

Supplementary Table 1 Summary of *Angiostrongylus cantonensis* Meningoencephalitis Cases in Children Under 2 Years of Age

Comprehensive literature review (1997–2024) with detailed treatment regimens

Ref.	Year	Country	Age	Clinical Presentation	Corticosteroid Regimen	Anthelmintic Regimen	Outcome
FATAL OUTCOME (n = 3)							
Graber et al. [1]	1997	Mayotte	11 mo	Seizures, coma, flaccid quadriplegia, areflexia, sphincter disturbance	None	None	Fatal (3 weeks; neurovegetative disturbances)
Cooke-Yarborough et al. [2]	1999	Australia (Fiji exposure)	11 mo	Flaccid quadriparesis, dysaesthesia, progressive bulbar dysfunction	IV methylprednisolone (dose NR); initial improvement, relapse after cessation	None	Fatal (day 41; bulbar dysfunction, respiratory failure)
Morton et al. [3]	2013	Australia (Sydney)	10 mo	Lethargy, decreased facial movements, hypotonia, areflexia, progressive weakness	Corticosteroids (dose NR)	Albendazole (dose NR); also received IVIG, imatinib	Fatal (day 27; autopsy confirmed <i>A. cantonensis</i> in brain, cord, lungs)
SURVIVED WITH SEQUELAE (n = 3)							
Morton et al. [3]	2013	Australia (Sydney)	14 mo	Fever, vomiting, dysconjugate eyes, intention tremor, bulging fontanelle, CSF eosinophilia 45%	Corticosteroids × 10 weeks (dose NR)	Albendazole (ceased due to transaminitis); EVD for hydrocephalus	Sequelae: Recurrent hydrocephalus requiring Rickham reservoir; persistent pleocytosis at 6 mo
Evans-Gilbert et al. [4]	2014	Jamaica	19 mo	Refusal to walk/crawl, excessive crying, fever, truncal ataxia, blood and CSF eosinophilia	IV dexamethasone 0.6 mg/kg/dose × 4 doses/day × 3 days; then oral prednisone 1 mg/kg/day × 4 weeks	Albendazole 10 mg/kg/day × 3 weeks	Sequelae: Residual neurological deficits at follow-up
Abe et al. [5]	2022	USA (Hawaii)	11 mo	Fever × 5 days, progressive ascending	IV methylprednisolone	Albendazole 150 mg/day × 6 weeks	Sequelae: Day 30 MRI showed new left frontal infarct, multifocal T2/FLAIR

Ref.	Year	Country	Age	Clinical Presentation	Corticosteroid Regimen	Anthelmintic Regimen	Outcome
				weakness; poor response to IVIG, quadriplegia requiring mechanical ventilation; CSF PCR+ and plasma mcfDNA NGS+	30 mg/kg/day × 16 days → tapered to 2 mg/kg/day × 2 weeks → oral prednisolone taper × 9 weeks (total 33 days IV + 9 weeks PO)		hyperintensities, medullary microhemorrhages. Discharged day 72. At 18 mo: proprioceptive abnormalities but normal language, social and fine motor skills. At 31 mo: walking, climbing, 2-3 word sentences, improved gross motor
COMPLETE RECOVERY (n = 11)							
Graber et al. [1]	1997	Reunion Island	10 mo	Persistent febrile vomiting, irritability, blood and CSF eosinophilia	None (supportive care only)	None	Complete recovery
Graber et al. [1]	1997	Reunion Island	11 mo	Febrile vomiting, irritability, unilateral VI nerve palsy, blood and CSF eosinophilia	None (supportive care only)	None	Complete recovery
Maurer et al. [6]	2001	USA (American Samoa)	10 mo	Fever, refusal to walk, eosinophilia, CSF pleocytosis with eosinophils	Corticosteroids (dose NR)	None	Complete recovery
Flerlage et al. [7]	2017	USA (Tennessee)	12 mo	18 days daily fever, irritability, decreased oral intake, emesis, CSF 36% eosinophils	Dexamethasone (dose NR); 3-week taper; serial LPs	Albendazole × 3 weeks (dose NR)	Complete recovery at 6-month follow-up
Ma et al. [8]	2018	China (Hainan)	15 mo	High fever × 3 wks, irritability × 2 wks, refusal to walk × 1 wk, CSF larvae and eosinophilia	Prednisone 1.5 mg/kg/day (17.5 mg qd) × 4 weeks	Levamisole 2.5 mg/kg/day (14 mg bid) × 4 weeks	Complete recovery at 6 mo; normal growth and development

