Table S2. Mapping genes targeted by SNP sets to Mental and Brain and Nervous System Disorder categories. (Information obtained fron Nextbio database)

2.55.0.0.0			Up/Down		
Gene	Disease	Score	regulated	References	
7SK	Autistic disorder	39	up-regulated	(1)	
75K 7SK	Encephalomyelopathy	32	up-regulated	(2)	
75K	Mood disorder	51	down-regulated	(3)	
75K	Multiple sclerosis	27	up-regulated	(4)	
ABCC12	Alzheimer's disease	55	down-regulated	(5)	
ABCC12	Dementia	55	down-regulated	(5)	
ABCC12	Disorder of basal ganglia	2	up-regulated	(6)	
ABCC12	Hypoxia of brain	8	up-regulated	(7)	
ABCC12	Meningitis	14	up-regulated	(8)	
ABCC12	Movement disorder	1	up-regulated	(6)	
ABCC12	Multiple sclerosis	37	down-regulated	(9) (10)	
ABCC12	Nerve injury	25	down-regulated	(11) (12) (13)	
ABCC12	Neuropathy	14	down-regulated	(11) (12) (13)	
ABCC12	Parkinson's disease	10	up-regulated	(6)	
ABCC12	Psychotic disorder	47	up-regulated	(15) (16) (17)	
ABCC12	Schizophrenia	47	up-regulated	(15) (16) (17)	
ARPC5L	Alzheimer's disease	26	down-regulated	(18)	
71111 C3L	Amyotrophic lateral	20	down regulated	(10)	
ARPC5L	sclerosis	14	down-regulated	(19)	
			30.1111	( - /	
ARPC5L	Anxiety disorder	73	up-regulated	(20)	
ARPC5L	Autistic disorder	45	down-regulated	(21) (22) (23)	
ARPC5L	Cerebrovascular disease	45	up-regulated	(24)	
ARPC5L	Chronic fatigue syndrome	100	down-regulated	(25)	
ARPC5L	Dementia	26	down-regulated	(18)	
	Developmental mental		_		
ARPC5L	disorder	41	up-regulated	(22)	
				(26) (27) (28) (29)	
ARPC5L	Disorder of basal ganglia	74	down-regulated	(30) (6)	
ARPC5L	Disorder of brain	38	up-regulated	(31) (32)	
ARPC5L	Huntington's disease	85	down-regulated	(27) (29)	
ARPC5L	Meningitis	69	down-regulated	(33) (8) (34) (35)	
ARPC5L	Mental retardation	38	up-regulated	(22)	
ARPC5L	Motor neuron disease	28	up-regulated	(36) (19)	
				(26) (27) (28) (29)	
ARPC5L	Movement disorder	71	down-regulated	(30) (6)	
ARPC5L	Nerve injury	1	down-regulated	(19)	
ARPC5L	Parkinson's disease	50	down-regulated	(26) (28) (30) (6)	

ARPC5L	Prion disease	26	down-regulated	(37)
ARPC5L	Psychotic disorder	36	down-regulated	(15) (38)
ARPC5L	Schizophrenia	36	down-regulated	(15) (38)
ATP8A2	Alzheimer's disease	44	down-regulated	(39) (18) (5) (40)
ATP8A2	Autistic disorder	23	up-regulated	(1)
ATP8A2	Cerebrovascular disease	29	down-regulated	(41)
ATP8A2	Dementia	43	down-regulated	(39) (18) (5) (40)
				(10) (26) (42) (28)
ATP8A2	Disorder of basal ganglia	84	down-regulated	(29) (6)
ATP8A2	Encephalitis	46	down-regulated	(43)
ATP8A2	Encephalomyelopathy	37	up-regulated	(44)
ATP8A2	Huntington's disease	80	down-regulated	(27) (29)
ATP8A2	Hypoxia of brain	32	down-regulated	(41)
ATP8A2	Meningitis	55	up-regulated	(33) (8) (35) (45)
				(10) (26) (42) (28)
ATP8A2	Movement disorder	81	down-regulated	(29) (6)
ATP8A2	Nerve injury	31	up-regulated	(12)
ATP8A2	Neuropathy	33	down-regulated	(46) (47)
				(10) (48) (26) (42)
ATP8A2	Parkinson's disease	84	down-regulated	(28) (6)
ATP8A2	Prion disease	40	down-regulated	(49)
ATP8A2	Psychotic disorder	30	0.0001 p-value	(38)
ATP8A2	Schizophrenia	30	0.0001 p-value	(38)
ATP8A2	Sleep disorder	34	down-regulated	(50)
C14orf102	Alzheimer's disease	48	up-regulated	(51) (5) (52)
C14orf102	Anxiety disorder	17	up-regulated	(20)
C14orf102	Autistic disorder	27	up-regulated	(1)
C14orf102	Cerebrovascular disease	20	down-regulated	(41)
C14orf102	Dementia	48	up-regulated	(51) (5) (52)
C14orf102	Disorder of basal ganglia	18	up-regulated	(10) (26) (27)
C14orf102	Huntington's disease	24	down-regulated	(10) (27)
C14orf102	Hypoxia of brain	22	down-regulated	(41)
C14orf102	Meningitis	51	up-regulated	(8) (53)
C14orf102	Movement disorder	15	up-regulated	(10) (26) (27)
C14orf102	Neural tube defect	42	down-regulated	(54)
C14orf102	Neuropathy	14	down-regulated	(55) (47)
C14orf102	Parkinson's disease	8	up-regulated	(26)
C14orf102	Psychotic disorder	20	0.0002 p-value	(38)
C14orf102	Schizophrenia	21	0.0002 p-value	(38)
C14orf102	Sleep disorder	42	down-regulated	(56)

