-line (corticosteroids, IV immunoglobulin [IVIG], or plasmapheresis [PE] alone or combined) and second-line (rituximab [RTX] and cyclophosphamide [CTX] alone or combined) immunotherapies.^{6,7} Long-term immunotherapy (mycophenolate mofetil [MMF] or azathioprine [AZA] >1 year) and other immunotherapy (intrathecal methotrexate [MTX]) were also administered.^{15,16}

Patients were followed regularly in local hospitals or PUMCH neurology clinics. Treatment effects and long-term outcomes were assessed using the modified Rankin Scale (mRS). A poor response was defined as no improvement in the mRS score or as an mRS score ≥ 4 for 4 weeks; clinical improvement was defined as a decrease in the mRS score ≥ 1 point from that at the previous visit; relapse was defined as an exacerbation of previous symptoms or the occurrence of new symptoms after being stable for 2 months. Long-term favorable outcome was defined as an mRS score ≤ 2 , and poor outcome was defined as an mRS score > 2 at the end of follow-up.

Statistical analysis

Statistical analyses were performed using SPSS IBM 20.0. GraphPad Prism 6.0 was used to generate figures. Quantitative data with normal distributions are presented as mean \pm SD, otherwise as medians with the interquartile range (IQR). The mRS scores before and after treatment were compared using the Wilcoxon test. Symptoms and demographic data were analyzed using the χ^2 test or Fisher exact test for categorical variables and Mann-Whitney *U* test for continuous variables. Factors affecting outcome were assessed using binary logistic regression analysis. Kaplan-Meier curves with log-rank were used to analyze relapse frequency. *p* < 0.05 was considered significant.

Standard protocol approvals, registrations, and patient consents

This study was approved by the Ethics Committee of PUMCH (JS-891), and informed consent was obtained from each patient. All the data analyzed in the study were strictly anonymous.

Data availability

Anonymized data not published within this article will be made available by request from any qualified investigator.

Results

Clinical characteristics

A total of 220 patients were enrolled, and all were tested for anti-NMDAR antibody with paired CSF and serum. All patients (100%) were positive for anti-NMDAR antibodies in CSF, and 157 (71.4%) were positive in serum. Review of the enrolled patients indicated that all patients met the diagnostic criteria proposed by Graus et al.³

The median age at onset was 21 (range 5–72) years, with 69 (31.4%) of the patients younger than 18 years. Overall, 143 (65.0%) patients were females, and 77 (35.0%) were males. Tumors were found in 43 (19.5%) patients: 42 females with ovarian teratomas and 1 male with lung cancer. All the female patients with ovarian teratomas underwent tumor removal, whereas the patient with lung cancer was treated with palliative therapy by an internist and later died of cancer. In addition, 8 patients were found to have prominent moles on the body surface, which were later resected. Pathologically, the moles were compound or intradermal nevi.^{17,18}

The most common clinical manifestations of the anti-NMDAR encephalitis were psychosis (182, 82.7%) and seizures (178, 80.9%). Movement disorder presented more often in patients younger than 18 years than adult patients (38/69, 55.1% vs 56/151, 37.1%, $p = 0.01$). Table 1 describes the patients' clinical characteristics in detail.

Figure 1 shows the distribution of female patients according to age and presence vs absence of tumor; the peak co-occurrence of ovarian teratoma was between 19 to 24 years.

Ancillary test results

All patients underwent brain MRI at onset, and 79 (35.9%) had abnormal fluid-attenuated inversion recovery sequence signals, including 31 (14.1%) in the medial temporal lobe. Other involved areas included the frontal, parietal, and occipital cortices, diencephalon, cerebellum, and brainstem. Eleven (5.0%) patients had demyelinating lesions, of whom 4 were positive for aquaporin-4 (AQP4) antibody and 5 for myelin oligodendrocyte glycoprotein (MOG) antibody. Abnormal EEG findings were seen in 113 (51.4%) patients: 102 (46.4%) had slow activity and 14 (6.4%) epileptic discharges. However, as most patients received short-duration EEG instead of video EEG of 24 hours or longer, the percentage of delta-brush abnormality was unable to assess.

