SLC6A1-RELATED DISORDERS: CLINICAL ASPECTS

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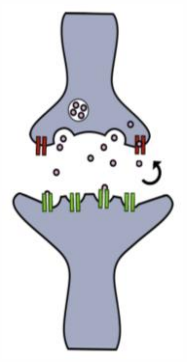

November 29, 2018

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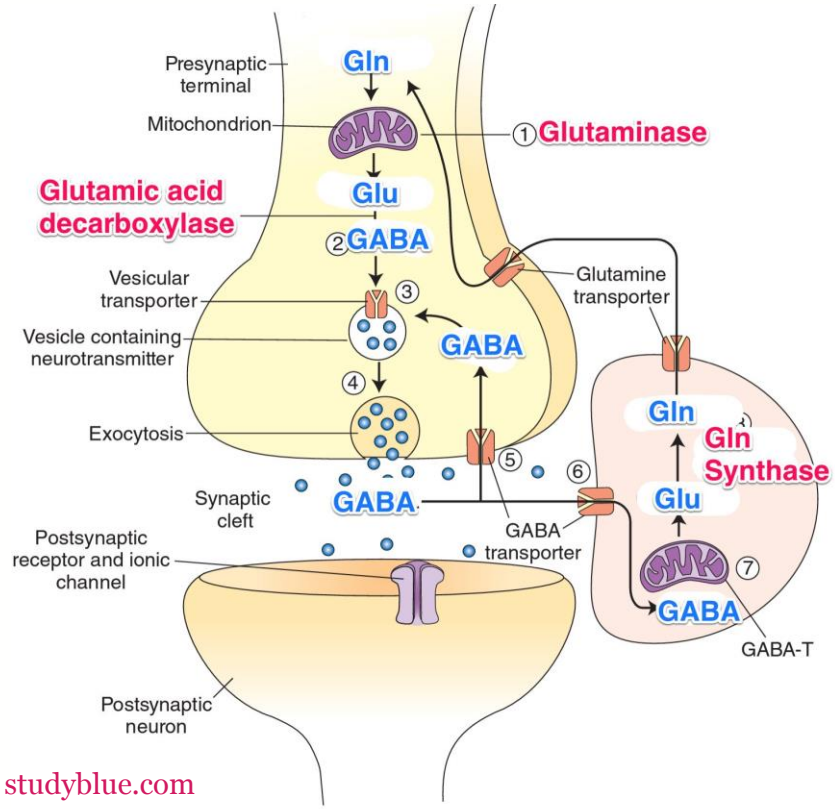
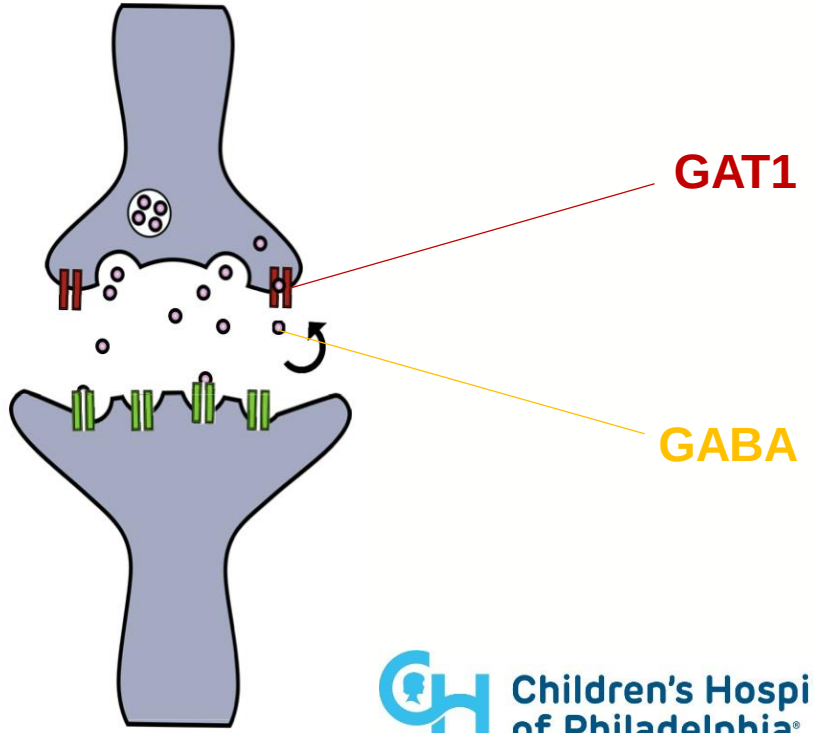
SLC6A1 GENE