C20orf78	Anxiety disorder	32	down-regulated	(20)
C20orf78	Disorder of basal ganglia	42	down-regulated	(29)
C20orf78	Huntington's disease	55	down-regulated	(29)
C20orf78	Movement disorder	39	down-regulated	(29)
C20orf78	Psychotic disorder	35	up-regulated	(57) (38)
C20orf78	Schizophrenia	35	up-regulated	(57) (38)
C4orf37	Autistic disorder	3	up-regulated	(23)
C4orf37	Meningitis	10	up-regulated	(8)
C4orf37	Multiple sclerosis	14	up-regulated	(9)
C4orf37	Psychotic disorder	1	down-regulated	(16)
C4orf37	Schizophrenia	1	down-regulated	(16)
C4orf37	Sleep disorder	16	up-regulated	(58)
C6orf138	Amnestic disorder	88	up-regulated	(59)
C6orf138	Cerebrovascular disease	48	down-regulated	(41)
C6orf138	Disorder of basal ganglia	62	down-regulated	(10)
C6orf138	Huntington's disease	54	down-regulated	(10)
C6orf138	Hypoxia of brain	51	down-regulated	(41)
C6orf138	Meningitis	75	down-regulated	(8) (45)
C6orf138	Movement disorder	59	down-regulated	(10)
C6orf138	Multiple sclerosis	71	down-regulated	(10) (4)
C6orf138	Nerve injury	46	down-regulated	(60) (12) (61)
C6orf138	Neuropathy	83	down-regulated	(47)
C6orf138	Parkinson's disease	63	down-regulated	(10)
CHST9	Alzheimer's disease	21	up-regulated	(5) (52)
CHST9	Amnestic disorder	79	down-regulated	(59)
	Amyotrophic lateral			
CHST9	sclerosis	37	down-regulated	(10)
CHST9	Dementia	21	up-regulated	(5) (52)
CHST9	Disorder of basal ganglia	33	up-regulated	(10) (29)
CHST9	Huntington's disease	47	up-regulated	(10) (29)
CHST9	Meningitis	31	up-regulated	(8)
CHST9	Motor neuron disease	46	down-regulated	(10) (62)
CHST9	Movement disorder	30	up-regulated	(10) (29)
CHST9	Multiple sclerosis	56	up-regulated	(63) (64) (65)
CHST9	Nerve injury	24	down-regulated	(11) (12) (13)
CHST9	Neuropathy	11	down-regulated	(11)
CHST9	Psychotic disorder	69	down-regulated	(38) (66)
CHST9	Schizophrenia	69	down-regulated	(38) (66)
CSMD1	Alzheimer's disease	38	8.7E-6 p-value	(40) (67) (68) (69)
	Attention deficit			
CSMD1	hyperactivity disorder	35		(70)
CSMD1	Autistic disorder	38	down-regulated	(71) (23)
CSMD1	Cerebrovascular disease	10	5.4E-5 p-value	(72)

CSMD1	Dementia	37	8.7E-6 p-value	(40) (67) (68) (69)
				(10) (26) (28) (29)
CSMD1	Disorder of basal ganglia	49	down-regulated	(73)
CSMD1	Huntington's disease	33	down-regulated	(10) (29)
CSMD1	Hypoxia of brain	13	5.4E-5 p-value	(72)
CSMD1	Meningitis	28	up-regulated	(33) (8)
CSMD1	Mood disorder	38	3.6E-6 p-value	(74) (75) (76) (77)
				(10) (26) (28) (29)
CSMD1	Movement disorder	46	down-regulated	(73)
CSMD1	Multiple sclerosis	45	up-regulated	(10) (64) (78)
CSMD1	Nerve injury	23	down-regulated	(11) (12)
CSMD1	Neuropathy	29	down-regulated	(60) (11) (47)
CSMD1	Parkinson's disease	49	down-regulated	(10) (26) (28) (73)
CSMD1	Psychotic disorder	71	down-regulated	(79) (38) (66) (80)
CSMD1	Schizophrenia	71	down-regulated	(79) (38) (66) (80)
DKK4	Autistic disorder	33	up-regulated	(23)
DKK4	Disorder of basal ganglia	1	up-regulated	(6)
DKK4	Encephalomyelopathy	3	up-regulated	(2)
DKK4	Meningitis	28	down-regulated	(8)
DKK4	Mood disorder	43	down-regulated	(17) (3)
DKK4	Movement disorder	1	up-regulated	(6)
DKK4	Multiple sclerosis	4	up-regulated	(81)
DUSP4	Alzheimer's disease	1	down-regulated	(5) (52)
DUSP4	Anxiety disorder	38	up-regulated	(20)
DUSP4	Cerebrovascular disease	6	up-regulated	(41)
DUSP4	Disorder of basal ganglia	38	down-regulated	(48) (26) (27) (29) (6)
DUSP4	Disorder of brain	46	down-regulated	(32)
DUSP4	Encephalitis	29	up-regulated	(82)
DUSP4	Encephalomyelopathy	31	down-regulated	(83)
DUSP4	Huntington's disease	46	down-regulated	(27) (29)
DUSP4	Hypoxia of brain	16	up-regulated	(84) (41)
DUSP4	Meningitis	53	up-regulated	(8) (35) (53) (45)
DUSP4	Mood disorder	23	down-regulated	(85) (86)
DUSP4	Movement disorder	35	down-regulated	(48) (26) (27) (29) (6)
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DUSP4	Multiple sclerosis	11	down-regulated	(63) (65)
				(12) (87) (88) (61)
DUSP4	Nerve injury	20	up-regulated	(13) (89)

