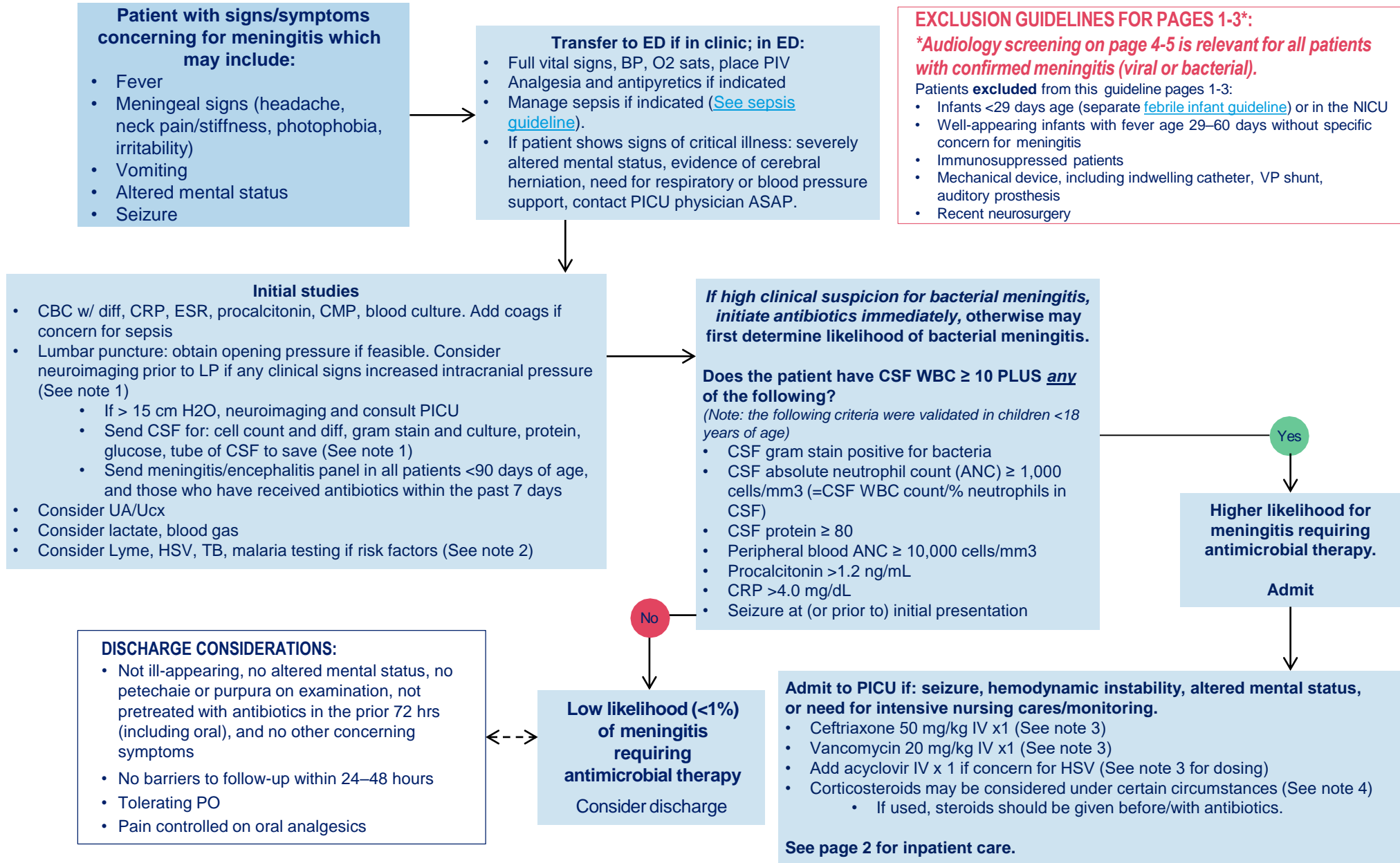