Repeated lumbar punctures were performed for diagnosis and evaluation, and the CSF results at onset before immunotherapy were collected and analyzed. The analysis showed a median opening pressure of 170 (IQR 150–280) mmH₂O. Of note, 81.3% of the patients had pleocytosis, the median white blood cell count was 14.0 (IQR 7.0–22.5) $\times 10^6/L$, and 90.9% were of mononuclear cells (i.e., lymphocyte and monocytes). The protein level was elevated in 29.7% of the patients. Table 2 summarizes the main ancillary tests of the patients.

Table 1 Clinical characteristics of patients with anti-NMDA receptor encephalitis

Clinical characteristics	No. of patients (percentage)			p Value
	All (n = 220)	Age <18 y (n = 69)	Age >18 y (n = 151)	
Median age, range (y)	21 (5–72)	15 (5–17)	26 (18–72)	
Female	143 (65.0)	41 (59.4)	102 (67.5)	0.24
Psychosis	182 (82.7)	57 (82.6)	125 (82.8)	0.98
Seizures	178 (80.9)	56 (81.2)	122 (80.8)	0.95
Fever	126 (57.3)	41 (59.4)	85 (56.3)	0.67
Decreased level of consciousness	117 (53.2)	35 (50.7)	82 (54.3)	0.62
Memory deficit	106 (48.2)	28 (40.6)	78 (51.7)	0.13
Speech disturbance	100 (45.5)	30 (43.5)	70 (46.4)	0.69
Movement disorder	94 (42.7)	38 (55.1)	56 (37.1)	0.01
Sleep disorder	93 (42.3)	27 (12.3)	66 (30.0)	0.52
Headache	86 (39.1)	21 (30.4)	65 (43.0)	0.08
Admission to the ICU	68 (30.9)	18 (26.1)	50 (33.1)	0.30
Mechanical ventilation	44 (20.0)	12 (17.4)	32 (21.2)	0.51
Central hypoventilation	39 (17.7)	9 (13.0)	30 (19.9)	0.22
Arrhythmia	34 (15.5)	12 (17.4)	22 (14.6)	0.59
Limb weakness	24 (10.9)	9 (13.0)	15 (9.9)	0.49
Ataxia	11 (5.0)	4 (5.8)	7 (4.6)	0.74
Ovarian teratoma of female patients	42 (29.4)	9 (22.0)	33 (32.4)	0.22

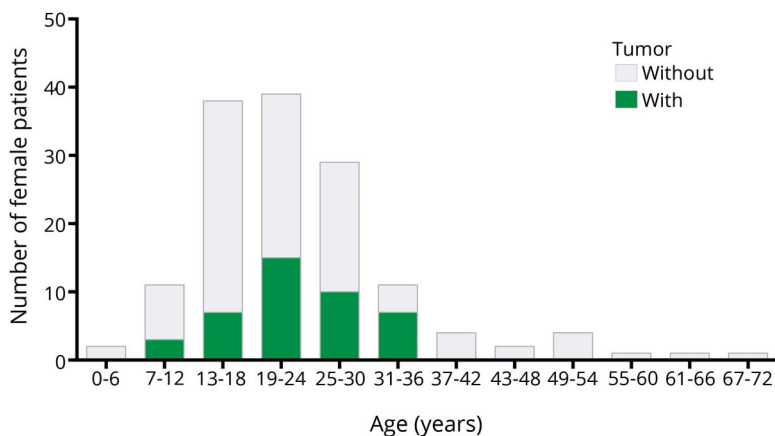
Abbreviation: ICU = intensive care unit.

Improvements in diagnosis accuracy and general hospitalization status

Underrecognition and misdiagnosis were inevitable in the past, occurring at the initial visit to a local physician or admission to our hospital. In 2011, 4 patients were identified, taking a median duration of 9 (IQR 1–36) months before diagnosis was made. The misdiagnosis rate then was 75.0%, with most cases misdiagnosed as viral encephalitis.