DUSP4	Neural tube defect	29	down-regulated	(54)
				(60) (90) (88) (47)
DUSP4	Neuropathy	17	down-regulated	(89)
DUSP4	Paralytic syndrome	24	up-regulated	(91)
DUSP4	Parkinson's disease	12	down-regulated	(48) (26) (6)
DUSP4	Psychotic disorder	22	down-regulated	(15) (66)
DUSP4	Schizophrenia	22	down-regulated	(15) (66)
DUSP4	Sleep disorder	91	up-regulated	(56) (50) (58)
DUSP4	Spinocerebellar ataxia	51	down-regulated	(83)
EML5	Alzheimer's disease	11	down-regulated	(52)
EML5	Amnestic disorder	45	up-regulated	(92)
EML5	Dementia	11	down-regulated	(52)
EML5	Disorder of basal ganglia	66	up-regulated	(10) (26) (29)
EML5	Huntington's disease	78	up-regulated	(10) (29)
EML5	Meningitis	73	down-regulated	(8) (93) (35) (53) (45)
EML5	Movement disorder	63	up-regulated	(10) (26) (29)
				(60) (11) (12) (94)
EML5	Nerve injury	77	down-regulated	(95)
EML5	Neuropathy	73	down-regulated	(60) (11) (94)
EML5	Parkinson's disease	30	up-regulated	(26)
EML5	Psychotic disorder	79	9.5E-7 p-value	(38) (96)
EML5	Schizophrenia	79	9.5E-7 p-value	(38) (96)
EML5	Sleep disorder	76	down-regulated	(56) (50)
EVI5	Amnestic disorder	65	up-regulated	(97)
EVI5	Anxiety disorder	14	up-regulated	(98)
EVI5	Autistic disorder	29	up-regulated	(71) (1)
EVI5	Cerebral palsy	17	up-regulated	(99)
				(10) (48) (26) (27)
EVI5	Disorder of basal ganglia	34	up-regulated	(29)
EVI5	Huntington's disease	39	up-regulated	(10) (27) (29)
				(100) (33) (8) (34)
EVI5	Meningitis	49	up-regulated	(35) (45)
EVI5	Mood disorder	25	down-regulated	(101) (3)
EVI5	Motor neuron disease	3	down-regulated	(36) (62)
E) #=				(10) (48) (26) (27)
EVI5	Movement disorder	31	up-regulated	(29)
EV (15	NA III and	400	6.55.43	(402) (402) (404)
EVI5	Multiple sclerosis	100	6.5E-12 p-value	(102) (103) (104)
EVI5	Nerve injury	72	up-regulated	(12) (88) (105) (106)

		1		(107) (95)
		1		, , ,
EVI5	Neural tube defect	25	up-regulated	(54)
EVI5	Neuropathy	4	up-regulated	(90) (88)
EVI5	Parkinson's disease	23	down-regulated	(48) (26)
EVI5	Psychotic disorder	61	up-regulated	(108) (38) (17)
EVI5	Schizophrenia	62	up-regulated	(108) (38) (17)
EVI5	Sleep disorder	42	up-regulated	(56) (58)
FAM69A	Alzheimer's disease	1	down-regulated	(18)
FAM69A	Autistic disorder	1	down-regulated	(71)
FAM69A	Cerebral palsy	32	down-regulated	(99)
FAM69A	Dementia	1	down-regulated	(18)
FAM69A	Disorder of basal ganglia	1	up-regulated	(28) (109)
FAM69A	Disorder of brain	29	up-regulated	(31) (32)
FAM69A	Encephalitis	44	down-regulated	(110) (82)
FAM69A	Encephalomyelitis	29	down-regulated	(110)
FAM69A	Encephalomyelopathy	9	down-regulated	(91) (110)
FAM69A	Meningitis	7	down-regulated	(33)
FAM69A	Mood disorder	1	down-regulated	(85)
FAM69A	Motor neuron disease	1	up-regulated	(62)
FAM69A	Movement disorder	1	up-regulated	(28) (109)
FAM69A	Multiple sclerosis	90	0.8E-7 p-value	(102) (103)
FAM69A	Myoneural disorder	40	up-regulated	(111)
FAM69A	Nerve injury	17	down-regulated	(112) (87) (95)
FAM69A	Neuropathy	11	up-regulated	(91) (111)
FAM69A	Paralytic syndrome	20	down-regulated	(91)
FAM69A	Parkinson's disease	5	up-regulated	(28) (109)
FAM69A	Prion disease	6	down-regulated	(113)
FAM69A	Psychotic disorder	51	0.0E-6 p-value	(108) (38)
FAM69A	Schizophrenia	51	0.0E-6 p-value	(108) (38)
FAM69A	Sleep disorder	39	down-regulated	(56) (50)
FOXR2	Nerve injury	83	up-regulated	(11) (89)
FOXR2	Neuropathy	86	up-regulated	(11) (89)
GOLGA1	Alzheimer's disease	24	0.0007 p-value	(40)
GOLGA1	Autistic disorder	44	down-regulated	(23)
GOLGA1	Dementia	24	0.0007 p-value	(40)
GOLGA1	Disorder of basal ganglia	55	up-regulated	(10) (26) (28) (29)
GOLGA1	Disorder of brain	50	down-regulated	(31)
GOLGA1	Encephalomyelopathy	51	down-regulated	(114)
GOLGA1	Huntington's disease	52	up-regulated	(29)
GOLGA1	Meningitis	51	down-regulated	(100)
GOLGA1	Movement disorder	52	up-regulated	(10) (26) (28) (29)
GOLGA1	Multiple sclerosis	33	down-regulated	(81)

