## CHARGE SYNDROME CHECKLIST: HEALTH SUPERVISION ACROSS THE LIFESPAN (FROM HEAD TO TOE)

\*Shaded boxes indicate key assessment points **ADOLESCENCE ADULTHOOD** INFANCY **CHILDHOOD** (0:2 years) (3:11 years) (12:17 years) (18+ years) Clinical diagnosis (Blake et al. or Verloes or Hale et al. criteria) Genetic testing - Genetics consult (CHD7 analysis, array CGH) Genetic counselling CNS malformations/hypoplasia olfactory bulb/temporal bone (semicircular canal) malformations – requires MRI/CT  $\,$ UROLOGY Seizures – more common at older ages – consider EEG Cranial nerve problems - monitor for absent sense of smell, facial nerve 핅 palsy, sensorineural hearing loss, vertigo, swallowing problems Coloboma, risk of retinal detachment E Ophthalmology consult (dilated eye exam in infancy, vision assessments) EARS, NOSE AND THROAT Corneal exposure – lubricating eye drops Photophobia - tinted glasses, sunhat Choanal atresia/cleft palate/tracheoesophageal fistula E ENT/Plastics consult Audiometry and tympanometry, monitor for recurrent ear infections Adaptive services for individuals with deafness/blindness Cochlear implant assessment if applicable Obstructive sleep apnea - monitor for tonsil/adenoid hypertrophy EYES, Excessive secretions - consider Botox, medication Dental issues - consider cleaning under anaesthetic Cardiac malformations common – major/minor defects, vascular ring or CARDIOLOGY RESPIROLOGY arrhythmias possible (echocardiogram, chest x-ray, ECG) E Cardiology consult Sinusitis, pneumonia, asthma E monitor Anesthesia risk (difficult intubations/postop airway obstruction/aspiration) extensive preoperative assessment, combine surgical procedures Gastroesophageal reflux - Gastroenterology consult - consider motility agents with proton pump inhibitor GASTROENTEROLOGY GENITOURINARY Poor suck/chew/swallow E feeding team assessment/intervention Aspiration risk, tracheoesophageal fistula - swallowing studies May need supplemental feeds - frequently requires gastrostomy tube or Gastro-jejunostomy tube Constipation – consider Senna glycoside with polyethylene glycol Renal anomalies – abdominal u/s +/E VCUG, blood pressure monitoring Hypogonadotropic hypogonadism - LH, FSH by 3 months Genital hypoplasia (if undescended testes E consider orchidoplexy) ENDOCRINOLOGY Delayed puberty - Endocrinology consult E gonadotropin levels, HRT Osteoporosis - DEXA scan Poor growth - Endocrinology consult - GH stimulation test, GH therapy Obesity E monitor Fertility and contraception E discuss Note presence of thymus at open heart surgery Routine immunizations/antibody titres to immunizations in adolescence Recurrent infections - Immunology consult Scoliosis/kyphosis monitor MSK Mobility (affected by ataxia, hypotonia) E evaluate Assess gross and fine motor skills – Occupational Therapy, Physiotherapy Communication, language, writing abilities - Speech Language Therapy Consider deatblind consultant Prepare for transitions to school, situations, places, systems

Family stress – offer supports and resources

Sleep disturbances – consider melatonin

obsessions, compulsions, anger Toileting skills E support

Address sexuality

DEVELOPMENTA

Psychoeducational assessment, Individualized Education Plan

Lite skills/adaptive behaviour/social skills/social play

conditions, seeing healthcare provider independently

Behavior management – self regulation, impulse control, anxiety,

Medical self-management - work on managing medications, understanding

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#### **Abbreviations Used in Checklist**

CGH E comparative genomic FSH E follicle stimulating hormone

hybridization GH E growth hormone

CNS E central nervous system HRT E hormone replacement therapy

CT E computed tomography LH E luteinizing hormone

DEXA scan E dual energy XEray MRI E magnetic resonance imaging

absorptiometry MSK E musculoskeletal

EEG E electroencephalogram U/S E ultrasound

ENT E ear, nose and throat VCUG E voiding cystourethrogram

#### Resources

 The CHARGE Syndrome Foundation (http://chargesyndrome.org/aboutE charge.asp)

- The CHARGE Informational Pack for Practitioners (SENSE UK) (https://www.sense.org.uk/content/chargeEinformationEpackEpractitioners)
- Book E CHARGE Syndrome (Genetics and Communication Disorders), 1<sup>st</sup> ed.
  Hartshorne TS, Hefner M, Davenport S, Thelin J. 2011
- OMIM Entry #214800 CHARGE Syndrome (http://www.omim.org/entry/214800)
- CHARGE Syndrome International Conference
- CHARGE Syndrome Listserv
- CHARGE Syndrome Facebook Group

Perkins School for the Blind eElearning

(<a href="http://www.perkinselearning.org/videos">http://www.perkinselearning.org/videos</a>)

- Deafblind International (<a href="http://www.deafblindinternational.org/index.htm">http://www.deafblindinternational.org/index.htm</a>)
- Open hands, open access: deafEblind intervener learning modules (http://moodle.nationaldb.org)

### **Key General References**

- 1. Blake K, Prasad C. 2006. CHARGE syndrome. Orphanet J Rare Dis 1: 34
- 2. Brown D. 2005. CHARGE syndrome "behaviors": challenges or adaptations? Am J Med Genet Part A 133A: 268E272
- 3. Hsu P, Ma A, Wilson M, Williams G, Curotta J, Munns CF, Mehr S. 2014. CHARGE syndrome: a review. J Pediatr Child Health 50: 504E511