- Encodes instructions for GABA transporter 1 (GAT1)
- Removes GABA from synaptic cleft
  - Major inhibitory neurotransmitter in the brain



Credit: studyblue.com



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SLC6A1-RELATED DISORDERS

- First implicated in neurological disease by Carvill et al. 2015
  - 6 individuals with MAE with pathogenic *SLC6A1* variants
  - 4% of individuals with MAE explained by *SLC6A1*

Mutations in the GABA Transporter *SLC6A1* Cause Epilepsy with Myoclonic-Atonic Seizures

Gemma L. Carvill,[1] Jacinta M. McMahon,[2] Amy Schneider,[2] Matthew Zemel,[1] Candace T. Myers,[1] Julia Saykally,[1] John Nguyen,[1] Angela Robbiano,[3] Federico Zara,[3] Nicola Specchio,[4] Oriano Mecarelli,[5] Robert L. Smith,[6] Richard J. Leventer,[7,8,9] Rikke S. Møller,[10,11] Marina Nikanorova,[10] Petia Dimova,[12] Albena Jordanova,[13,14,15] Steven Petrou,[16] EuroEPINOMICS Rare Epilepsy Syndrome Myoclonic-Astatic Epilepsy & Dravet working group, Ingo Helbig,[17,18] Pasquale Striano,[19] Sarah Weckhuysen,[13,14,20] Samuel F. Berkovic,[2] Ingrid E. Scheffer,[2,7,16,21,*] and Heather C. Mefford[1,21,*]

The American Journal of Human Genetics *96*, 808–815, May 7, 2015

SLC6A1-RELATED DISORDERS

- Follow up study by Johannesen et al. 2018

FULL-LENGTH ORIGINAL RESEARCH

Epilepsia®

Defining the phenotypic spectrum of *SLC6A1* mutations

Katrine M. Johannesen[1,2] | Elena Gardella[1,2] | Tarja Linnankivi[3] | Carolina Courage[4,5] | Anne de Saint Martin[6,7] | Anna-Elina Lehesjoki[4,5] | Cyril Mignot[8] | Alexandra Afenjar[9] | Gaetan Lesca[10,11,12] | Marie-Thérèse Abi-Warde[6,7] | Jamel Chelly[13,14] | Amélie Piton[13,14] | J. Lawrence Merritt II[15] | Lance H. Rodan[16,17] | Wen-Hann Tan[16,17] | Lynne M. Bird[18] | Mark Nespeca[19] | Joseph G. Gleeson[20] | Yongjin Yoo[21] | Murim Choi[21] | Jong-Hee Chae[22] | Desiree Czapansky-Beilman[23] | Sara Chadwick Reichert[24] | Manuela Pendziwiat[25] | Judith S. Verhoeven[26] | Helenius J. Schelhaas[26] | Orrin Devinsky[27] | Jakob Christensen[28] | Nicola Specchio[29] | Marina Trivisano[29] | Yvonne G. Weber[30] | Caroline Nava[31,32] | Boris Keren[31,32] | Diane Doummar[33] | Elise Schaefer[34] | Sarah Hopkins[35] | Holly Dubbs[36] | Jessica E. Shaw[36] | Laura Pisani[36] | Candace T. Myers[15] | Sha Tang[37] | Shan Tang[38] | Deb K. Pal[38] | John J. Millichap[39,40] | Gemma L. Carvill[40] | Kathrine L. Helbig[37] | Oriano Mecarelli[41] | Pasquale Striano[42] | Ingo Helbig[25,35] | Guido Rubboli[1,43] | Heather C. Mefford[15] | Rikke S. Møller[1,2]

Epilepsia. 2018;59:389–402.