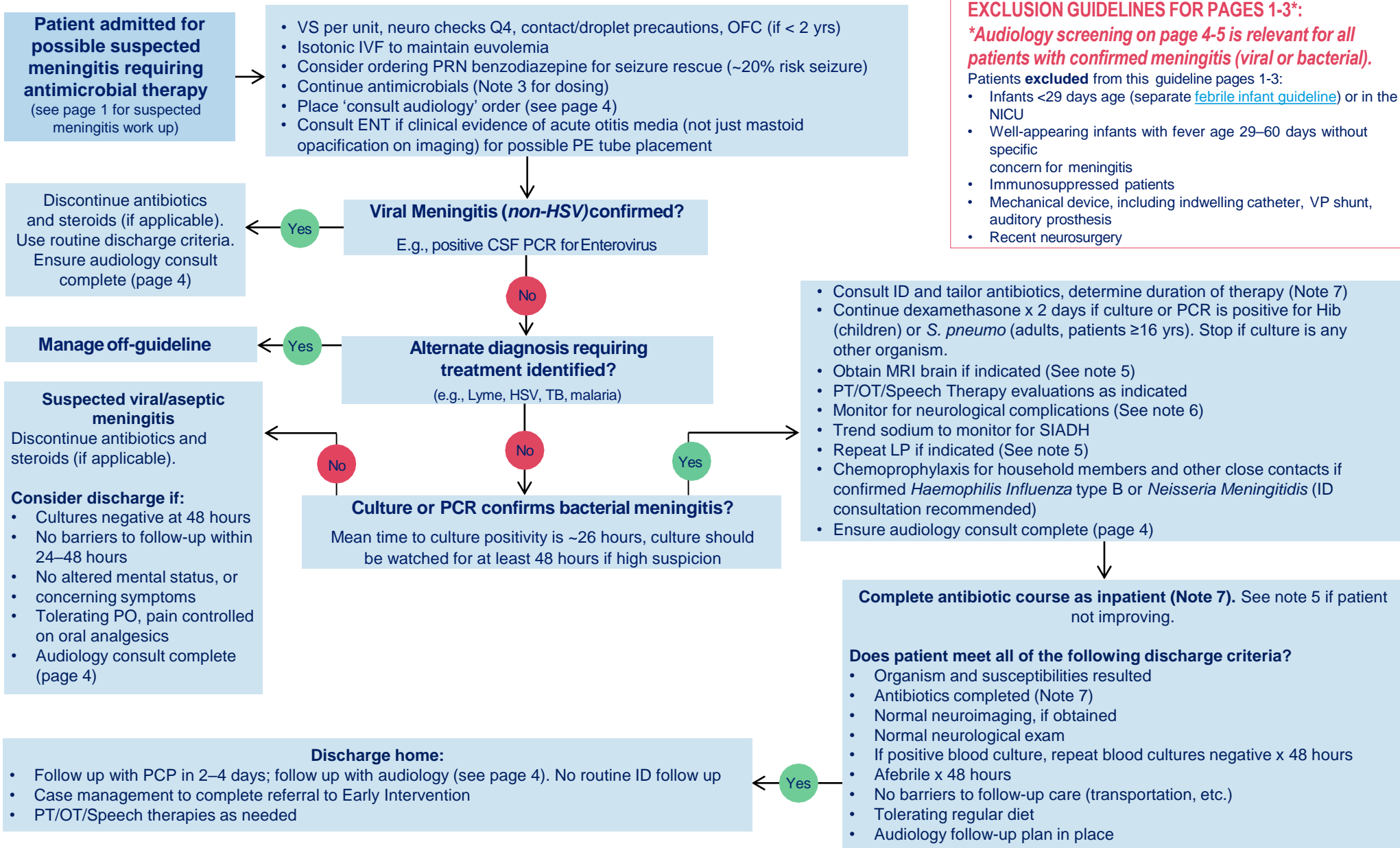


