

# Etiology, early and late term neurological prognosis in cases with meningoencephalitis/encephalitis

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#### **INTRODUCTION**

Encephalitis is defined as inflammation of the brain parenchyma associated with neurological dysfunction.(1) It has a wide etiological spectrum of various infectious and non-infectious causes. It is an important neurological emergency due to high mortality and long-term neurological sequelae.(2,3) Inflammation in encephalitis may result directly from infection of the brain parenchyma or from an autoimmune response to various neuronal receptors. The clinical picture can range from mild nonspecific symptoms such as fever, headache, vomiting, behavioural changes to more severe symptoms such as seizures, confusion and coma with altered consciousness. Timely and accurate diagnosis of encephalitis is very important in terms of enabling early and effective treatment.

### **OBJECTIVES**

In our study, we aimed to reveal the etiological causes of encephalitis in pediatric patients and investigate its long-term neurological outcomes.

## MATERIAL & METHOD

Our research was retrospective and conducted in a single center. The patient population consisted of 93 patients older than 1 month and younger than 18 years old, who were treated as inpatients with the diagnosis of meningoencephalitis/encephalitis at the Department of Pediatrics of Kocaeli University School of Medicine, between January 2011 and January 2021. These patients fulfill the 2013 International Encephalitis Consortium diagnostic criteria.

## **RESULTS**

Boys were 58.1% (n=54) of the patients, girls were 41.9% (n=39), and the median age at diagnosis was 65 months (36-117 months). In two main groups, 54.8% (n=51) were diagnosed with infectious and 45.2% (n=42) with immune-based encephalitis. The etiological cause could be determined in 33% (n=31) with laboratory examinations. The most common infectious agent identified was Herpes simplex virus type 1 (10.8%, n=10). Autoimmune encephalitis was diagnosed in 30.1% (n=28) of the patients and anti-NMDAR antibodies were detected in 4.3% (n=4). ADEM was observed in 15.1% (n=14) of the cases. While four patients (4.3%) experienced a fatal outcome, it was observed that 54.8% (n=51) had developed one or more neurological sequelae during a tracking period of median 59 months (33.5-100 months). The most common sequelae were behavioral/psychiatric disorders (30.6%), epilepsy (24.4%) and motor deficits (20%). In our research, we characterized the presence of intractable seizures requiremented of  $\geq 2$ antiepileptic drugs (OR=19.953, 95%CI [2.252-176.759], p=0.007), and an infectious etiology (OR=10.096, 95%CI [1.18-86.358], p= 0.035) as negative prognosis factors.

		OR	95% CI	р
Age	1 year difference	0,994	0,997-1,011	0,467
Glascow coma	1 point difference	0,641	0,34-1,207	0,168
score				
Fever on appeal	No	1,0 (reference)	0,195-7,59	0,834
	Yes	1,216		
Difficulty in	<2 antiepileptic	1,0 (reference)	2,552-176,759	0,007
control of seizure	drug		-	
	2≥ antiepileptic	19,953		
	drug			
Status epilepticus	No	1,0 (reference)	0,100-11,437	0,956
	Yes	1,068		
Stay in intensive	No	1,0 (reference)	0,165-9,0339	0,845
care unit	Yes	1,221		
Etiologic origin	Immune origin	1,0 (reference)	1,18-86,358	0,035
	Infectious origin	10,096		
OR (odds ratio) CI( confiden	tial index)			

OR (odds ratio) CI( confidential index

in patients with encephalitis

#### **CONCLUSION**

We conclude that viral agents are the main cause, but increased awareness of autoimmunity and understanding the complex nature of encephalitis with serious consequences is an important step in developing effective diagnostic and treatment strategies.

#### **REFERENCES**

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#### Table 1: Logistic regression model analysis of factors associated with neurological prognosis

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