GOLGA1	Nerve injury	66	down-regulated	(12) (89)
GOLGA1	Neuropathy	35	down-regulated	(47) (89)
GOLGA1	Paralytic syndrome	61	up-regulated	(91)
GOLGA1	Parkinson's disease	55	up-regulated	(10) (26) (28)
GOLGA1	Psychotic disorder	50	0.0002 p-value	(38)
GOLGA1	Schizophrenia	51	0.0002 p-value	(38)
GOLGA1	Sleep disorder	91	down-regulated	(50)
GP2	Alzheimer's disease	1	up-regulated	(18)
GP2	Amnestic disorder	20	up-regulated	(115)
GP2	Anxiety disorder	1	down-regulated	(20)
GP2	Dementia	1	up-regulated	(18)
GP2	Disorder of basal ganglia	1	down-regulated	(27)
GP2	Huntington's disease	1	down-regulated	(27)
GP2	Meningitis	9	down-regulated	(8) (34)
GP2	Movement disorder	1	down-regulated	(27)
GP2	Nerve injury	35	down-regulated	(88)
GP2	Neuropathy	38	down-regulated	(88)
GP2	Psychotic disorder	12	up-regulated	(17)
GP2	Schizophrenia	12	up-regulated	(17)
GPR119	Alzheimer's disease	59	7.8E-5 p-value	(116)
GPR119	Anxiety disorder	48	down-regulated	(20)
GPR119	Dementia	58	7.8E-5 p-value	(116)
GPR119	Nerve injury	27	up-regulated	(11)
GPR119	Neuropathy	29	up-regulated	(11)
HACE1	Alzheimer's disease	1	down-regulated	(39) (5) (40) (69)
HACE1	Autistic disorder	1	up-regulated	(21)
HACE1	Cerebrovascular disease	1	up-regulated	(24)
HACE1	Dementia	1	down-regulated	(39) (5) (40) (69)
				(117) (26) (27) (28)
HACE1	Disorder of basal ganglia	11	down-regulated	(29)
HACE1	Encephalitis	1	down-regulated	(82)
HACE1	Huntington's disease	16	down-regulated	(117) (27) (29)
HACE1	Meningitis	3	up-regulated	(8) (45)
HACE1	Mood disorder	1	0.0003 p-value	(74)
114.051	NA over one discorder	0	dayya waaydatad	(117) (26) (27) (28)
HACE1	Movement disorder	8	down-regulated	(29)
HACE1	Multiple sclerosis	1	up-regulated	(78)
HACE1	Nerve injury	6	up-regulated	(12) (88) (105) (107)
HACE1	Neuropathy	1	down-regulated	(60) (88)
HACE1	Parkinson's disease	1	down-regulated	(26) (28)
HACE1	Psychotic disorder	7	0.5E-6 p-value	(108)