Ref.	Year	Country	Age	Clinical Presentation	Corticosteroid Regimen	Anthelmintic Regimen	Outcome
Xie et al. [9]	2019	China (Guangdong)	15 mo	Cough × 1 mo, fever × 2 days, mental fatigue, poor diet, hypereosinophilia	Methylprednisolone 20 mg/day × 10 days	Albendazole 100 mg/day × 10 days	Complete recovery without relapse
Xie et al. [9]	2019	China (Guangdong)	19 mo	Cough × 1 mo, fever × 10 days, mental fatigue, poor diet, hypereosinophilia, abnormal MRI	Methylprednisolone 20 mg/day × 10 days	Albendazole 100 mg/day × 10 days	Complete recovery without relapse
Pham Thu et al. [10]	2020	Vietnam (North)	9 mo	Fever × 7 days, seizures, neck stiffness, ↑ICP; blood eosinophilia 16.1%, CSF 26% eosinophils; LIVE WORM visualized in CSF; ELISA+	IV dexamethasone 0.6 mg/kg/day ÷ q8h × 3 days → PO prednisolone 2 mg/kg/day × 5 days; also mannitol 1.5 g/kg/day × 3 days	Albendazole 200 mg/day × 2 weeks	Complete recovery after 12 days; MRI normalized
Cattaneo et al. [11]	2021	Mayotte	14 mo	Strabismus, psychomotor regression, axial hypotonia, hemiparesis, tetraventricular hydrocephalus, CSF eosinophilia 25%	IV methylprednisolone 1 mg/kg/day × 2 weeks → oral prednisolone taper; repeated large-volume LPs	Albendazole 15 mg/kg/day (duration NR)	Complete recovery at 1-month follow-up; normal neurological exam
Chancey et al. [12]	2024	USA (Florida-Orlando)	19 mo	Fever 103°F, 11 days malaise, refusal to walk, Kernig+, no spontaneous LE movement, geophagia history	IV methylprednisolone 2 mg/kg/day × 7 days → oral prednisone taper × 3 weeks	None (rapid improvement with steroids alone)	Complete recovery; at baseline at 3-month follow-up

Ref.	Year	Country	Age	Clinical Presentation	Corticosteroid Regimen	Anthelmintic Regimen	Outcome
Chancey et al. [12]	2024	USA (Florida-St. Petersburg)	8 mo	Vomiting × 6 days, lethargy, fever 103°F, left esotropia, CSF eosinophilia 41%, hypotonia, ataxia	IV methylprednisolone 2→5 mg/kg/day × 7 days → oral prednisone taper × 4 weeks; acetazolamide added	None	Complete recovery; at baseline at 6-week follow-up; required PT

Abbreviations: CSF = cerebrospinal fluid; EVD = external ventricular drain; IVIG = intravenous immunoglobulin; IV = intravenous; LP = lumbar puncture; LE = lower extremity; mcfDNA NGS = microbial cell-free DNA next-generation sequencing; mo = months; NR = not reported; PO = per oral; PT = physical therapy; wks = weeks.

Notes: This table includes only cases with confirmed or highly probable *A. cantonensis* infection in children <24 months of age ($n = 17$ cases from 12 publications). Cases are organized by outcome severity (fatal → sequelae → complete recovery), then chronologically by publication year. Standard anthelmintic dosing for neuroangiostrongyliasis: Albendazole 15 mg/kg/day (max 400 mg bid) × 2–3 weeks. Standard corticosteroid dosing varies: Methylprednisolone 1–2 mg/kg/day (conventional) to 10–30 mg/kg/day (pulse); Prednisolone/Prednisone 1–2 mg/kg/day; Dexamethasone 0.15–0.6 mg/kg/day.