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SLC6A1-RELATED DISORDERS

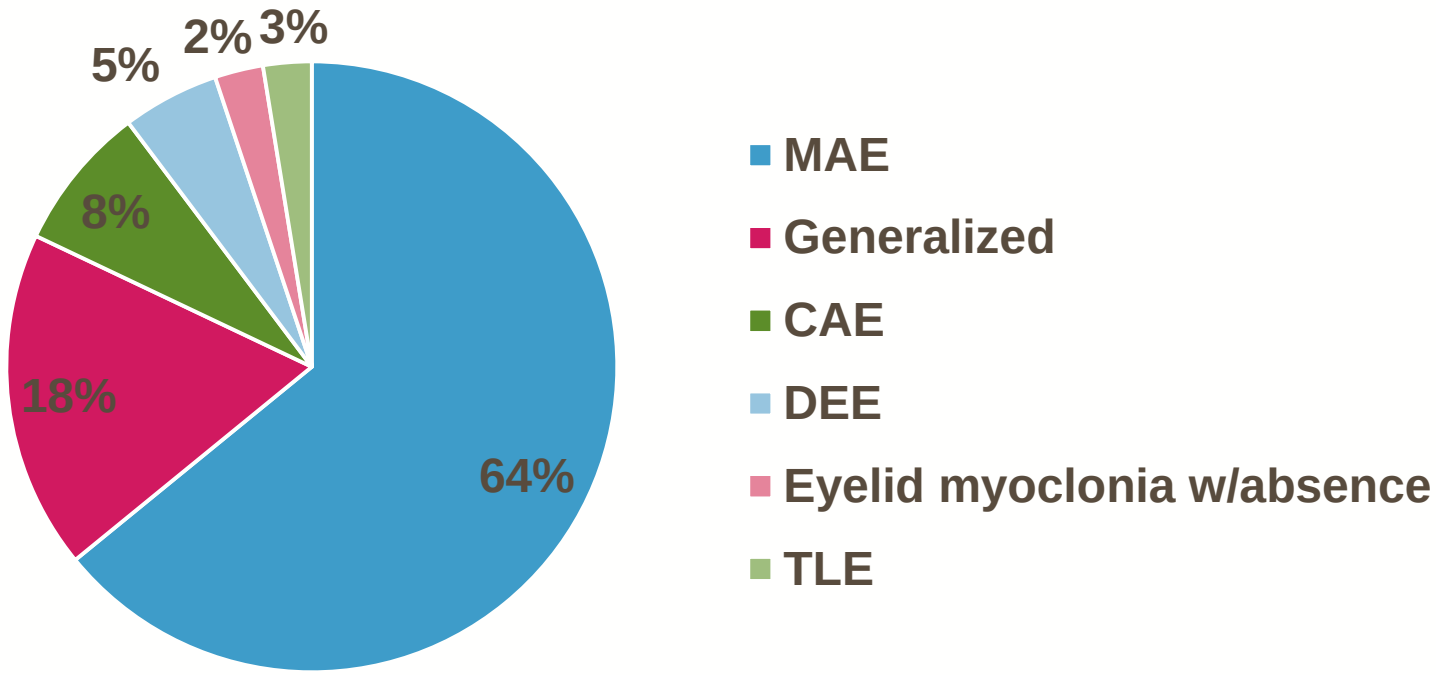
- As of November 2018:
  - 48 individuals published in the literature
  - 41 unique *SLC6A1* variants reported in HGMD
  - 39 (likely) pathogenic *SLC6A1* variants in ClinVar

- Phenotypic spectrum has expanded beyond Epilepsy with Myoclonic-Atonic Seizures (MAE/Doose syndrome)

- What do *SLC6A1*-related disorders look like now?

