

<p>Neurologic, Psychiatric and/or Rheumatologic Plagiocephaly and/or craniosynostosis (<i>n</i>=5) Sensory sensitivity (<i>n</i>=2) Tic (<i>n</i>=1) Ehlers-Danlos Syndrome (<i>n</i>=1) Postural orthostatic tachycardia syndrome (<i>n</i>=1) Dyslexia (<i>n</i>=1) Attention deficit hyperactivity disorder (<i>n</i>=1) Foot pronation(<i>n</i>=1)</p>	<p>Otolaryngologic Hearing loss (<i>n</i>=3) Tympanostomy (<i>n</i>=2)</p> <p>Prematurity and Associated Complications Gestational age 26 weeks with chronic lung disease (<i>n</i>=1) Gestational age 33 weeks due to fetal hydrops (<i>n</i>=1) Gestational age 25 weeks with intraventricular hemorrhage (<i>n</i>=1)</p>
<p>Ophthalmologic Pupil asymmetry (<i>n</i>=2) Amblyopia (<i>n</i>=1) Ptosis (<i>n</i>=1) Intermittent diplopia (<i>n</i>=1) Coloboma with microphthalmia (<i>n</i>=1)</p>	<p>Genetic KMT2C gene mutation associated with congenital heart disease, developmental delay and autism spectrum disorder (Kleestra syndrome) SACS gene mutation associated with periodic fevers, tic, tethered cord and leg dystonia (<i>n</i>=1) Multiple congenital anomalies (short long bones, hypospadias, bifid uvula, sacral dimple, joint hypermobility, tics and learning difficulties without identified mutation (<i>n</i>=1)</p>
<p>Cardiac Ventricular septal defect requiring surgical repair (<i>n</i>=2) Unspecified congenital heart defect (<i>n</i>=1) Murmur with normal echocardiogram (<i>n</i>=1) Heart palpitations (<i>n</i>=1)</p>	
<p>Gastrointestinal Gastroesophageal reflux (<i>n</i>=23) Inguinal hernia (<i>n</i>=1) Milk protein allergy (<i>n</i>=1) Viral hepatitis (<i>n</i>=1)</p>	

Supplemental Table S1: Medical comorbidities of children with BPT as reported by their parents/caregivers. Total number of comorbidities is greater than the number of children with any comorbidity as some reported more than one condition.

Testing and Evaluations Performed	Abnormal Findings
Neuroimaging MRI brain (<i>n</i> =40) MRI spine (<i>n</i> =12) MRI orbit (<i>n</i> =1) CT head (<i>n</i> =6) CT neck (<i>n</i> =1) CT unspecified (<i>n</i> =2) X-ray neck (<i>n</i> =1) X-ray spine (<i>n</i> =1) Head ultrasound (<i>n</i> =4) Neck ultrasound (<i>n</i> =1)	MRI brain: Chiari malformation (<i>n</i> =1); dysgenesis corpus callosum (<i>n</i> =1); “incidental small cyst” (<i>n</i> =1); white matter injury of unknown etiology (<i>n</i> =1); enlarged vestibular aqueduct (<i>n</i> =1) MRI spine: Tethered cord in patient with SACS gene mutation (variant of unknown significance) (<i>n</i> =1) Neck ultrasound: Lateral neck cyst (<i>n</i> =1)
EEG (<i>n</i> =39)	Posterior slowing at 12 months of age, felt to be “normal variant” (<i>n</i> =1)
Lumbar puncture (<i>n</i> =4)	None reported
Genetic testing (<i>n</i> =8) Specific testing for CACNA1A (<i>n</i> =2)	KMT2C mutation (<i>n</i> =1): encodes histone methyltransferase; associated with Kleefstra syndrome 2 characterized by intellectual disability, hypotonia, speech delay and facial dysmorphism SACS gene variant of unknown significance (<i>n</i> =1): associated with Autosomal recessive spastic ataxia of Carlevoix-Saguenay (ARSACS), characterized by dystonia, ataxia, spasticity, amyotrophy, nystagmus and dysarthria
Ophthalmologic evaluation (<i>n</i> =20) By an ophthalmologist (<i>n</i> =14) By an optometrist (<i>n</i> =2) By a neuro-ophthalmologist (<i>n</i> =1) Vision screen, NOS (<i>n</i> =3) “Eye exam”, NOS (<i>n</i> =2)	Amblyopia (<i>n</i> =1); pupil asymmetry (<i>n</i> =2); ptosis (<i>n</i> =1); coloboma with microphthalmia (<i>n</i> =1); glasses prescription needed in left eye (<i>n</i> =1)
Otolaryngologic evaluation (<i>n</i> =14) Hearing test (<i>n</i> =5) Vestibular testing (<i>n</i> =2) “ENT” (<i>n</i> =7)	Abnormal hearing, felt to be unrelated to BPT per parental report (<i>n</i> =3; 1 with KMT2C mutation); right-sided vestibular deficit, felt to be unrelated to BPT per parental report (<i>n</i> =1)
Lab testing (<i>n</i> =8)	Elevated urinary dopamine (normal MIBG scan and renal ultrasound) (<i>n</i> =1)
Gastroenterological evaluation (<i>n</i> =4) Abdominal ultrasound (<i>n</i> =2) Abdominal MRI (<i>n</i> =1) Upper GI series (<i>n</i> =1)	None reported
Orthopedic evaluation (<i>n</i> =4)	None reported
Allergy testing (<i>n</i> =2)	None reported
Cognitive testing (<i>n</i> =1)	None reported
Hair mineral testing (<i>n</i> =1)	None reported
Neurosurgical evaluation (<i>n</i> =1)	None reported

Supplemental Table S2: Diagnostic evaluations performed and associated abnormal findings as reported

by their parents/caregivers. Total number of evaluations is greater than the number of patients as many

patients had more than one test performed. MRI: magnetic resonance imaging; CT: computed tomography; EEG: electroencephalogram; NOS: not otherwise specified; ENT: Ear, nose and throat; GI: gastrointestinal; MIBG: metaiodobenzylguanidine