Supplementary Table 2 Pediatric Cases Using High-Dose/Pulse Corticosteroids (10-30 mg/kg/day) for Treatment of *A. cantonensis* Meningoencephalomyelitis

No.	Ref.	Age/Sex	Country	Clinical Severity	High-Dose Corticosteroid Regimen	Anthelmintic	Outcome
1	Abe et al., 2022 [5]	11 mo/M	Hawaii, USA	Coma, complete quadriplegia, respiratory failure requiring mechanical ventilation, longitudinal myelitis C3–C7, cerebral hemorrhage, poor response to IVIG	IV Methylprednisolone 30 mg/kg/day × 16 days → 2 mg/kg/day × 2 weeks → oral prednisolone taper × 9 weeks (Total: 33 days IV + 9 weeks PO)	Albendazole 150 mg/day × 6 weeks	Good recovery with mild proprioceptive abnormality; discharged day 72; at 31 mo: walking, climbing, 2-3 word sentences

No.	Ref.	Age/Sex	Country	Clinical Severity	High-Dose Corticosteroid Regimen	Anthelmintic	Outcome
2	Chancey et al., 2024 [12]	10 yo/M	Tampa, Florida, USA	Coma, central apnea requiring mechanical ventilation, elevated ICP (OP 45 mmHg), obstructive hydrocephalus requiring EVD, abnormal brain MRI with FLAIR hyperintensity and leptomeningeal enhancement	IV Methylprednisolone 10 mg/kg/day × 5 days → tapered over 20 days with prednisolone at day 12	Albendazole 400 mg twice daily × 14 days	Complete recovery; extubated day 12, EVD removed day 18; discharged to inpatient rehabilitation day 22; returned to normal activities without gross neurological deficits
3	Defo et al., 2018 [13]	10 yo/M	French Guiana (Brazil origin)	Eosinophilic meningitis, transverse myelitis T2-T10, paraparesis of lower limbs, dysuria (urinary retention), CSF 92% eosinophils	IV Methylprednisolone 30 mg/kg/day × 5 days → oral prednisolone 2 mg/kg/day tapered over 1 month	Ivermectin 200 µg/kg/day × 10 days	Complete recovery; no sequelae at 3 months; MRI normalized by day 38

Key Observations:

- All 3 cases had severe presentations with neurological compromise (myelitis, coma, respiratory failure)
- High-dose pulse methylprednisolone (10-30 mg/kg/day) was used for 5-16 days, followed by tapering regimens
- All 3 patients achieved good to complete recovery despite initial severe presentations
- The most severely affected infant (Abe 2022) required the longest treatment duration (30 mg/kg/day × 16 days)
- Anthelmintic agents varied: albendazole (2 cases) and ivermectin (1 case)

Abbreviations: mo = months old; yo = years old; M = male; F = female; MV = mechanical ventilation; ICP = intracranial pressure; OP = opening pressure; EVD = external ventricular drain; IV = intravenous; PO = per oral; IVIG = intravenous immunoglobulin; CSF = cerebrospinal fluid; MRI = magnetic resonance imaging; FLAIR = fluid-attenuated inversion recovery.

Notes: This table includes pediatric cases where high-dose/pulse corticosteroid therapy (≥10 mg/kg/day methylprednisolone) was used. Standard-dose corticosteroid regimens (1-2 mg/kg/day) are not included. Pulse steroid therapy is typically reserved for severe cases with significant neurological involvement such as myelitis, encephalitis with altered consciousness, or respiratory compromise.