PHENOTYPIC FEATURES - EPILEPSY

- Epilepsy is present in 81% of individuals
  - Median age of onset 24 months (range 5m – 7y)
  - 65% of individuals become seizure free



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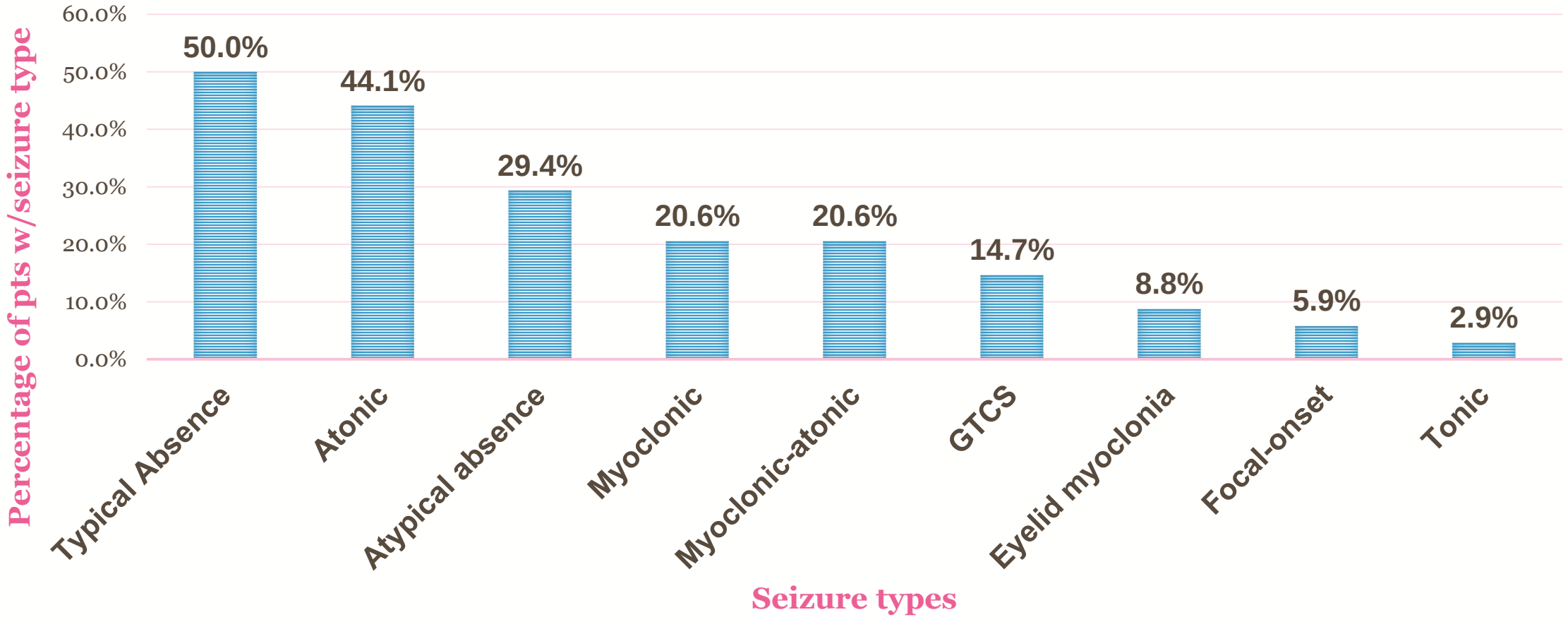
PHENOTYPIC FEATURES - EPILEPSY

