

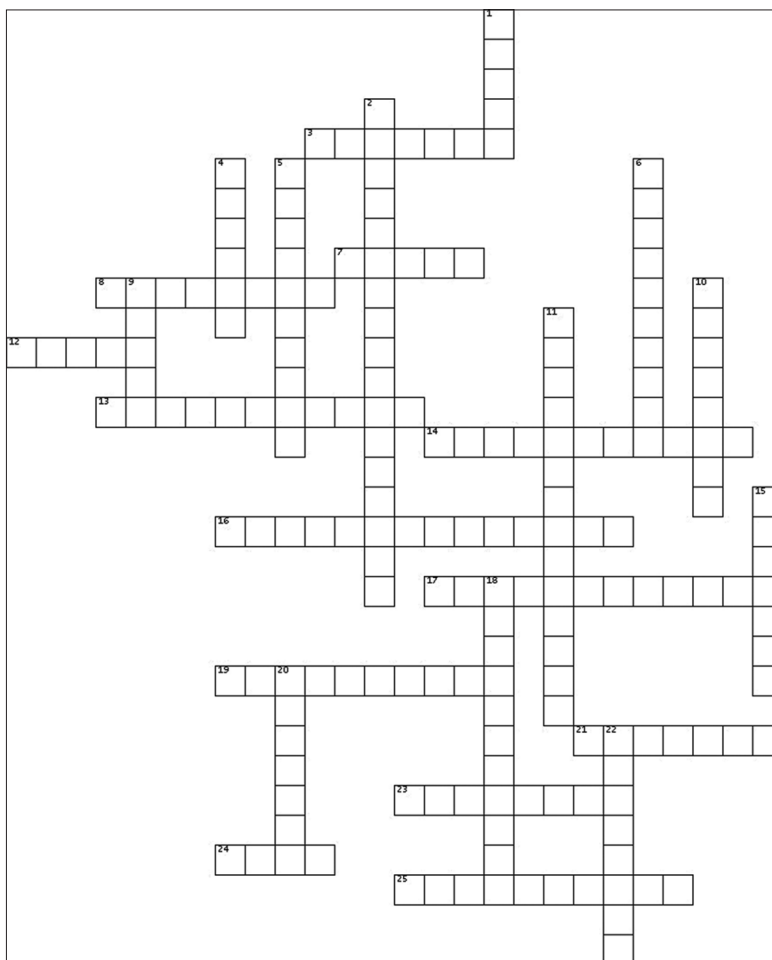
Crossword

A Syndrome Scrabble

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SYNDROME CROSSWORD



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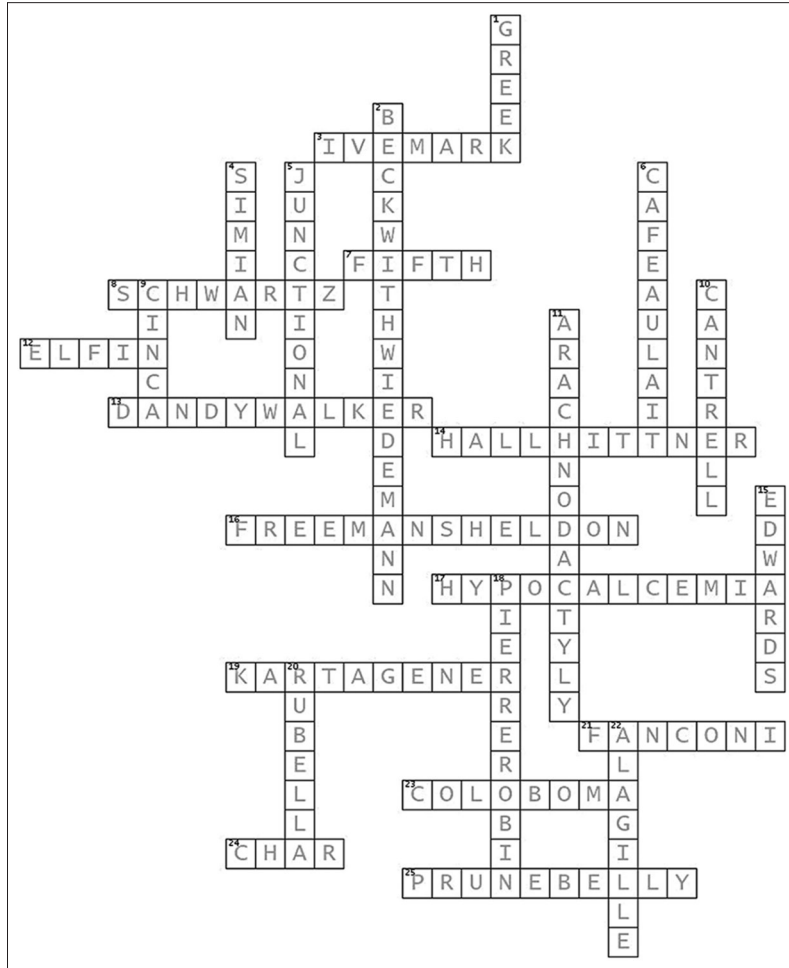
DOWN

- 1 The word "Syndrome" derives its origin from this language.
- 2 Macrosomia, hemihypertrophy, and macroglossia form a triad of this syndrome.
- 4 Single Palmar Crease seen in Trisomy 21.
- 5 Congenital His Bundle Tachycardia, otherwise known as _____ Ectopic Tachycardia, which is a rare arrhythmia observed in neonates.
- 6 >6 of these spots constitute a dermatological sign of Neurofibromatosis type 1.
- 9 Acronym for this syndrome which also happens to be the other name for neonatal onset multisystem inflammatory disease.
- 10 A pentalogy formed by the constellation of five congenital midline birth anomalies.
- 11 A term used to describe long, slender, and spider-like abnormality of the fingers.
- 15 Trisomy 18 Syndrome.
- 18 Sequence causing micrognathia, glossoptosis, and cleft palate, resulting in a difficult airway.
- 20 Classic Triad of cataract, patent ductus arteriosus, and sensorineural deafness, seen in.
- 22 An autosomal dominant disease linked to arteriohepatic dysplasia.

ACROSS

- 3 A type of heterotaxy syndrome with findings of right isomerism, like asplenia.
- 7 Benign idiopathic neonatal seizures are called as _____ day fits.
- 8 Diagnostic criteria for congenital long QT syndrome.
- 12 Named facies in William's Syndrome.
- 13 A congenital malformation featuring hydrocephalus, posterior fossa cyst, and the absence of the cerebellar vermis.
- 14 "CHARGE" association is also known as _____ syndrome.
- 16 Whistling Face syndrome having features of microstomia.
- 17 Chromosome 22q11.2 deletion may present with this electrolyte abnormality due to primary hypoparathyroidism.
- 19 Immotile cilia syndrome.
- 21 Anemia associated with abnormalities of the gastrointestinal system such as esophageal atresia and tracheoesophageal fistula.
- 23 Congenital absence or defect of ocular tissue, frequently seen across neonatal syndromes.
- 24 Syndrome associated with a mutation in neural crest transcription factor TFAP2B, characterized by typical facial and hand anomalies with patent ductus arteriosus.
- 25 Syndrome deriving its name from the wrinkled appearance of abdominal skin.

SYNDROME CROSSWORD ANSWERS



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