With increased awareness of AE, screening for anti-NMDAR antibodies at admission is now required for all cases with suspected encephalitis. By 2017, the misdiagnosis rate had decreased to 15.4%. Overall, 30 patients were misdiagnosed with viral encephalitis, 10 with schizophrenia, 2 with epilepsy, and 1 each with cerebral angiitis, cerebral vascular events, and tuberculosis. The percentages of correct diagnosis at the initial hospital visit over misdiagnosis by calendar year were shown in figure 2, indicating a growing

Figure 1 Distribution of female patients by age and presence or absence of ovarian teratoma



number of referred patients and increased correct diagnosis identified over the years.

The median duration from onset to diagnosis was 2 (IQR 1–4) weeks. Immunotherapy was initiated the same day diagnosis was made and sometimes even before diagnosis at local hospitals as empirical treatment. The median length of hospitalization was 26 (IQR 14–42) days. However, severe patients who required intensive care stayed in the hospital for up to 117 days.

Treatment outcomes

Overall, 219 (99.5%) patients received first-line immunotherapy, in most cases a combined regimen of repeated steroids and IVIG. A total of 208 (94.5%) patients received steroids, of whom 103 (46.8%) received pulsed IV methylprednisolone. IVIG was administered to 199 (90.5%) patients, and 7 (3.2%) patients underwent PE. Second-line immunotherapy was administered to only a small proportion of the patients, usually because of the off-label use of RTX in AE, cost, IV and hospitalization requirements, and concerns about side effects. Twelve (5.5%) patients received RTX, and 4 (1.8%) received CTX. Long-term immunotherapy was administered mainly in patients who were enrolled later, as an add-on therapy for severe or refractory patients in the acute phase, or as maintenance therapy to prevent and manage relapses. In general, MMF was administered in 109 (49.5%) patients, 55 of whom at onset and 54 after relapse, and AZA was administered in 8 (3.6%) patients. In addition, intrathecal MTX was given to 8 (3.6%) severe patients.

During the first 12 months, 207 (94.1%) patients experienced improvement, 8 (3.6%) were stable, and 5 (2.3%) patients died. All survival patients were followed for at least a year (range 12–72 months). At 12-month follow-up, 204 (92.7%) patients had attained satisfactory neurologic function (mRS score of 0, 1, and 2 in 144, 47, and 13 patients, respectively) compared with 23 (10.5%) patients with an mRS score ≤ 2 at onset, as

shown in figure 3. The median mRS score at the last follow-up was 0 (IQR 0–1), which was significantly lower than the score of 4 (IQR 3–5) at onset ($Z = -12.67$, $p < 0.0001$).

Table 3 summarizes the comparisons between patients with favorable and poor clinical outcomes. Of interest, more patients with speech disturbance were identified in the favorable outcome group ($p = 0.03$). Further analysis indicated that patients with speech disturbance presented to the neurologist earlier than those without (20 days [IQR 12–34 days] vs 31 days [IQR 19–58 days], $p = 0.008$, $Z = -2.665$). However, age at onset, the rates of decreased consciousness level, central hypoventilation, ICU admission, mechanical ventilation, ovarian teratoma, and relapse were higher in the poor outcome subgroup, although none of these differences reached significance.

Relapse

During the first 12 months, 38 (17.3%) patients experienced a first relapse. Their median age was 21 (IQR 16–37) years, and the median duration from onset to the first relapse was 7 (IQR 5–10) months. Of the relapsed patients, 25 (65.8%) were female, and 5 had ovarian teratomas at disease onset.

Thorough clinical and laboratory examinations were conducted to rule out other etiologies and to validate the diagnosis. Two patients were treated at local hospitals before antibodies were tested, whereas 10/36 (27.8%) patients showed elevated antibody titer at relapse during serial serum antibody monitoring. Of the 17 patients who underwent lumbar puncture, CSF antibodies were detected in 15 (88.2%) patients. MRI was abnormal in 8 (21.1%) patients, and an ovarian teratoma was detected in only 1 (2.6%) patient at relapse. A delay in treatment was associated with relapse ($p < 0.01$).

