SUCCESSFUL NEUROLOGY TRAINEE RESEARCH NETWORK COLLABORATIVE AUDIT

PO001

SUCCESSFUL NEUROLOGY TRAINEE RESEARCH NETWORK COLLABORATIVE AUDIT
Eiman Abdelgadir, Sharmini Patel, Michael O’Gara, Oliver Leach, Omar Al Masri, Shalaja Bhattacharjee, Alexander Shah, Jamie Talbot, Hasaan Zafar, Camille Carroll, Plymouth Hospital NHS Trust, UK; Royal Devon and Exeter Hospital, UK; Royal Cornwall Hospital, UK

Introduction Successful Trainee Clinical Research Networks have been established since 2007. Our network in the peninsula, the SOUTHWEST Neurology Audit and Research Group (SONAR) is the first such Neurology trainee network in the UK. To enable development of cohesive collaborative working of the network we designed an audit which would be deliverable across three neurology centres within the peninsula.

Method We audited management of suspected acute meningitis and meningococcal sepsis against national guidelines within a 4 week period in December. A standardised anonymised data collation tool was used across the three centres and results were analysed at one centre.

Results All 9 registrars on the rotation contributed to audit methodology design and data analysis; seven contributed cases (from all three centres). Ten cases were included in the audit, 6 (Exeter), 3 (Plymouth), and 1 (Truro). Our audit highlighted deficiencies in timely senior review, delivery of antibiotics and steroids, inappropriate administration of acyclovir and delay in lumbar puncture.

Conclusion This was SONAR’s first collaborative project and demonstrated that as a group of trainees we can successfully conduct a project across multiple hospital sites. We plan to extend the scope and ambition of our future undertakings.

BACTERIAL MENINGITIS WITH MYELOPATHY AND CRANIAL NEUROPATHY

PO002

Darsh Ahbakin, Samiksha bundles, Krishna Chinthapalli, Gemma Cummins, Robin Howard, Edward J Wild, Department of Neurology, National Hospital for Neurology and Neurosurgery, Queen Square, London, UK

Acute meningococcal meningitis in adults can be complicated by cranial neuropathies and more rarely by myelopathy. A 55-year-old woman presented with acute bacterial meningitis requiring intubation and was treated with intravenous antibiotics and dexamethasone. Cerebrospinal fluid 16S PCR was positive for Neisseria meningitidis and latex agglutination confirmed the W135 serotype. Ten days after presentation she developed mild upper limb and severe lower limb weakness with hypertreflexia, despite resolution of meningism and improvement of inflammatory markers. On the next day she developed complete bilateral hearing loss and bilateral facial palsy. Magnetic resonance imaging with contrast showed bilateral enhancement of VIIth and VIIIth cranial nerves, with focal signal change within the thoracic spinal cord. Audiometry confirmed complete sensorineural deafness. She was treated with five days of intravenous methylprednisolone and continued a further nine days of intravenous antibiotics. Four weeks after onset, she remained completely deaf with mild improvement of limb and facial weakness. We report a case of acute meningococcal meningitis complicated by presumed extensive vasculitis leading to myelopathy and delayed onset of multiple cranial neuropathies. There was an unusual biphasic presentation and the vasculitis was apparently ameliorated by glucocorticoids, suggesting conventional steroid recommendations may occasionally be inadequate.

ILIOPSOAS HAEMATOMA PRESENTING AS PAINLESS WEAKNESS

PO003

Omar Almasri, Chench Reddy Kannaka, Stuart Weatherby, Derriford Hospital, UK

Spontaneous iliopsoas haematoma is an uncommon complication of anticoagulation. Such bleeding can be life threatening and therefore early recognition is important. In the published literature the universal presenting symptom is pain. We report on the unusual case of an elderly patient who was anticoagulated with Warfarin for Atrial fibrillation, and who presented with painless weakness of the right lower limb. Anticoagulation was reversed and her condition was managed conservatively. Her case is presented in the context of the published literature.

PAINLESS AND PROGRESSIVE THINK COMPRESSIVE

PO004

Sukaina Asad, Geoffrey Stone, Dan Morris, Caroline Hayhurst, Ann Johnston, University Hospital of Wales, Cardiff, UK

A 47-year-old woman was referred by her ophthalmologist with a working diagnosis of optic neuritis. She presented with a 6 week history of progressive, painless loss of vision in her left eye associated with complete loss of colour vision. She was otherwise well. On examination she was normotensive. Visual acuity on the right with pinhole was 6/9 + 3 and 6/0 on the left. Field testing on the right eye was full but in her left eye she was just able to appreciate grey shadows and object outlines. There was a left RAPD and bilateral disc pallor. There was no papilloedema or ophthalmoplegia. There were no other neurological signs or endocrinological abnormalities. Inflammatory markers and an autoimmune screen were normal. Neuro-imaging demonstrated a large 50 × 53 mms mass in sphenoid sinus eroding the skull base into the pituitary floor, the optic canals and displacing both cavernous sinuses. Urgent drainage of this sphenoid mucocele was performed in an attempt to preserve vision in both eyes. In conclusion painless and progressive visual loss is out of keeping for an