Aim: To reduce unwarranted resource utilization and reduce variation in management of patients with suspected meningitis.



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NOTE 1: LP/CSF testing.

- Recommended to preform cranial imaging prior to LP in patient with: focal neurologic deficits (excluding cranial nerve palsies), new-onset seizures, severely altered mental status (defined as a score on the Glasgow Coma Scale of <10) and severely immunocompromised state.
- Minimum CSF volumes: cell count and diff 0.5 ml min, glucose 0.6 ml, protein 0.6 ml, culture/gram stain min 0.5 ml meningitis/encephalitis panel 0.25 ml. Saved sample should be refrigerated.
- CSF WBC can be adjusted for a suspected bloody tap using a 1:500 ratio (WBC:RBC).
- The meningitis/encephalitis PCR panel should be sent in all patients <90 days of age, and those who have received antibiotics within the past 7 days.

NOTE 2: Risk factors for specific meningitis pathogens:

- Lyme: tick bite several weeks to a few months ago. Lyme meningitis will be covered by ceftriaxone.
- HSV meningitis is associated with primary genital HSV. HSV encephalitis presents with fever, altered mental status, altered level of consciousness, new onset seizure, or focal neurologic deficits.
- TB: subacute presentation (often more than 1 week), altered consciousness, personality changes, cranial nerve palsies common. May not have history of pulmonary disease.
- Parechovirus: cause of meningoencephalitis in infants < 3 months. May have rash; tachycardia out of proportion to fever; concurrent URI symptoms, abdominal distension, diarrhea; skin mottling; apnea (especially in premature infants); seizures common in < 3 months. Expect no CSF pleocytosis.

NOTE 3: Antimicrobial regimens

- Ceftriaxone: 50 mg/kg IV every 12 hours (max 2000 mg/dose). If cephalosporin allergy or type I penicillin allergy (anaphylaxis), use meropenem instead and consult ID.
 - Meropenem: 40 mg/kg IV q8h (max 2000 mg/dose)
- Vancomycin:
 - 20 mg/kg IV q8h if < 18 years old (Max 1250 mg/dose initially)
 - 15 mg/kg IV q8h if 18–24 years old (Max 1250 mg/dose initially)
- Acyclovir IV:
 - < 3 months: 20 mg/kg IV q8h
 - 3 months to 11 years old: 15 mg/kg IV every 8 hours
 - ≥12 years old: 10 mg/kg IV every 8 hours

NOTE 4: Adjunctive Steroids:

- Dexamethasone is beneficial for treatment of infants and children with HiB meningitis to diminish the risk of hearing loss, if administered before or concurrently with the 1st dose of antimicrobial agents
- For infants and children 6 weeks and older with pneumococcal meningitis, adjunctive therapy with dexamethasone is controversial and data are not sufficient to make a routine recommendation for children. Dexamethasone is recommended in adults (patients ≥16 yrs) with suspected or proven pneumococcal meningitis, as there is a mortality benefit.
- Dose: Dexamethasone 0.15 mg/kg/dose IV q6h for 2 days (max 10 mg per dose)
- First dose should be administered 10–20 minutes prior to, or concomitantly with the first dose of antibiotics.

