ClinGen Epilepsy Gene Curation Expert Panel						
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
2019-04-30 10:13:29	Kelly Toner	No	No		No	N/A
2019-04-25 11:51:00	Madeline Hughes	No	No		No	
2019-02-05 17:34:18	Erica Smith	Yes	No		I participate in gene-condition curation activities for my company as part of my employment.	none

2019-02-05 9:29:04	Saadet Andrews	No	Yes	NEUROD2 (PIMD 30323019) TLK2 (PIMD 29861108) NTRK2, GABRB2, CLTC, DHDDS, NUS1, RAB11A, GABBR2, and SNAP25 (PIMD 29100083) MED23 (PIMD 27311965) FRRS1L (PIMD 27236917)		We have creatine deficiency disorders expert panel that I chair, recently accepted by ClinGen/ClinVar as expert panel.
2019-01-03 17:26:47	Colin Ellis	No	No		No	
2019-01-02 15:52:36	Benjamin Kang	Yes	No		No	I do not have any relevant conflicts of interest
2018-05-08 15:37:08	Tanya Bardakjian	No	No		no	none
2018-05-07 13:55:32	Ingo Helbig	No	Yes	SCN2A, SCN3A, KCNA2, CHD2, GRIN1, GRIN2A, STXBP1, STX1B	no	none
2018-04-26 19:16:59	Philippe Campeau	No	Yes	TBC1D24, ATP6V1B2, PIGN, PIGC, PIGP, GPAA1, FHF1, KCNH1	I curate LOVD databases for some of the genes above: TBC1D24 and http://www.gpibiosynthesis.org/	

2018-04-20 9:24:38	Bekim Sadikovic	Yes	No			I work for an academic hospital (Western University, London Health Sciences) as a lab director. Lab performs testing for numerous genetic disorders including genes involved in
2018-03-23 13:20:50	David Dyment	No	Yes	ASAH1 SYNJ1 NALCN SCN1A	No	I have no conflicts
2018-03-23 11:44:01	Annapurna Poduri	No	Yes	PCDH19 KCNQ2 KCNT1 SCN2A PRRT2 GABRG2 DNM1	brain malformation panel--minimal overlap (DEPDC5, for example) may exist	none
2018-03-22 6:32:06	Maria Roberta Cilio	No	Yes	KCNQ2/3 KCNT1	No	
2018-03-20 16:45:02	Ghayda Mirzaa	Yes	Yes	ARX, CDKL5, PIK3CA, MTOR, AKT3, DEPDC5, CENPE, STAMBP, PIK3R2, CCND2	The European COST consortium has a project aimed at curating genes associated with brain malformations - many of which overlap with isolated epilepsy phenotypes.	
2018-03-19 18:05:26	Slavé Petrovski	No	Yes	Involved in gene discoveries of GABRB3, ALG13, DNM1, SLC1A2 and GNB1 through consortia efforts.	No	Employee of AstraZeneca (outside of epilepsy research).

2018-03-19 14:49:22	Erin Riggs	No	No		N/A	N/A
2018-03-12 14:09:55	Carrie-Anne Barry	No	No		N/A	N/A
2018-03-12 8:18:19	Rebecca K Siegert	No	No			Currently also a member of the ClinGen Hearing Loss Gene and Variant Curation efforts, the ClinGen ID/Autism Gene Curation efforts, as well as Variant Discrepancy Resolution efforts.
2018-03-09 13:42:21	Elizabeth Butler	Yes	No		I assist with variant curation for clinical testing sent to GeneDx. I don't have any other existing or planned curation efforts outside of my normal daily job responsibilities.	
2018-03-09 11:10:42	Karl Martin Klein	No	Yes	CDKL5, DEPDC5, LGI1, ARHGEF9 and multiple genes within the EuroEPINOMICS consortium	no	none
2018-03-07 15:54:58	Erika Axeen	No	No		no	none
2018-03-07 14:49:38	Jacy Wagnon	No	Yes	SCN8A	No	Nothing to disclose.

2018-03-07 14:07:46	Heather Mefford	No	Yes	CHD2, SLC6A1, GRIN2A, CACNA1A, SLC1A2, PPP3CA, GABRB3, DNM1, ALG13, SYNGAP1	No	
2018-03-07 13:04:34	Tristan T. Sands	No	Yes	KCNQ2 and KCNQ3 variants for benign familial neonatal epilepsy		
2018-03-07 11:02:14	Katherine Helbig	No	Yes	DNM1 KCNA2 GRIN1 NEXMIF/KIAA2022 SCN3A SLC6A1 SPTAN1 GRIN2B PCDH19 SCN2A GABRA1 STXBP1 SYNJ1	N/A	N/A
2018-03-07 10:54:06	Andrew Grant	Yes	No		I have done/will do curation for the EIEE, ID/Autism, RASopathy, and Hearing Loss working groups	
2018-03-07 10:50:35	Kristen Park	No	Yes	KCNQ2	No	
2018-03-07 10:50:04	Pasquale Striano	No	Yes	SXTBP1, KCNQ2, SCN2A	NA	
2018-03-07 10:47:34	Courtney Thaxton	No	No			
2018-03-07 10:46:25	Khalida Liaquat	Yes	No		NO	I work for Quest Diagnostics.