HACE1	Schizophrenia	7	0.5E-6 p-value	(108)
HACE1	Sleep disorder	8	up-regulated	(58)
HPGDS	Alzheimer's disease	37	4.0E-5 p-value	(118)
HPGDS	Amnestic disorder	49	up-regulated	(115) (59)
HPGDS	Anxiety disorder	27	up-regulated	(20)
HPGDS	Cerebral palsy	54	up-regulated	(99)
	Childhood disorder of			
HPGDS	conduct and emotion	59	down-regulated	(119) (120)
HPGDS	Dementia	37	4.0E-5 p-value	(118)
HPGDS	Disorder of basal ganglia	37	down-regulated	(48) (27) (121)
HPGDS	Disorder of brain	44	down-regulated	(122)
HPGDS	Huntington's disease	42	down-regulated	(27) (121)
HPGDS	Meningitis	23	down-regulated	(33)
HPGDS	Movement disorder	34	down-regulated	(48) (27) (121)
HPGDS	Multiple sclerosis	13	up-regulated	(78)
				(112) (12) (87) (95)
HPGDS	Nerve injury	78	up-regulated	(13) (89)
HPGDS	Neuropathy	43	down-regulated	(60) (90) (46) (89)
HPGDS	Parkinson's disease	29	down-regulated	(48)
HPGDS	Prion disease	75	up-regulated	(37) (113)
HPGDS	Psychotic disorder	16	0.0003 p-value	(38)
HPGDS	Schizophrenia	16	0.0003 p-value	(38)
HPGDS	Sleep disorder	45	down-regulated	(50)
IGSF1	Amnestic disorder	39	up-regulated	(92) (97)
IGSF1	Autistic disorder	20	up-regulated	(23) (1)
				(10) (48) (42) (27)
IGSF1	Disorder of basal ganglia	60	up-regulated	(28) (29)
IGSF1	Disorder of brain	16	up-regulated	(32)
IGSF1	Encephalitis	47	down-regulated	(43)
IGSF1	Encephalomyelopathy	20	up-regulated	(2)
IGSF1	Epilepsy	14	up-regulated	(123)
IGSF1	Huntington's disease	70	up-regulated	(10) (27) (29)
IGSF1	Meningitis	31	up-regulated	(33) (8)
IGSF1	Mood disorder	6	up-regulated	(124)
IGSF1	Motor neuron disease	21	up-regulated	(62)
				(10) (48) (42) (27)
IGSF1	Movement disorder	57	up-regulated	(28) (29)
IGSF1	Multiple sclerosis	1	up-regulated	(9)
				(112) (11) (12) (87)
IGSF1	Nerve injury	48	down-regulated	(107) (89)

IGSF1	Neuropathy	32	down-regulated	(11) (88) (47) (89)
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IGSF1	Parkinson's disease	29	down-regulated	(48) (42) (28)
IGSF1	Psychotic disorder	17	up-regulated	(38) (17)
IGSF1	Schizophrenia	18	up-regulated	(38) (17)
IGSF1	Sleep disorder	84	down-regulated	(56) (58)
ITFG1	Alzheimer's disease	44	down-regulated	(18) (5) (52)
ITFG1	Autistic disorder	12	down-regulated	(1)
ITFG1	Cerebral palsy	27	up-regulated	(125)
ITFG1	Cerebrovascular disease	9	down-regulated	(24) (41)
ITFG1	Chronic fatigue syndrome	78	up-regulated	(25)
ITFG1	Dementia	43	down-regulated	(18) (5) (52)
ITFG1	Disorder of basal ganglia	78	down-regulated	(10) (26) (28) (29) (6)
ITFG1	Disorder of brain	20	up-regulated	(31) (32)
ITFG1	Encephalomyelopathy	21	down-regulated	(83) (91)
ITFG1	Epilepsy	8	down-regulated	(123)
ITFG1	Huntington's disease	86	down-regulated	(10) (117) (29)
ITFG1	Hypoxia of brain	2	down-regulated	(41)
ITFG1	Meningitis	44	up-regulated	(33) (8) (34)
ITFG1	Mood disorder	37	down-regulated	(85) (3)
ITFG1	Movement disorder	75	down-regulated	(10) (26) (28) (29) (6)
ITFG1	Multiple sclerosis	24	down-regulated	(9) (10) (63) (65)
				(112) (12) (88) (106)
ITFG1	Nerve injury	28	down-regulated	(95) (13)
ITFG1	Neuropathy	10	down-regulated	(11) (91) (88) (47)
ITFG1	Paralytic syndrome	42	down-regulated	(91)
ITFG1	Parkinson's disease	62	down-regulated	(26) (28) (6)
ITFG1	Prion disease	20	down-regulated	(37) (113)
ITFG1	Psychotic disorder	22	down-regulated	(15)
ITFG1	Schizophrenia	23	down-regulated	(15)
ITFG1	Sleep disorder	1	down-regulated	(56)
ITFG1	Spinocerebellar ataxia	16	up-regulated	(83)
MAGEH1	Anxiety disorder	46	up-regulated	(20)
MAGEH1	Autistic disorder	22	down-regulated	(1)
MAGEH1	Disorder of basal ganglia	44	up-regulated	(26) (27) (6)
MAGEH1	Encephalomyelopathy	33	down-regulated	(83) (2)
MAGEH1	Huntington's disease	48	up-regulated	(27)
MAGEH1	Meningitis	41	up-regulated	(8) (34)
MAGEH1	Mood disorder	8	down-regulated	(126)
MAGEH1	Movement disorder	41	up-regulated	(26) (27) (6)