However, neither tumor status ($p = 0.58$) nor treatment regimen ($p = 0.34$) was statistically associated with relapse frequency (figure e-1, links.lww.com/NXI/A157).

Table 2 Summary of the main ancillary tests results

	Number (percentage)	
Brain MRI		
Total with abnormal findings	79 (35.9)	
Medial temporal lobe	31 (14.1)	
Frontal lobe	20 (9.1)	
Parietal lobe	13 (5.9)	
Occipital lobe	13 (5.9)	
Diencephalon	2 (0.9)	
Cerebellum	3 (1.4)	
Brainstem	6 (2.7)	
Demyelinating lesion	11 (5.0)	
Anti-AQP4 Ab(+)	4 (1.8)	
Anti-MOG Ab(+)	5 (2.3)	
EEG		
Total with abnormal findings	113 (51.4)	
Epileptic discharges	14 (6.4)	
Slow activity	102 (46.4)	
CSF analysis (information of 64 patients)	Median (IQR)	Number (percentage)
Opening pressure (mmH₂O)	170 (150–280)	
WBC (×10⁶/L)	14.0 (7.0–22.5)	
Pleocytosis		52 (81.3)
Percentage of mononuclear cells (%)	90.9 (83.1–99)%	
Protein (g/L)	0.30 (0.21–0.42)	
	≥0.45	19 (29.7)
Positive SOB		16 (25.0)
Glucose (mmol/L)		3.5 (3.1–4.2)
Chloride (mmol/L)		123 (120–125)

Abbreviations: AQP4 = aquaporin-4; IQR = interquartile range; MOG Ab = myelin oligodendrocyte glycoprotein antibody; SOB = specific oligoclonal band; WBC = white blood cell count.
 Mononuclear cells: lymphocytes and monocytes.
 CSF-SOB is defined as bands that are present in CSF but are absent in serum.
 Positive SOB is defined as at least 1 specific band in the CSF.

All relapsed patients underwent reinitiation of the first-line immunotherapy, and 18 patients were also given long-term MMF. Subsequently, 12 (31.6%) patients experienced further relapses (range 2–4 episodes). There was no significant difference in the occurrence of subsequent relapse between MMF-treated patients and other patients (4/18, 22.2% vs 8/20, 40.0%, $p = 0.31$).

Subsequent relapses were similar to, or worse than, the initial episodes in only 3 (7.9%) patients. The initial hospitalization

duration was 25 (IQR 14–43) days, and the subsequent hospitalizations were all shorter than 14 days.

Between 2011 and 2017, 80 (36.4%) patients experienced relapse, and 21 (26.3%) experienced multiple relapses (range 2–4 episodes). Most patients had a first relapse during the first 24 months (64/80, 80.0%). However, relapses up to 6 years after onset were also reported in our cohort.

Discussion

To our knowledge, this is the largest Chinese anti-NMDAR encephalitis cohort to date. In our study, anti-NMDAR encephalitis is predominantly found in females (65.0%) with the median age at onset of 21 years. Most patients presented with psychiatric symptoms and seizures, and younger patients presented more often with movement disorders, which are consistent with previous studies.^{6,7}

However, our study reported a low ICU admission rate. The large cohort study of Titulaer et al.⁷ reported that 75% (435/563) of patients with mRS score ≥ 4 were admitted to the ICU. In our study, because of limited availability of medical recourses and concern of expenses, only 68/133 (51.1%) of the severe patients (mRS score ≥ 4) were admitted to the ICU.

