were absent in both lower extremities. Babinski’s signs were not noted. The Romberg sign was present. Magnetic resonans imaging (MRI) of the spinal cord revealed enhancing T₂ hyperintensity within the spinal cord, from T₁₀–L₁. (Figure 1). Cerebrospinal fluid (CSF) showed no pleocytosis and normal protein and glucose concentrations. The CSF did not show oligoclonal banding and immunoglobulin (Ig)G index was normal. Serum serologic evaluation of Borrelia burgdorferi was positive for IgM but negative for IgG. CSF serologic evaluation was negative (both ELISA and Western blot). The patient was diagnosed as manifesting acute transverse myelitis. He was treated with intravenous methylprednisolone pulse therapy (1000 mg/day for 5 consecutive days), followed by oral prednisolone (1 mg/kg per day). After the start of steroid therapy, the patient showed gradual clinical improvement and was able to walk on the 30th day of illness. We also administered doxycycline 4 mg/kg per day for 14-days for borreliosis.

Conclusion This case serves as a reminder that acute transverse myelitis can be a rare clinical manifestation of Lyme disease. 

Results In the group receiving probiotics, mostly colonizing the stool cultures bacteria were Klebsiella spp, Escherichia coli, Enterococcus spp, Enterobacter spp, Staphylococcus spp. respectively, and in the group not receiving probiotics mostly colonizing the stool cultures bacteria were Klebsiella spp, Enterococcus spp, Staphylococcus spp., Escherichia coli, Enterobacter spp. respectively. When probiotic receiving group compared was with not receiving group, proliferation rate of stool cultures was higher in probiotic group. In the groups receiving and not receiving probiotic, proliferation of the nose cultures were similar. Increase in the proliferation rates of weekly stool cultures in probiotic receiving group was statistically significant but there was no statistically difference in the proliferation rates of nose and other cultures that were taken weekly. There was no statistical difference in both groups in the development of resistant organisms.

Conclusions The use of probiotics in neonatal intensive care unit for premature infants who received treatment with antibiotics, did not prevent the colonization of pathogenic microorganisms.

Background The empirical use of antibiotics in children with suspected meningitis is a common clinical practice worldwide that often leads to drug resistance. It is difficult to clinically differentiate bacterial when compared to viral meningitis until a culture study ofcerebral spinal fluid (CSF) Or CSF viral PCR study is performed. A ‘wait and see’ approach may lead to undesirable outcome. Bacterial Meningitis Score (BMS) is a tool that was developed to help physicians to differentiate between viral versus bacterial meningitis.

Aim To determine the usefulness if any of BMS for discriminating between bacterial or viral meningitis is young children.

Methodology We retrospectively reviewed the charts of all children (from birth till 14 years old) who were admitted with the diagnosis of meningitis to Hamad general hospital in last 2 years period.

Result A total 120 patients (68% boys) with confirmed meningitis were reviewed during the study period. The mean age was (6.3±2.7 year). The majority of patients 112 (93.3%) had viral type meningitis while the remaining had bacterial meningitis (Strep Pneumia, Neisseria meningitis and H. Influenza). The sensitivity of BMS tool revealed a sensitivity of 100% (95% CI: 75.1 to 100.0) and a specificity of 60.9% (95% CI:50.1–69.7).

Conclusion Our study shows that BMS is a simple, easy and highly sensitive tool that can differentiate bacterial from viral meningitis and it is use may limit the use of unnecessary antibiotic s and hospitalizations.

Background and Aims To investigate the effect of probiotics on colonization of resistant organisms in preterm infants receiving antibiotics.

Methods This study comprised of preterm infants who were born < 36 weeks and received antibiotic treatment or prophylaxis. Preterm infants were divided into two groups according to receiving probiotic (Lactobacillus reuteri). Stool culture and nasal swab cultures were taken to determine colonization.

Results In the group receiving probiotics, mostly colonizing the stool cultures bacteria were Klebsiella spp, Escherichia coli, Enterococcus spp, Enterobacter spp, Staphylococcus spp. respectively, and in the group not receiving probiotics mostly colonizing the stool cultures bacteria were Klebsiella spp, Enterococcus spp, Staphylococcus spp., Escherichia coli, Enterobacter spp. respectively. When probiotic receiving group compared was with not receiving group, proliferation rate of stool cultures was higher in probiotic group. In the groups receiving and not receiving probiotic, proliferation of the nose cultures were similar. Increase in the proliferation rates of weekly stool cultures in probiotic receiving group was statistically significant but there was no statistically difference in the proliferation rates of nose and other cultures that were taken weekly. There was no statistical difference in both groups in the development of resistant organisms.

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