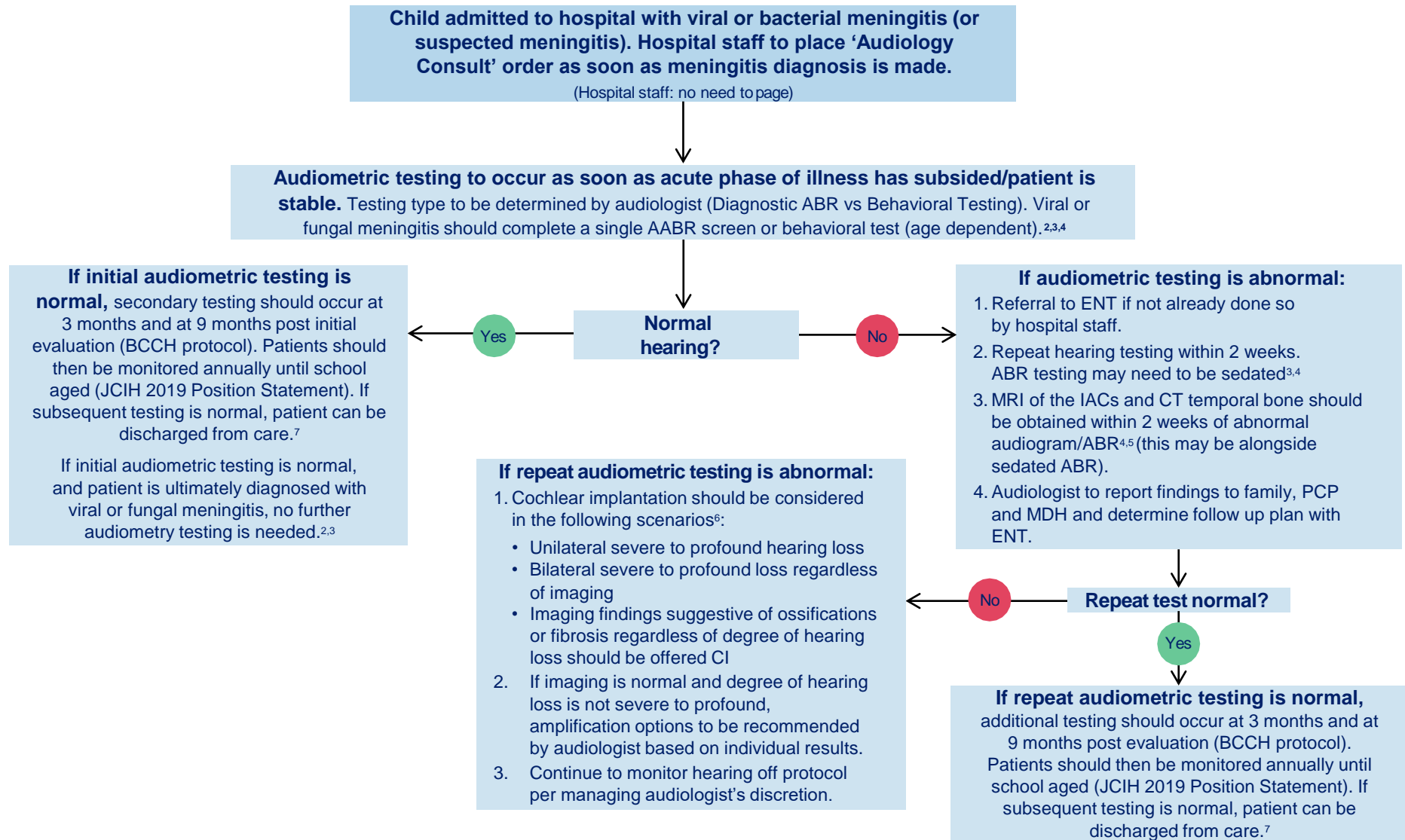
NOTE 5: Brain imaging and repeat LP

- MRI: not routinely needed except if: complicated course, certain pathogens (e.g., Cronobacter, Citrobacter, S. aureus), persistently positive CSF, neonates or older infants with typical neonatal pathogens (GBS, enteric gram negs, listeria) since clinical clues can be limited.
- LP: Repeat at 24–48 hours if gram negative meningitis. Consider repeat at 24–48 hrs in GBS meningitis. For pneumococcal meningitis, repeat LP at 48–72 hours if the isolate is cefotaxime and ceftriaxone nonsusceptible, or if the patient's condition has worsened/not improved, or if the patient has received dexamethasone (which may interfere with ability to mount fever to trend clinical response).

NOTE 6. Potential neurological complications of meningitis include cerebral edema, subdural effusion, seizures, hearing loss, cranial nerve palsy, motor impairment, cerebrovascular complications, hydrocephalus, intellectual disability, mood disorder, attention deficit disorder, hypothalamic dysfunction.

NOTE 7. Duration of therapy: ranges denote typical duration of IV antibiotics. Should be determined for each patient by primary team and ID consultants and other involved specialists.

- S. pneumoniae – 10 to 14 days
- N. meningitidis – 7 days
- H. influenzae – 10 days
- L. monocytogenes – 21 to 28 days
- S. agalactiae – 14 days
- S. aureus – At least two weeks
- Gram-negative bacilli – Three weeks or a minimum of two weeks beyond the first sterile CSF culture, whichever is longer



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AUDIOLOGY/ENT NOTES

NOTE A. Incidence of post-meningitic hearing loss varies from 2-35% (Rodenburg-Vlot et al., 2015). A systematic review of the literature showed an incidence of hearing loss (>25 dB) of 14% and an incidence of 5% for profound hearing loss (>90 dB) (Rodenburg-Vlot et al., 2016). Although hearing loss may improve, fluctuate, or deteriorate, the majority of hearing losses are likely to be stable.

Limited by poor methodology/standardized audiometric care in most studies

NOTE B. Bacteria most likely to cause hearing loss and ossifications are: strep pneumo, neisseria meningitides, haemophilus influenza (BCCH protocol). The evidence does not suggest that viral meningitis or fungal meningitis have high incidence of hearing loss, however, a single screening ABR (age appropriate) or behavioral testing should be performed on these patients until better data exists (they do not need to follow the next steps for f/u in this protocol if normal hearing confirmed).