MAGEH1	Myoneural disorder	54	up-regulated	(111)
MAGEH1	Nerve injury	57	down-regulated	(11) (88) (127) (105) (106) (13)
MAGEH1	Neuropathy	41	up-regulated	(11) (90) (88) (127) (128) (111)
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MAGEH1	Paralytic syndrome	40	up-regulated	(91)
MAGEH1	Parkinson's disease	36	down-regulated	(26) (6)
MAGEH1	Prion disease	30	down-regulated	(37)
MAGEH1	Psychotic disorder	22	down-regulated	(57)
MAGEH1	Schizophrenia	23	down-regulated	(57)
MAGEH1	Spinocerebellar ataxia	43	down-regulated	(83)
NALCN	Alzheimer's disease	68	down-regulated	(5) (40) (67)
NALCN	Amnestic disorder	54	down-regulated	(115) (129)
NALCN	Anxiety disorder	56	up-regulated	(20)
NALCN	Cerebrovascular disease	23	down-regulated	(41)
NALCN	Dementia	67	down-regulated	(5) (40) (67)
NALCN	Disorder of basal ganglia	44	up-regulated	(10) (26) (121) (6)
NALCN	Epilepsy	76	3.6E-6 p-value	(130)
NALCN	Huntington's disease	47	up-regulated	(10) (121)
NALCN	Hypoxia of brain	25	down-regulated	(41)
NALCN	Meningitis	48	down-regulated	(100)
NALCN	Mood disorder	45	3.3E-5 p-value	(131) (74)
NALCN	Movement disorder	41	up-regulated	(10) (26) (121) (6)
NALCN	Multiple sclerosis	8	down-regulated	(78)
NALCN	Myoneural disorder	39	down-regulated	(132)
NALCN	Nerve injury	55	down-regulated	(112) (12) (87) (88) (106) (95)
NALCN	Neuropathy	40	down-regulated	(132) (88) (47) (89)
NALCN	Parkinson's disease	39	up-regulated	(26) (6)
NALCN	Prion disease	30	down-regulated	(113)
NALCN	Psychotic disorder	51	up-regulated	(38) (66)
NALCN	Schizophrenia	52	up-regulated	(38) (66)
NCAM1	Amnestic disorder	1	down-regulated	(115)
NCAM1	Autistic disorder	1	down-regulated	(23)
NCAM1	Dementia	1	up-regulated	(51) (40)
NCAM1	Disorder of basal ganglia	32	down-regulated	(10) (48) (26) (28) (29) (109)
NCAM1	Huntington's disease	36	up-regulated	(29) (121)
NCAM1	Meningitis	33	up-regulated	(33) (8) (34)

1	Ī	1		(10) (48) (26) (28)
NCAM1	Movement disorder	29	down-regulated	(29) (109)
110/11/11	Wievernene disorder		down regulated	(10) (48) (26) (42)
NCAM1	Parkinson's disease	23	up-regulated	(28) (109)
			ap regulated.	(10) (100)
NCAM1	Psychotic disorder	16	down-regulated	(57) (38) (66)
NCAM1	Schizophrenia	17	down-regulated	(57) (38) (66)
NCAM1	Sleep disorder	11	down-regulated	(56) (58)
NETO2	Amnestic disorder	41	down-regulated	(59)
NETO2	Anxiety disorder	36	up-regulated	(133) (20)
NETO2	Dementia	43	down-regulated	(18) (5) (52)
		1.0	a a mini ragana caa	(10) (26) (28) (29)
NETO2	Disorder of basal ganglia	79	down-regulated	(121) (6)
				7(-7
NETO2	Huntington's disease	90	down-regulated	(10) (29) (121)
NETO2	Mood disorder	21	down-regulated	(85)
				(10) (26) (28) (29)
NETO2	Movement disorder	76	down-regulated	(121) (6)
				(12) (88) (106) (107)
NETO2	Nerve injury	54	down-regulated	(95)
	,			
NETO2	Parkinson's disease	48	down-regulated	(10) (48) (26) (28) (6)
NETO2	Psychotic disorder	32	up-regulated	(15) (16)
NETO2	Schizophrenia	32	up-regulated	(15) (16)
NETO2	Sleep disorder	52	up-regulated	(56) (50)
NTRK3	Alzheimer's disease	26	up-regulated	(39)
NTRK3	Amnestic disorder	59	up-regulated	(92) (115)
NTRK3	Autistic disorder	48	down-regulated	(22) (23)
NTRK3	Cerebral palsy	65	down-regulated	(125)
NTRK3	Cerebrovascular disease	33	down-regulated	(41)
NTRK3	Chronic fatigue syndrome	85	down-regulated	(25)
NTRK3	Dementia	26	up-regulated	(39)
	Developmental mental			
NTRK3	disorder	50	down-regulated	(22)
				(48) (26) (27) (28)
NTRK3	Disorder of basal ganglia	69	down-regulated	(29) (30)
NTRK3	Encephalitis	68	down-regulated	(82)
NTRK3	Huntington's disease	76	down-regulated	(27) (29)
NTRK3	Hypoxia of brain	36	down-regulated	(41)
NTRK3	Meningitis	80	down-regulated	(33) (8) (34) (35) (45)
			_	
NTRK3	Mental retardation	48	down-regulated	(22)
NTRK3	Movement disorder	66	down-regulated	(48) (26) (27) (28)