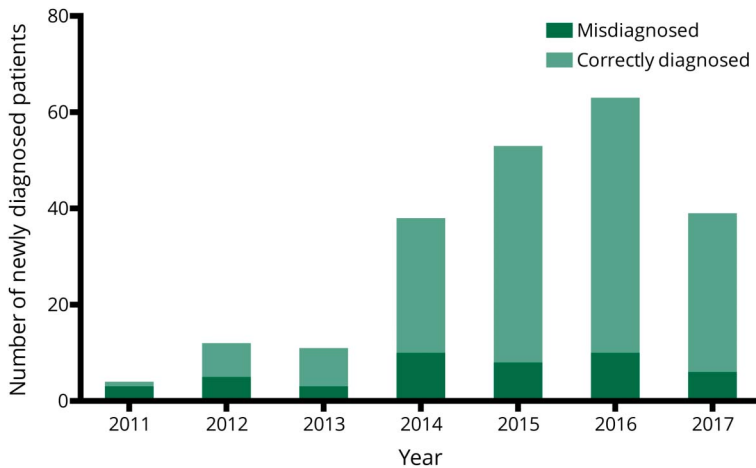
The prevalence of an underlying neoplasm varied among studies. Titulaer et al.⁷ reported that 38% of the patients had a tumor, and Asian patients were more likely (45%) to have a teratoma. However, only 19.5% of the patients in our cohort had a tumor, with 29.4% of the females had an ovarian teratoma. Other studies of Chinese or Asian patients have also reported low prevalence of tumors (Lim et al., 22.7%; Wang et al., 8%; Liu et al., 6.7%; and Zhang et al., 8.1%).^{8,11,13,14} The heterogeneity among reports may be due to sample sizes and selection bias or other factors including genetic backgrounds and epidemiologic reasons, and future studies are required.

In tumor-negative patients, herpes simplex virus (HSV) infection has been reported to be the possible trigger. However, as the gold standard for the infection is PCR testing, which is expensive and time consuming, most patients received empirical treatment before or even without a definite diagnosis of HSV encephalitis. Thus post-HSV anti-NMDAR encephalitis was unable to assess in our current study.

Brain MRI findings provide further evidence that anti-NMDAR encephalitis is a “diffuse encephalopathy.”⁶ Abnormal signals were reported in 35.9% of the patients, predominantly in the medial temporal lobe. Signals in other areas of the cortex, diencephalon, brainstem, and cerebellum were also reported.

Notably, “overlapping syndrome” was identified in 11 (5.0%) patients with both MRI demyelinating lesions and anti-AQP4 or anti-MOG antibodies.¹⁹ Studies have suggested more

Figure 2 Number of patients of correct diagnosis over misdiagnosis at the initial hospital visit by calendar year



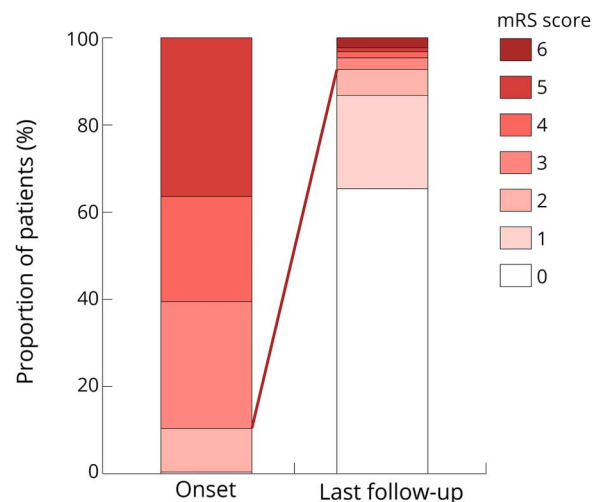
intense immunotherapy requirements and more residual deficits in these patients.^{19,20} In our series, all these patients received long-term immunotherapy with MMF, and 10 (90.9%) had favorable clinical outcomes.

In the management of anti-NMDAR encephalitis, repeated first-line immunotherapy was frequently used in our cohort, whereas second-line immunotherapy was administered in a small portion of patients owing to the off-label use of RTX for AE in China, cost, hospitalization requirements, and concerns about side effect.^{9,11,14,15} However, long-term immunotherapy was administered to 117 (53.2%) patients, including MMF to 109 (49.5%). With combined therapy of re-enforced first-line therapy and long-term immunotherapy, 204 (92.7%) patients reached favorable clinical outcomes, and the median mRS score decreased significantly from 4 to 0 at a follow-up of 12 months. Compared with other reports (Dalmau et al., 77%; Tituaer et al., 79%; Wang et al., 80.4%; Liu et al., 64%; and Zhang et al., 89.2%),^{6,7,11,13,14} we observed more satisfactory clinical outcomes. Speech disturbance was found to be more frequent in the group with favorable outcome. Further analysis indicated that patients with speech distance were diagnosed earlier. Thus, this could be a confounder reflecting better recognition and therefore quicker treatment.

