Authors Allen et al ¹	Age range (years) 30-40 ^a	Median age of onset (years) 30-40 ^a	Total number of patients (n) 1	Gender: number of patients (% of n) (See Allen et al)	Number of patients with positive anti- GFAP antibodies in serum (% of n) Not tested	Number of patients with positive anti- GFAP antibodies in CSF (% of n) 1 (100%)	Symptom/clinical syndrome: number of patients (% of n) Flaccid paralysis: 1 (100%)	Treatments used: number of patients (% of n) Image: Control of the second s	Outcomes: number of patients (% of n) At 6 months of onset, still in need of intermittent mechanical ventilation and help with all the activities of daily living
Fang et al ²	21-73	43	16	Male: 8 (50%) Female: 8 (50%)	16 (100%)	10 (62.5%)	Meningoencephalitis: 6 (37.5%) Meningoencephalomyelitis: 5 (31.2%) Encephalomyelitis: 2 (12.5%) Encephalitis: 2 (12.5%) Chronic meningitis: 1 (6.2%)	High dose corticosteroid treatment (acute): 11 (68.75%) Mycophenolate (chronic): 5 (31.2%) Azathioprine (chronic): 1 (6.2%) Unknown: 5 (31,2%)	Relapse during corticosteroid dose tapering: 7 (43.7%) No relapse: 6 (37.5%) Unknown: 3 (31.2%)
Flanagan et al ³	8-103	44	102	Male: 20 (52.7%) Female: 18 (47.3%)	102 (100%)	64 (62.7%)	Encephalitis: 43 (42%) Meningoencephalitis: 13 (12.5%) Myelitis: 11 (10.5%) Encephalomyelitis: 8 (8%) Neuropathy: 8 (8%) Meningitis: 5 (5%) Ataxia: 5 (5%) Meningoencephalomyelitis: 3 (3%) Encephalopathy: 2 (2%) Myasthenia gravis: 2 (2%) Epilepsy: 1 (1%) Dysautonomia: 1 (1%)	High dose corticosteroid treatment (acute): 16 (42.1%) ^b Intravenous immunoglobulins (acute): 6 (15.7%) ^b Plasma exchange (acute): 3 (7.8%) ^b Mycophenolate (chronic): 5 (13.15%) ^b Azathioprin (chronic): 3 (7.8%) ^b Rituximab (chronic): 3 (7.8%) ^b	Unspecified clinical improvement: 18 (47.3%) ^b No improvement: 20 (52.7%) ^b

Supplementary table 1. Literature of autoimmune GFAP astrocytopathy.

Authors	Age range (years)	Median age of onset (years)	Total number of patients (n)	Gender: number of patients (% of n)	Number of patients with positive anti- GFAP antibodies in serum (% of n)	Number of patients with positive anti- GFAP antibodies in CSF (% of n)	Symptom/clinical syndrome: number of patients (% of n)	Treatments used: number of patients (% of n)	Outcomes: number of patients (% of n)
Iorio et al ⁴	6-80	56.5	22	Male: 9 (0.40%) Female: 13 (0.60%)	22 (100%)	8 (36.3%)	Seizures: 7 (31.8%) Altered consciousness: 4 (18.1%) Ataxia: 3 (13.6%) Movement disorder: 2 (9%) Paraparesis: 2 (9%) Behaviour disturbance: 2 (9%). Tetraparesis: 2 (9%) Paraplegia: 1 (4%) Psychosis: 1 (4%) Headache: 1 (4%) Rigor: 1 (4%) Dyskinesias: 1 (4%) Sensory loss: 1 (4%) Sphincter disfunction: 1 (4%) Blurred vision: 1 (4%)	Intravenous methylprednisolone (acute): 15 (68.1%) Intravenous immunoglobulins (acute): 2 (9%) Oral corticosteroid treatment (acute): 1 (4.5%) Plasma exchange (acute): 1 (4.5%)	Unspecified clinical improvement: 16 (72.7%) (Pretreatment mean mRss: 3.26 (95% CI 2.78 to 3.74); last follow-up mRss: 1.47 (95% CI 0.82 to 2.12); p=0.0001) Death: 2 (9%)
Ip et al ⁵ Lee et al ⁶	40-50ª 70-80ª	40-50 ^a 70-80 ^a	1	(See Ip et al) (See Lee et al)	1 Not tested	1	Meningoencephalitis and bilateral sensorineural deafness: 1 (100%) Rigidity, bradykinesia, arm myoclonus and hyperreflexia: 1 (100%)	Intravenous immunoglobulin, intravenous methylprednisolone, and plasma exchange (acute). Rituximab (chronic) Intravenous methylprednisolone (acute).	Treatment improved hearing and upper limb power. Successful weaning of tracheostomy and nasogastric feeding. Walking with little assistance. Promptly and unspecified response to methylprednisolone

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Authors	Age range (years)	Median age of onset (years)	Total number of patients (n)	Gender: number of patients (% of n)	Number of patients with positive anti- GFAP antibodies in serum (% of n)	Number of patients with positive anti- GFAP antibodies in CSF (% of n)	Symptom/clinical syndrome: number of patients (% of n)	Treatments used: number of patients (% of n)	Outcomes: number of patients (% of n)
Long et al ⁷	23-73	54	19	Male: 6 (31.5%) Female: 13 (68.5%)	Not tested	19	Myelitis: 13 (68.4%) Headache: 12 (63.2%) Abnormal vision: 12 (63.2%) Fever: 10 (52.6%) Ataxia: 7 (36.8%) Psychosis: 6 (31.6%) Dyskinesia: 3 (15.8%) Dementia: 3 (15.8%) Seizures: 2 (10.5%) Coma: 1 (5.3%) SIADH: 1 (5.3%) Hiccup and nausea: 1 (5.3%)	Intravenous methylprednisolone (acute): 18 (94.7%) Intravenous immunoglobulins (acute): 11 (57.8%) Oral methylprednisolone (chronic): 16 (84.2%) Mycophenolate (chronic): 2 (10.5%) Azathioprine (chronic): 2 (10.5%)	Not described
Yang et al ⁸	27-69	56	7	Male: 3 (42.8%) Female: 4 (57.2%)	Not tested	7	Headache: 6 (85%) Ataxia: 6 (85%) Dyskinesia: 6 (85%) Dementia: 6 (85%) Coma: 6 (85%) Abnormal vision: 3 (42%) Longitudinally extensive transverse myelitis: 3 (42%)	Intravenous methylprednisolone (acute): 7 (100%) Intravenous immunoglobulins (acute): 6 (85.7%) Plasma exchange and mycophenolate (acute): 1 (14.2%) Mycophenolate (chronic stage): 2 (28.5%) Azathioprine (chronic stage): 2 (28.5%)	Unspecified good outcome: 1 (14.2%) Poor prognosis: 6 (85.8%) Death: 2 (28.5%)

^a Exact data available in the original articles.

 $^{\rm b}\,\%$ of n calculated for 38 patients which have available data.

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