

The Simons Searchlight Gene List

The Simons Searchlight gene list contains 147 gene changes (orange) and 20 copy number variants (purple) that are known to be associated with autism and other neurodevelopmental disorders.

Genetic Changes We Study

1q21.1	ACTB	DDX3X	MED13L	SETBP1
2p16.3 deletion	ACTL6B	DEAF1	MEF2C	SETD2
2q34 duplication	ADNP	DHCR7	MEIS2	SETD5
2q37 deletion	ADSL	DLG4	MYT1L	SHANK2
2q37.3 deletion	AFF2	DMPK	NAA15	SIN3A
5q35	ALDH5A1	DNMT3A	NBEA	SLC6A1
6q16 deletion	AHDC1	DSCAM	NCKAP1	SLC9A6
7q11.23 duplication	ANK2	DST	NEXMIF	SMARCA4
9q34 duplication	ANK3	DYRK1A	NIPBL	SMARCC1
15q11.2 BP1-BP2 deletion	ANKRD11	EBF3	NLGN2	SMARCC2
15q13.3 deletion	ARHGEF9	EHMT1	NLGN3	SON
15q15 deletion	ARID1B	EIF3F	NLGN4X	SOX5
15q24 deletion	ARX	EP300	NR3C2	SPAST
16p11.2*	ASH1L	FOXP1	NR4A2	SRCAP
16p12.2 deletion**	ASXL3	GIGYF1	NRXN1	STXBP1
16p13.11 deletion	ATRX	GIGYF2	NRXN2	SYNCRIP
16p13.3 deletion	AUTS2	GRIN1	NRXN3	SYNGAP1
17q21.3	BAZ2B	GRIN2A	NSD1	TANC2
Xp11.22 duplication	BCKDK	GRIN2B	PACS1	TAOK1
Xq28 duplication	BCL11A	GRIN2D	PBRM1	TBR1
	BRSK2	HIVEP2	PHF21A	TCF20
	CACNA1C	HNRNPH2	PHF3	TLK2
	CAPRIN1	HNRNPU	PHIP	TRIO
	CASK	IQSEC2	POMGNT1	TRIP12
	CASZ1	IRF2BPL	PPP3CA	TSHZ3
	CHAMP1	KANSL1	PPP2R1A	UPF3B
	CHD2	KATNAL2	PPP2R5D	USP9X
	CHD3	KCNB1	PSMD12	VPS13B
	CHD8	KDM3B	PTCHD1	WAC
	CIC	KDM5B	RELN	WDFY3
	CNOT3	KDM6B	RERE	ZBTB20
	CREBBP	KMT2A	REST	ZNF292
	CSDE1	KMT2C	RFX3	ZNF462
	CSNK2A1	KMT2E	RIMS1	
	CTBP1	KMT5B	RORB	
	CTCF	MBD5	SCN1A	
	CTNNB1	MBOAT7	SCN2A	
	CUL3	MED13	SCN8A	

* Includes deletions and duplications that include at a minimum the BP4 - BP5 region (proximal) or BP2 - BP3 (distal) region

** Formerly known as 16p12.1 deletion