Relapses were relatively common in our cohort. The definition of relapse in our study, along with other proposed definitions,^{7,21,22} is based more on observations and descriptions of clinical symptoms. Nevertheless, thorough examinations are needed to rule out other disorders and validate the diagnosis. When monitoring and evaluating the relapses, MRI was frequently unremarkable. The serum antibody titer did not correlate with the clinical severity perfectly, and some relapsed antibodies were detected only in the CSF, as previously reported.^{6,22} However, serial CSF monitoring may be impractical during follow-up, and better indicators should be identified in future studies. Neither tumor status nor treatment regimen

was associated with relapse frequency statistically, possibly due to the low prevalence of tumor and variety in the treatment regimens used in our cohort. Long-term MMF did not prevent further relapses, possibly because of the relatively small sample size, and the role of long-term immunotherapy warrants further investigation. Although most patients experienced a first relapse within 24 months, relapse 6 years after onset was also reported. Other reports also suggested that AE relapse could occur years after the initial episode.^{21,23,24} Therefore, extended follow-up is essential. Our study has several limitations. As the national referral center for complicated disease, our cohort may be biased by more refractory cases. The analysis for each individual treatment, particularly for long-term immunotherapy, and the assessment of relapses warrants further study. Parameters other

Figure 3 Distribution of mRS scores at onset and last follow-up



mRS = modified Rankin Scale.

Table 3 Comparisons of clinical data of patients with anti-NMDA receptor encephalitis

	Favorable outcome (n = 204)	Poor outcome (n = 16)	Z/ χ^2 value	p Value
Age (y)	21 (16–28)	27 (20–32)	–1.79	0.07
Sex (n, %)				
Male	71 (34.8)	6 (37.5)	0.05	0.83
Female	133 (65.2)	10 (62.5)		
Symptoms (n, %)				
Fever	118 (57.8)	7 (43.8)	1.20	0.27
Headache	80 (39.2)	5 (31.3)	0.40	0.53
Psychosis	170 (83.3)	11 (68.8)	2.16	0.17
Memory deficit	100 (49.0)	5 (31.3)	1.88	0.17
Seizures	165 (80.9)	12 (75.0)	0.33	0.52
Decreased level of consciousness	106 (52.0)	10 (62.5)	0.66	0.42
Movement disorder	87 (42.7)	6 (37.5)	0.16	0.69
Speech disturbance	96 (47.1)	3 (18.8)	4.80	0.03
Central hypoventilation	35 (17.2)	4 (25.0)	0.63	0.49
ICU admission	61 (29.9)	7 (43.8)	1.33	0.27
Mechanical ventilation	40 (19.6)	4 (25.0)	0.27	0.53
Arrhythmia	32 (15.7)	2 (12.5)	0.12	0.73
Ataxia	9 (4.4)	2 (12.5)	2.04	0.19
Limb weakness	22 (10.8)	2 (12.5)	0.05	0.69
Ovarian teratoma^a	37 (27.8)	5 (50.0)	2.21	0.16
CSF analysis of acute phase (n, %)				
Pleocytosis^b	88 (58.7)	4 (44.4)	0.70	0.40
Increased level of protein^c	33 (24.6)	3 (33.3)	0.34	0.56
Positive SOB^d	48 (50.5)	2 (28.6)	1.26	0.26
Abnormal brain MRI signal (n, %)	72 (35.3)	7 (43.8)	0.46	0.50
Abnormal EEG findings (n, %)^e	110 (80.9)	6 (75.0)	0.17	0.68
Diagnosis duration (mo)	1.0	1.0	/	/
Immunotherapy (n, %)				
Steroids	194 (95.1)	14 (87.5)	1.66	1.20
IVIg	184 (90.2)	15 (93.8)	0.22	0.64
MMF	103 (50.5)	6 (37.5)	1.51	0.17
Baseline mRS score >2 (n, %)	183 (89.7)	14 (87.5)	0.08	0.78
Relapsed patient (n, %)	73 (35.8)	7 (43.8)	0.41	0.52

Abbreviations: ICU = intensive care unit; IVIG = IV immunoglobulin; MMF = mycophenolate mofetil; mRS = modified Rankin Scale; SOB = specific oligoclonal band.