- Generalized seizure types predominate



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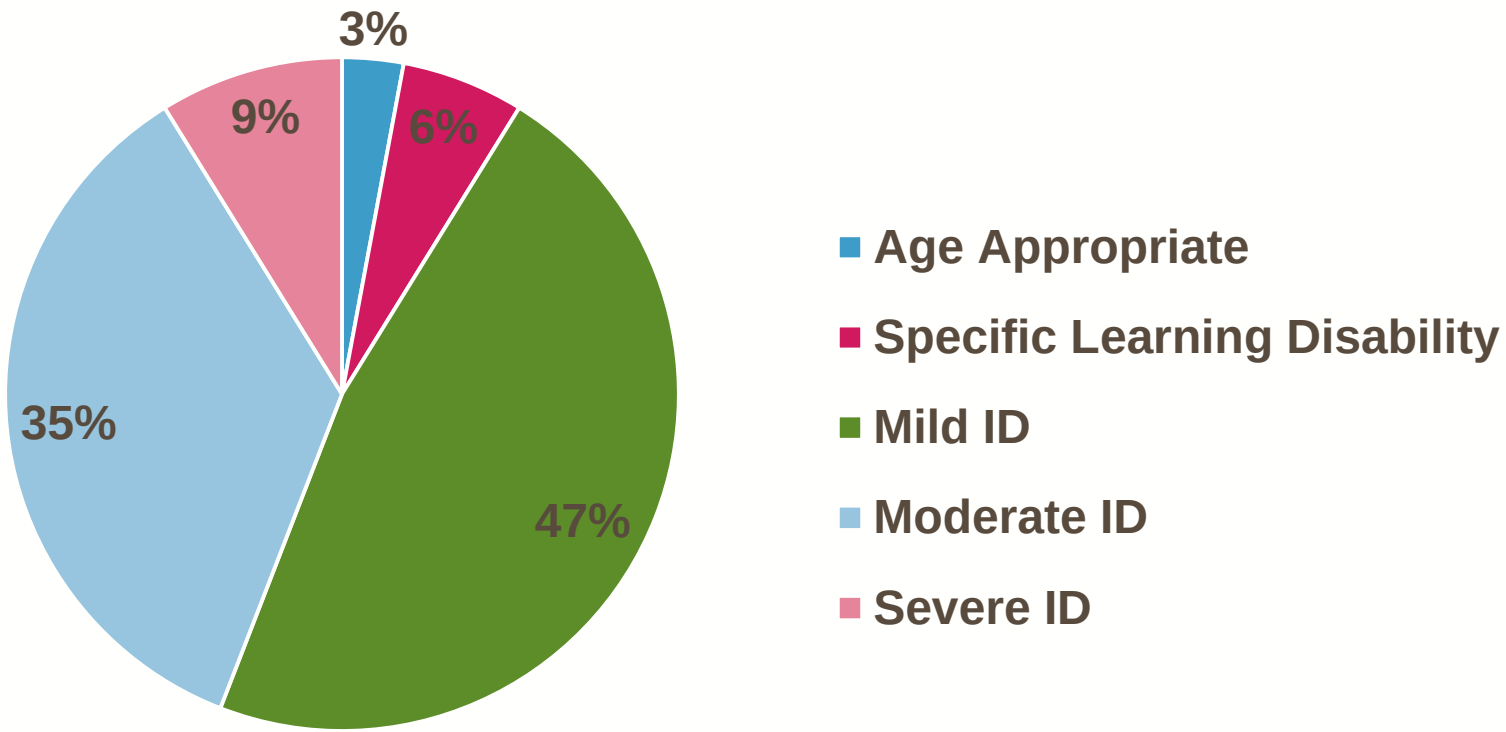
PHENOTYPIC FEATURES - DEVELOPMENT

- Developmental delays in 91% of individuals
- No correlation between seizure control and developmental outcome