References

1. Graber D, Jaffar-Bandjee MC, Attali T, Poisson J, Renouil M, Alessandri JL, Combes JC. Angiostrongylosis in infants in Réunion and Mayotte: Apropos of three cases of eosinophilic meningitis including one fatal radiculo-myeloencephalitis with hydrocephalus. *Archives de Pédiatrie*. 1997;4: 424–429. doi:10.1016/S0929-693X(97)86666-3.
2. Cooke-Yarborough CM, Kornberg AJ, Hogg GG, Spratt DM, Forsyth JR. A fatal case of angiostrongyliasis in an 11-month-old infant. *Medical Journal of Australia*. 1999;170:541–543.
3. Morton NJ, Britton P, Palasanthiran P, Bye A, Sugo E, Kesson A, Ardern-Holmes S, Snelling TL. Severe hemorrhagic meningoencephalitis due to *Angiostrongylus cantonensis* among young children in Sydney, Australia. *Clinical Infectious Diseases*. 2013;57:1158–1161. doi:10.1093/cid/cit444.
4. Evans-Gilbert T, Lindo JF, Henry S, Brown P, Christie CDC. Severe eosinophilic meningitis owing to *Angiostrongylus cantonensis* in young Jamaican children: case report and literature review. *Paediatrics and International Child Health*. 2014;34:148–152. doi:10.1179/2046905513Y.0000000106.
5. Abe K, Tanaka Y, Takahashi T, Yamada M, Saito Y, Suzuki H. A rare etiology for ascending paralysis in an infant. *Journal of the Pediatric Infectious Diseases Society*. 2022;11:448–451.
6. Maurer DM, Greene JP, Vincent JM, et al. Fever, refusal to walk, and eosinophilia in a ten-month-old Samoan boy. *The Pediatric Infectious Disease Journal*. 2001;20:230–231, 232–233.
7. Flerlage T, et al. *Angiostrongylus cantonensis* eosinophilic meningitis in an infant, Tennessee, USA. *Emerging Infectious Diseases*. 2017;23:1756–1758.
8. Ma M, Zhang M, Qiu Z. Eosinophilic meningitis caused by *Angiostrongylus cantonensis* in an infant: a case report. *Medicine (Baltimore)*. 2018;97:e10975.
9. Xie M, Zhou Z, Guo S, Li Z, Zhao H, Deng J. Next-generation sequencing specifies *Angiostrongylus* eosinophilic meningoencephalitis in infants: two case reports. *Medicine (Baltimore)*. 2019;98:e16985.
10. Pham Thu Hien H, Dao Huu N, Le Thi Thu T, Nguyen Van L. Case Report: *Angiostrongylus cantonensis* meningoencephalitis in a 9-month-old baby in Vietnam. *American Journal of Tropical Medicine and Hygiene*. 2020;103:723–726. doi:10.4269/ajtmh.20-0166.
11. Cattaneo C, Viterbo A, Piga S, et al. Tetraventricular hydrocephalus following eosinophilic meningitis due to *Angiostrongylus cantonensis* in a 14-month-old boy from Mayotte: a case report. *Open Forum Infectious Diseases*. 2021;8:ofab031.
12. Chance MD, Noel AD, Thompson AB, Marrero N, Bula-Rudas F, Horvat CM, Green J, Armstrong JE, Levent F, Dudas RA, Shaffren S, Samide A, Martinez K, Stockdale K, Chancey RJ. *Angiostrongylus cantonensis* meningoencephalitis in three pediatric patients in Florida, USA. *Journal of the Pediatric Infectious Diseases Society*. 2024;13:639–642.
13. Defo AL, Lachaume N, Cuadro-Alvarez E, Maniassom C, Martin E, Njuieyon F, Henaff F, Mrsic Y, Brunelin A, Epelboin L, Blanchet D, Harrois D, Desbois-Nogard N, Qvarnstrom Y, Demar M, Dard C, Elenga N. *Angiostrongylus cantonensis* infection of central nervous system, Guiana Shield. *Emerging Infectious Diseases*. 2018;24:1153–1155. doi:10.3201/eid2406.180168