				(29) (30)
				, , , ,
NTRK3	Multiple sclerosis	56	up-regulated	(9) (81) (65)
				(11) (12) (127) (107)
NTRK3	Nerve injury	91	down-regulated	(95) (89)
NTRK3	Neural tube defect	53	up-regulated	(54)
				(11) (134) (127) (47)
NTRK3	Neuropathy	68	down-regulated	(14) (89)
NTRK3	Parkinson's disease	53	down-regulated	(48) (26) (28) (30)
NTRK3	Prion disease	63	up-regulated	(37)
NTRK3	Psychotic disorder	94	up-regulated	(57) (38) (80)
NTRK3	Schizophrenia	94	up-regulated	(57) (38) (80)
NTRK3	Sleep disorder	64	down-regulated	(56) (50)
OPN5	Disorder of basal ganglia	27	down-regulated	(10)
OPN5	Meningitis	70	up-regulated	(8) (34)
OPN5	Movement disorder	24	down-regulated	(10)
OPN5	Neuropathy	29	down-regulated	(60)
OPN5	Parkinson's disease	35	down-regulated	(10)
OPN5	Psychotic disorder	68	up-regulated	(17)
OPN5	Schizophrenia	68	up-regulated	(17)
PAGE3	Disorder of basal ganglia	77	down-regulated	(48)
PAGE3	Movement disorder	74	down-regulated	(48)
PAGE3	Parkinson's disease	85	down-regulated	(48)
PAGE5	Disorder of basal ganglia	52	down-regulated	(48) (29) (6)
PAGE5	Huntington's disease	36	down-regulated	(29)
PAGE5	Meningitis	47	down-regulated	(34)
PAGE5	Movement disorder	49	down-regulated	(48) (29) (6)
PAGE5	Multiple sclerosis	36	up-regulated	(4)
PAGE5	Parkinson's disease	56	down-regulated	(48) (6)
PAGE5	Psychotic disorder	86	up-regulated	(17)
PAGE5	Schizophrenia	87	up-regulated	(17)
PHKB	Alzheimer's disease	2	down-regulated	(39) (18)
PHKB	Anxiety disorder	12	up-regulated	(133)
PHKB	Autistic disorder	7	up-regulated	(21) (23)
PHKB	Cerebral palsy	36	down-regulated	(125) (99)
	Childhood disorder of			
PHKB	conduct and emotion	16	up-regulated	(119)
PHKB	Chronic fatigue syndrome	67	up-regulated	(25)
PHKB	Dementia	2	down-regulated	(39) (18)
				(48) (117) (26) (27)
PHKB	Disorder of basal ganglia	35	down-regulated	(29) (6)

РНКВ	Disorder of brain	2	up-regulated	(32)
PHKB	Encephalomyelopathy	26	down-regulated	(83) (91)
РНКВ	Epilepsy	1	down-regulated	(123)
PHKB	Huntington's disease	29	up-regulated	(117) (27) (29)
PHKB	Meningitis	35	down-regulated	(100) (8) (35) (53)
				(48) (117) (26) (27)
PHKB	Movement disorder	32	down-regulated	(29) (6)
PHKB	Multiple sclerosis	1	down-regulated	(81)
				(87) (88) (61) (106)
PHKB	Nerve injury	25	down-regulated	(107) (89)
PHKB	Neuropathy	23	down-regulated	(11) (91) (88) (89)
PHKB	Paralytic syndrome	46	down-regulated	(91)
				(48) (26) (28) (109)
PHKB	Parkinson's disease	36	down-regulated	(30) (6)
PHKB	Prion disease	15	up-regulated	(49) (113)
PHKB	Sleep disorder	1	up-regulated	(50)
PHKB	Spinocerebellar ataxia	9	up-regulated	(83)
	Attention deficit			
PPP1R1C	hyperactivity disorder	1	0.0003 p-value	(135)
	Developmental mental			
PPP1R1C	disorder	11	down-regulated	(22)
PPP1R1C	Disorder of basal ganglia	1	up-regulated	(26)
PPP1R1C	Meningitis	8	up-regulated	(8)
PPP1R1C	Mental retardation	9	down-regulated	(22)
PPP1R1C	Mood disorder	1	0.0008 p-value	(74)
PPP1R1C	Movement disorder	1	up-regulated	(26)
PPP1R1C	Multiple sclerosis	11	up-regulated	(64) (78)
PPP1R1C	Myoneural disorder	20	down-regulated	(132)
PPP1R1C	Nerve injury	26	up-regulated	(112) (11) (105) (13)
PPP1R1C	Neural tube defect	27	down-regulated	(54)
PPP1R1C	Neuropathy	17	down-regulated	(11) (132) (47)
PPP1R1C	Parkinson's disease	1	up-regulated	(26)
PPP1R1C	Psychotic disorder	4	7.9E-5 p-value	(38)
PPP1R1C	Schizophrenia	4	7.9E-5 p-value	(38)
PSMC1	Alzheimer's disease	41	up-regulated	(39) (51)
PSMC1	Anxiety disorder	40	up-regulated	(98)
PSMC1	Autistic disorder	23	down-regulated	(1)
PSMC1	Cerebrovascular disease	54	down-regulated	(24)