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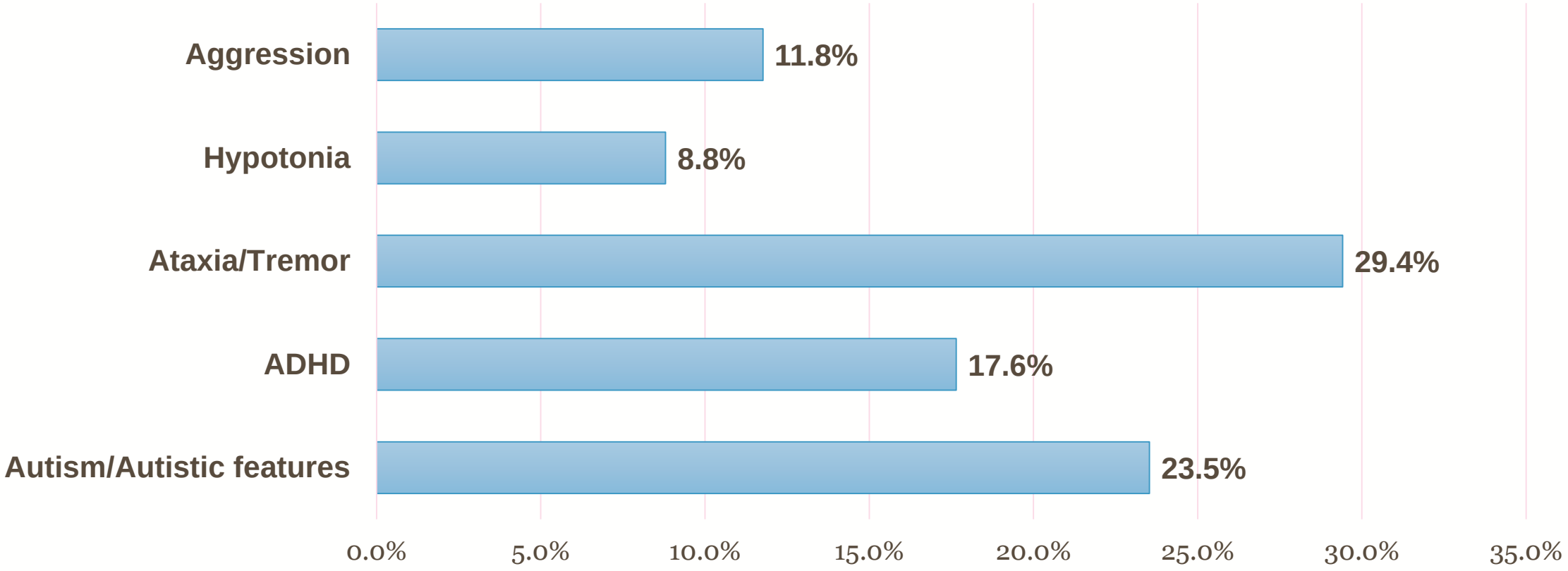
OTHER NEUROLOGICAL FEATURES