^a Analysis of 143 female patients.

^b Analysis of 159 patients with results available.

^c Analysis of 143 patients with results available.

^d Analysis of 102 patients with results available.

^e Analysis of 144 patients with results available.

than the mRS score are required to describe fully the outcomes of anti-NMDAR encephalitis, especially in evaluating the cognitive or behavioral function, which usually remains last in these patients. Despite these limitations, our study adds to the present knowledge of anti-NMDAR encephalitis, and acts as a precursor for future multicenter studies with more comprehensive evaluations, and studies to further evaluate the efficacy of each individual treatment.

We describe the clinical characteristics, immunotherapy regimens, and long-term outcomes of patients with anti-NMDAR encephalitis in China. Repeated first-line therapy is effective in managing acute phase encephalitis, and the efficacy of long-term immunotherapy warrants further study. Although relapses are relatively common, most patients reached favorable outcomes. Further multicenter studies with more advanced study design, more detailed evaluation, and extended follow-up are required.

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Disclosure

The authors report no disclosures. Go to Neurology.org/NN for full disclosures.

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Appendix Authors

Name	Location	Role	Contribution
Xiaolu Xu, MD	Peking Union Medical College Hospital, Beijing, China	Author	Analyzed the data and drafted the manuscript for intellectual content
Qiang Lu, MD	Peking Union Medical College Hospital, Beijing, China	Author	Analyzed the data and drafted the manuscript for intellectual content

Appendix (continued)

Name	Location	Role	Contribution
Yan Huang, MD	Peking Union Medical College Hospital, Beijing, China	Author	Major role in the acquisition of data and analyzed the data
Siyuan Fan, MD	Peking Union Medical College Hospital, Beijing, China	Author	Major role in the acquisition of data and analyzed the data
Lixin Zhou, MD	Peking Union Medical College Hospital, Beijing, China	Author	Analyzed the data and revised the manuscript for intellectual content
Jing Yuan, MD	Peking Union Medical College Hospital, Beijing, China	Author	Analyzed the data and revised the manuscript for intellectual content
Xunzhe Yang, MD	Peking Union Medical College Hospital, Beijing, China	Author	Interpreted the data and revised the manuscript for intellectual content
Haitao Ren	Peking Union Medical College Hospital, Beijing, China	Author	Major role in the acquisition of data; interpreted the data; and revised the manuscript for intellectual content
Dawei Sun, MD	Peking Union Medical College Hospital, Beijing, China	Author	Major role in the acquisition of data; interpreted the data; and revised the manuscript for intellectual content
Yi Dai, MD	Peking Union Medical College Hospital, Beijing, China	Author	Major role in the acquisition of data and interpreted the data
Huadong Zhu, MD	Peking Union Medical College Hospital, Beijing, China	Author	Major role in the acquisition of data and interpreted the data
Yinan Jiang, MD	Peking Union Medical College Hospital, Beijing, China	Author	Major role in the acquisition of data
Yicheng Zhu, MD	Peking Union Medical College Hospital, Beijing, China	Author	Designed and conceptualized study and revised the manuscript for intellectual content
Bin Peng, MD	Peking Union Medical College Hospital, Beijing, China	Author	Designed and conceptualized study and revised the manuscript for intellectual content
Liyang Cui, MD	Peking Union Medical College Hospital, Beijing, China	Author	Designed and conceptualized the study and revised the manuscript for intellectual content
Hongzhi Guan, MD	Peking Union Medical College Hospital, Beijing, China	Author	Designed and conceptualized the study; interpreted the data; and revised the manuscript for intellectual content

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