NOTE C. The Rodenburg-Vlot et al. 2016 systemic review found that hearing loss after bacterial meningitis predominantly seems to be of early onset, therefore all patients should be tested as soon as the acute phase is over (and at the latest at discharge). Ossifications can occur within 2-4 weeks after initial infection (BCCH protocol). Rodenburg-Vlot et al. (2018) suggested long-term audiological follow-up is only needed for the patients who develop a hearing loss during the meningitis, as patients with initial normal hearing after meningitis showed stable normal hearing over time; however at Children's Minnesota, we will monitor more conservatively based on the BCCH protocol and JCIH recommendations listed below.

NOTE D. Ear specific behavioral testing including speech testing should be completed on all meningitis patients, or a diagnostic ABR if age or compliance are limited (De Barros et al. 2014). Auditory brainstem response testing (ABR) can often be completed without sedation under 3 months age. However, sedation is typically required for this test after age 3 months. Consider coordinating with upcoming sedations. Otoacoustic emissions can be used as a supplementary objective test during auditory function tests, but should not be used in isolation (De Barros et al. 2014).

NOTE E. Both CT and MRI have a role

- In five cases with normal CT results, signs of ossification were seen on MRI (20%), whereas ossification was seen on CT in three cases with normal MRI (12%). Thus, the accuracy of both modalities appears inadequate (Caye-Thomasen, 2012)
- If early signs of fibrosis/ossifications, this can progress (Caye-Thomasen, 2012). Cochlear ossifications can occur within 4 weeks after meningitis and can significantly influence surgical complexity in cochlear implantation (Durisin, et al, 2010).
- MRI + CT = 94% sensitivity (Isaacson et al., 2009)
 - CT alone is 50%

NOTE F. Cochlear implantation should be considered in the following scenarios:

- Unilateral cases of severe to profound hearing loss
- Bilateral or unilateral severe to profound loss regardless of imaging
- Imaging findings suggestive of ossifications or fibrosis regardless of degree of hearing loss should be offered CI

NOTE G. British Columbia Children's Hospital (BCCH) recommends all patients diagnosed with bacterial meningitis who are initially follow to have normal hearing, repeat testing at 3 months post-infection, and then again 6 months after the first repeated test (9 months post-infection). The Joint Commission on Infant Hearing (JCIH) recommends all patients who are diagnosed with bacterial meningitis in infancy, have an initial diagnostic test and are monitored annually until school aged.