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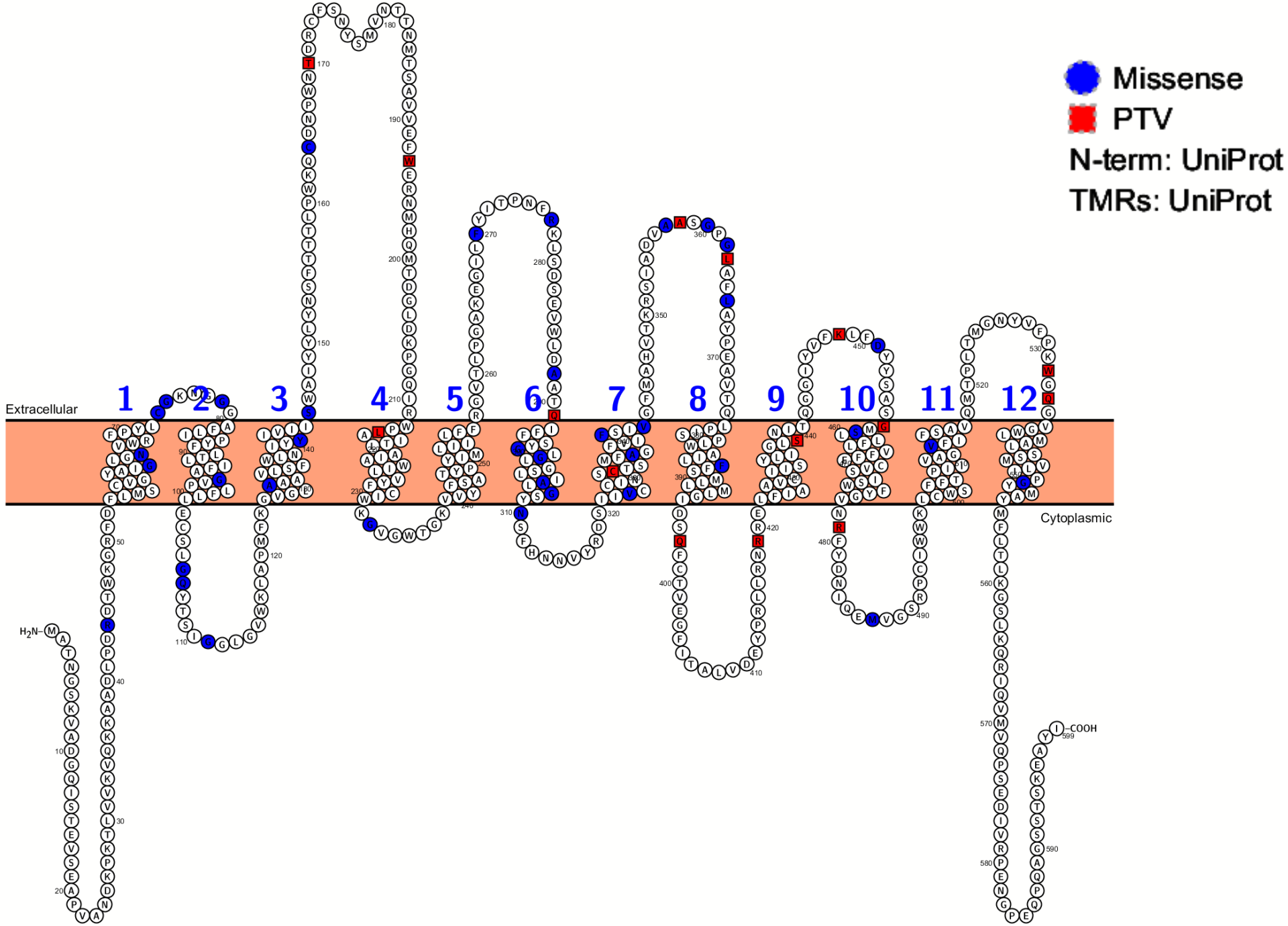
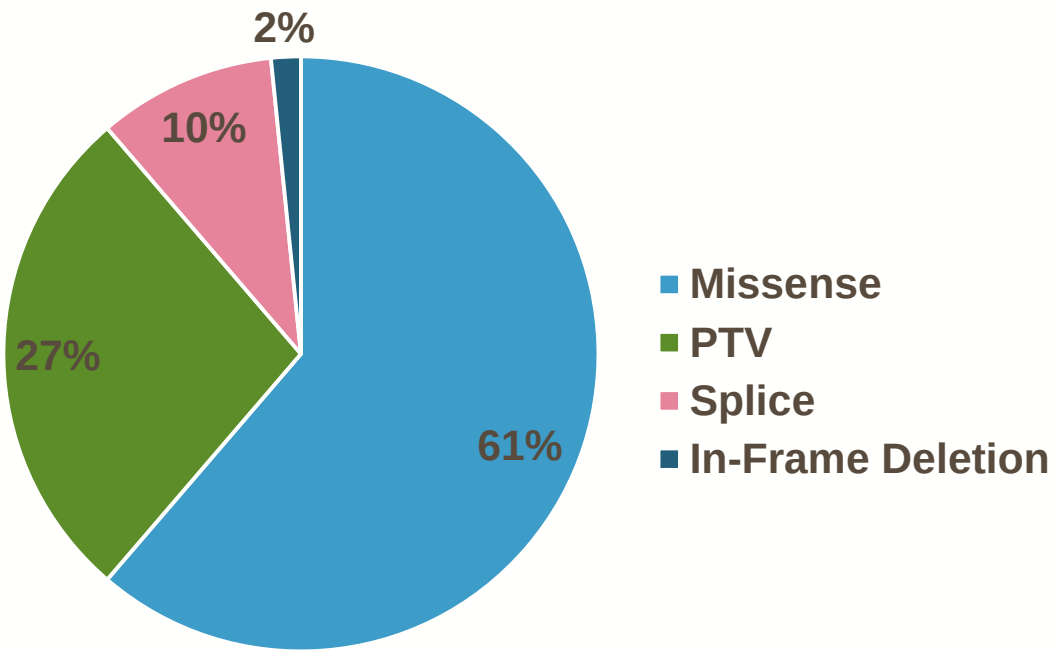
SLC6A1 GENETIC SPECTRUM

