ClinGen Brain Malformations Variant Curation Expert Panel						
Start Date	Your Name	Do you work for a laboratory that offers fee-for-service testing related to the work of your Expert Panel?	Have you made substantial contributions to the literature implicating a gene:disease relationship that relates to the work of your Expert Panel?		Do you have any other existing or planned independent curation efforts that will potentially overlap with the scope of your ClinGen work? If so, please describe:	Please disclose any other relevant conflicts of interest (e.g. patents, intellectual property ownership, or paid consultancies related to any variants or genes associated with the work of your Expert Panel):
	Open-Ended Response	Response	Response	If yes, please list the genes:	Open-Ended Response	Open-Ended Response
2018-09-17 11:55:42	Ryan Doan	No	No		No	
2018-07-24 8:39:41	Michael Evenson	Yes	No		No	
2018-06-26 14:52:06	Ghayda Mirzaa	Yes	Yes	PIK3CA, MTOR, PIK3R2, CCND2, AKT3, CENPE, ARX, CDKL5, STAMBP, KIF11, CDC42	I'm a member of the ClinGen Epilepsy Group.	
2018-06-19 10:22:16	Yi-Shan Lee	Yes	No		No	No
2018-05-04 16:19:35	Abbe Lai	No	No			

2018-05-01 12:00:42	Jamel	No	Yes	DCX, TUB1A1, TUBB2B, TUBB3, TUBB5, TUBG1, KIF2A, KIF5C, TUBG1, DYNC1H1, TBC1D23, NEDD4L		
2018-04-24 21:00:34	Christopher Walsh	No	No			
2018-04-24 10:58:47	Heather Mefford	No	Yes	DEPDC5 (in epilepsy)	No	
2018-04-16 12:28:40	Lance Rodan	No	No		No	
2018-04-04 7:22:32	CHRISTELLE MOUFAWAD EL ACHKAR	No	No			
2018-03-27 13:45:05	David Miller	No	No		Only the overlap between my involvement with ClinGen ID/Autism and ClinGen Brain Malformations groups	

2018-03-27 8:21:32	Elizabeth C Engle	No	Yes	KIF21A, TUBB3, TUBB2B, PHOX2A, SALL4, CHN1, HOXA1, HOXB1, ECEL1, MYMK, others.	Our research effort is to continue to define new genetic causes of the CCDDs and determine the spectrum of mutations underlying new syndromes.	Any IP is determined by Boston Children's Hospital.
2018-03-27 6:32:33	Renzo Guerrini	No	Yes	MTOR, PIK3R2, AKT3, PIK3CA	No	
2018-03-26 17:24:01	A. James Barkovich	No	Yes	MTOR, AKT, PI3K, DEPDCs	No	
2018-03-20 11:43:39	Annapurna Poduri	No	Yes	AKT3 PIK3CA DEPDC5 mTOR	Epilepsy Panel (Pediatric Neurology)	n/a
2018-03-20 11:14:59	Tim Yu	No	Yes	I have contributed to the identification of many genes for brain malformations and for autism and intellectual disability	No	I do scientific consulting for gene discovery and clinical genomic interpretation, but not specifically for any genes within the work of our Expert panel

2018-03-20 9:18:46	Ganeshwaran H. Mochida	No	Yes	ASPM, VPS13B, TRAPPC9, SPART, WDR62, JAM3, NDE1, CHMP1A, GRID2, METTL23, QARS, KATNB1, PCLO, PYCR2, DIAPH1, GPT2, AARS, DONSON	No	
2018-03-05 21:41:35	Christopher Yuskaitis	No	No			
2018-03-06 9:00:37	Eric Marsh	No	Yes	Arx	No	
2018-03-06 8:45:52	Lacey Smith	No	Yes	PCDH19, RHOBTB2, PACS2	No	
2018-03-05 18:35:34	Anne Rochtus	No	No			
2018-03-05 16:47:36	Edward Yang	No	No		No/N.A.	Have done consulting for a company (Corticometrics LLC) that develops computer software for detection of cortical dysplasia from brain MRI. Genetics of brain malformations is well outside of the scope of the company's work.