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REFERENCES

1. Audiology Clinical Practice Guideline: Meningitis. (2013, June). British Columbia Children's Hospital (BCCH), Canada. <https://pdf4pro.com/cdn/audiology-clinical-practice-guideline-2013-june-42e15b.pdf>.
2. Caye-Thomasen, P., Seidelin Dam, M., Haukali Omland, S., & Mantoni, M. (2012) Cochlear ossification in patients with profound hearing loss following bacterial meningitis. *Acta Oto-Laryngologica*, 132(7):720-725. doi:10.3109/00016489.2012.656323.
3. De Barros, A., Roy, T., Amstutz Montadert, I., Marie, J.P., Marcolla, A., Obstoy, M.F., Choussy, O., Dehesdin, D., & Lerosey, Y. (2014) Rapidly progressive bilateral postmeningitic deafness in children: Diagnosis and management. *Eur Ann Otorhinolaryngol Head Neck Dis*. 131(2):107-112. doi: 10.1016/j.anorl.2013.04.006. Epub 2014 Feb 18. PMID: 24559741.
4. Durisin, M., Bartling, S., Arnoldner, C., Ende, M., Prokein, J., Lesinski-Schiedat, A., Lanfermann, H., Lenarz, T., & Stöver, T. (2010) Cochlear Osteoneogenesis After Meningitis in Cochlear Implant Patients, *Otology & Neurotology*. 31(7):1072-1078. doi: 10.1097/MAO.0b013e3181e71310.
5. Isaacson, B., Booth, T., Kutz, J.W., Lee, K.H., & Roland, P.S. (2009) Labyrinthitis ossificans: how accurate is MRI in predicting cochlear obstruction? *Otolaryngol Head Neck Surg*. 140(5):692-6. doi: 10.1016/j.otohns.2008.12.029.
6. Leazer, R., Erickson, N., Paulson, J., Zipkin, R., Stemmler, M., Schroeder, A.R., Bendel-Stenzel, M., & Fine, B.R. (2017) Epidemiology of Cerebrospinal Fluid Cultures and Time to Detection in Term Infants. *Pediatrics*. 139(5):e20163268. doi: 10.1542/peds.2016-3268.
7. Merkus, P., Free, R.H., Mylanus, E.A.M., Stokroos, R., Metselaar, M., van Spronsen, E., Grolman, W., & Frijns, J.H.M. (2010) Dutch Cochlear Implant Group (CI-ON) Consensus Protocol on Postmeningitis Hearing Evaluation and Treatment. *Otology & Neurotology*. 31(8):1281-1286. doi: 10.1097/MAO.0b013e3181f1fc58.
8. Nigrovic, L.E., Kuppermann, N., Macias, C.G., Cannavino, C.R., Moro-Sutherland, D.M., Schremmer, R.D., Schwab, S.H., Agrawal, D., Mansour, K.M., Bennett, J.E., Katsogridakis, Y.L., Mohseni, M.M., Bulloch, B., Steele, D.W., Kaplan, R.L., Herman, M.I., Bandyopadhyay, S., Dayan, P., Truong, U.T., Wang, V.J., Bonsu, B.K., Chapman, J.L., Kanegaye, J.T., & Malley, R. (2007) Pediatric Emergency Medicine Collaborative Research Committee of the American Academy of Pediatrics. Clinical prediction rule for identifying children with cerebrospinal fluid pleocytosis at very low risk of bacterial meningitis. *JAMA*. 297(1):52-60. doi: 10.1001/jama.297.1.52.
9. Nigrovic, L.E., Malley, R., & Kuppermann, N. (2012) Meta-analysis of bacterial meningitis score validation studies. *Arch Dis Child*. 97(9):799-805. doi: 10.1136/archdischild-2012-301798.
10. Rees, C.A., Cruz, A.T., Freedman, S.B., Mahajan, P., Uspal, N.G., Okada, P., Aronson, P.L., Thompson, A.D., Ishimine, P.T., Schmidt, S.M., Kuppermann, N., & Nigrovic, L.E. (2019) HSV Study Group of the Pediatric Emergency Medicine Collaborative Research Committee. Application of the Bacterial Meningitis Score for Infants Aged 0 to 60 Days. *J Pediatric Infect Dis Soc*. 8(6):559-562. doi: 10.1093/jpids/ piy126.
11. Rodenburg-Vlot, M.B.A., Ruytjens, L., Oostenbrink, R., Goedegebure, A., & van der Schroeff, M.P. (2016) Systematic Review: Incidence and Course of Hearing Loss Caused by Bacterial Meningitis: In Search of an Optimal Timed Audiological Follow-up. *Otology & Neurotology*. 37(1):1-8.
12. Rodenburg-Vlot, M.B.A., Ruytjens, L., Oostenbrink, R., & van der Schroeff, M.P. (2018) Repeated Audiometry After Bacterial Meningitis: Consequences for Future Management. *Otol Neurotol*. 39(5):e301-e306. doi: 10.1097/MAO.0000000000001808.
13. Mintegi S, García S, Martín MJ, Durán I, Arana-Arri E, Fernandez CL, Benito J, Hernández-Bou S. Clinical prediction rule for distinguishing bacterial from aseptic meningitis. *Pediatrics*. 2020 Sep 1;146(3).
14. Brouwer MC, McIntyre P, de Gans J, Prasad K, van de Beek D. Corticosteroids for acute bacterial meningitis. *Cochrane database of systematic reviews*. 2010(9).
15. Long SS. Brain imaging and bacterial meningitis. *The Journal of Pediatrics*. 2014 Jul 1;165(1):1-3.
16. Tunkel AR, Hartman BJ, Kaplan SL, Kaufman BA, Roos KL, Scheld WM, Whitley RJ. Practice guidelines for the management of bacterial meningitis. *Clin Infect Dis*. 2004 Nov 1;39(9):1267-84.
17. van de Beek D, Cabellos C, Dzupova O, Esposito S, Klein M, Kloek AT, Leib SL, Mourvillier B, Ostergaard C, Pagliano P, Pfister HW, Read RC, Sipahi OR, Brouwer MC; ESCMID Study Group for Infections of the Brain (ESGIB). ESCMID guideline: diagnosis and treatment of acute bacterial meningitis. *Clin Microbiol Infect*. 2016 May;22 Suppl 3:S37-62.
18. Nelson's Pediatric Antimicrobial Therapy 2021, 27th Ed., edited by John S. Bradley, MD, Elizabeth D. Barnett, MD and Joseph B. Cantey, MD
19. [Bacterial Meningitis Score for Children – MDCalc](#)

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