- 39 (likely) pathogenic variants reported in ClinVar
- 41 variants reported in HGMD
- Altogether 62 unique *SLC6A1* variants reported
  - Most commonly reported variant c.863C>T; p.(Ala288Val)

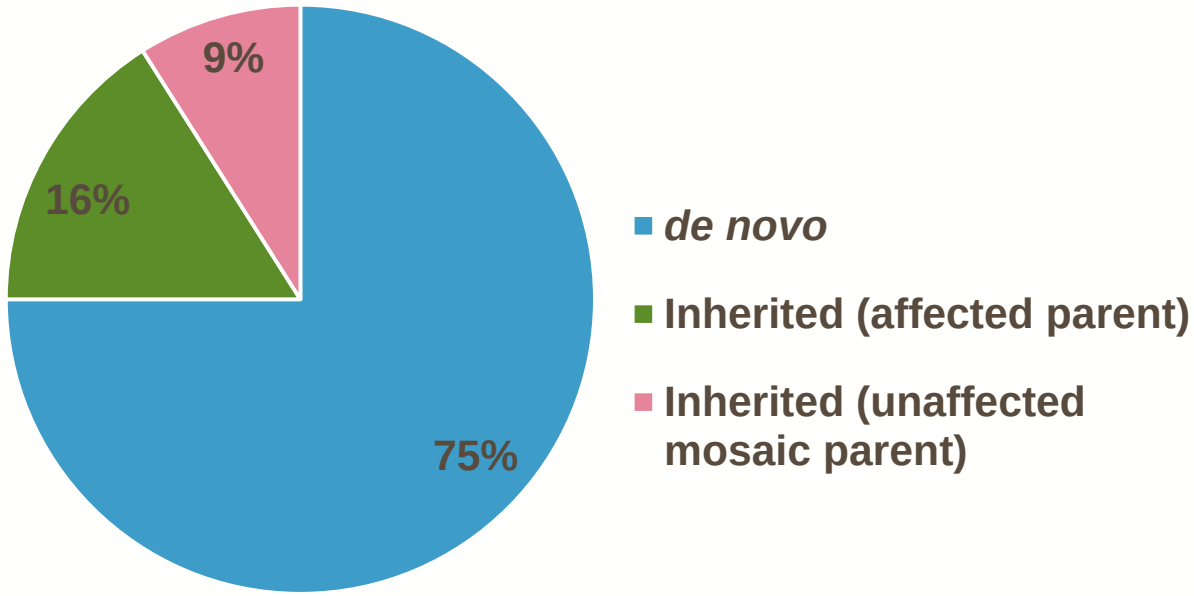

Missense
PTV
N-term: UniProt
TMRs: UniProt
Extracellular
Cytoplasmic
1
2
3
4
5
6
7
8
9
10
11
12
H2N–
–COOH

SLC6A1 GENETIC SPECTRUM

Variant Type



Inheritance of *SLC6A1* Variant



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GENOTYPE-PHENOTYPE CORRELATIONS?

- Not explored in the published literature

- Based on available data, no correlation between genotype and phenotype
  - Systematic studies of genotype-phenotype correlations needed
  - Correlation between function and phenotype?

SUMMARY

- Childhood-onset generalized epilepsy in 80%
  - Median onset 24 months
  - Most common seizure types: absence (typical and atypical), atonic
  - >60% Epilepsy with Myoclonic-Atonic Seizures (MAE, Doose syndrome)
  - Seizures can usually be well-controlled with AEDs
- Developmental delay in >90%
  - Often apparent before seizure onset
  - Most often mild to moderate developmental impairment
- Ataxia and coordination difficulties in 30%
- Autism spectrum disorders in 25%
- No clear genotype-phenotype correlations

CHOP NEUROGENETICS

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Neurogenetics Program

Epilepsy Genetics Research Project