PSMC1	Dementia	41	up-regulated	(39) (51)
				(10) (26) (42) (27)
PSMC1	Disorder of basal ganglia	59	down-regulated	(30) (6)
DCN 4C4	Htin et en la disease	40	d	(27)
PSMC1	Huntington's disease	48	down-regulated	(27)
PSMC1	Hypoxia of brain	40	up-regulated	(136)
				(10) (26) (42) (27)
PSMC1	Movement disorder	56	down-regulated	(30) (6)
PSMC1	Nerve injury	34	down-regulated	(106)
PSMC1	Neuropathy	67	down-regulated	(90) (47)
			are the transfer of the transf	(10) (26) (42) (28)
PSMC1	Parkinson's disease	62	down-regulated	(30) (6)
PSMC1	Prion disease	82	down-regulated	(49) (37)
PSMC1	Psychotic disorder	39	down-regulated	(57) (38)
PSMC1	Schizophrenia	40	down-regulated	(57) (38)
PSMC1	Sleep disorder	27	down-regulated	(56)
PTBP2	Amnestic disorder	6	down-regulated	(115)
	Amyotrophic lateral			
PTBP2	sclerosis	10	down-regulated	(10)
DTDD3	A suist dissudes	45		(20)
PTBP2	Anxiety disorder	45	up-regulated	(20)
PTBP2	Autistic disorder	14	up-regulated	(21) (71)
PTBP2	Cerebral palsy	28	up-regulated	(125)
				(117) (26) (27) (28)
PTBP2	Disorder of basal ganglia	51	down-regulated	(109) (6)
PTBP2	Encephalomyelopathy	11	down-regulated	(2) (91)
PTBP2	Epilepsy	23	0.0002 p-value	(137)
PTBP2	Huntington's disease	31	up-regulated	(117) (27)
PTBP2	Meningitis	51	down-regulated	(8) (93) (35) (45)
1 151 2			aowii regulatea	(0) (33) (33) (13)
PTBP2	Mood disorder	56	down-regulated	(85) (77) (126) (138)
PTBP2	Motor neuron disease	22	down-regulated	(10) (36) (62)
11012	Wotor Hearth disease		down regulated	(117) (26) (27) (28)
PTBP2	Movement disorder	48	down-regulated	(109) (6)
1 1012	Wiovernent disorder	40	down regulated	· · · · ·
PTBP2	Nerve injury	47	down-regulated	(12) (87) (88) (105) (107)
				(60) (90) (91) (88)
PTBP2	Neuropathy	26	down-regulated	(128) (47)
PTBP2	Paralytic syndrome	32	up-regulated	(91)
			-	· ' · '
PTBP2	Parkinson's disease	57	down-regulated	(26) (28) (109) (6)

PTBP2	Prion disease	17	down-regulated	(113)
PTBP2	Psychotic disorder	42	up-regulated	(38) (16)
PTBP2	Schizophrenia	42	up-regulated	(38) (16)
PTBP2	Sleep disorder	1	down-regulated	(56)
RP11	Amnestic disorder	30	up-regulated	(92)
RP11	Anxiety disorder	64	down-regulated	(98) (20)
RP11	Autistic disorder	52	up-regulated	(21) (22)
RP11	Cerebrovascular disease	27	down-regulated	(24)
	Developmental mental			
RP11	disorder	68	up-regulated	(22)
				(117) (27) (28) (29)
RP11	Disorder of basal ganglia	70	down-regulated	(109)
RP11	Disorder of brain	49	down-regulated	(139)
RP11	Encephalomyelopathy	39	up-regulated	(83) (91)
RP11	Huntington's disease	82	down-regulated	(117) (27) (29)
RP11	Hypoxia of brain	24	up-regulated	(7)
RP11	Meningitis	81	down-regulated	(100) (8) (53) (45)
RP11	Mental retardation	65	up-regulated	(22)
RP11	Mood disorder	17	up-regulated	(3)
				(117) (27) (28) (29)
RP11	Movement disorder	67	down-regulated	(109)
RP11	Nerve injury	25	up-regulated	(60) (12) (13)
				(60) (90) (46) (91)
RP11	Neuropathy	43	up-regulated	(128) (140)
RP11	Paralytic syndrome	49	up-regulated	(91)
RP11	Parkinson's disease	34	down-regulated	(28) (109)
RP11	Prion disease	48	down-regulated	(49)
RP11	Psychotic disorder	41	up-regulated	(17)
RP11	Schizophrenia	41	up-regulated	(17)
RP11	Sleep disorder	59	down-regulated	(50)
RP11	Spinocerebellar ataxia	44	up-regulated	(83)
RP13	Alzheimer's disease	51	down-regulated	(39) (51) (40) (52)
	Attention deficit			
RP13	hyperactivity disorder	79		(70)
RP13	Autistic disorder	68	down-regulated	(21) (71)
RP13	Cerebrovascular disease	19	down-regulated	(24)
RP13	Dementia	51	down-regulated	(39) (51) (40) (52)
	Developmental mental			
RP13	disorder	99		(141)

RP13	Disorder of basal ganglia	25	up-regulated	(29) (30)
RP13	Encephalitis	55	down-regulated	(82)
RP13	Encephalomyelopathy	24	up-regulated	(91)
RP13	Huntington's disease	27	up-regulated	(29)
RP13	Hypoxia of brain	33	down-regulated	(136)
RP13	Meningitis	71	up-regulated	(100) (33) (34) (45)
RP13	Mental retardation	97		(141)
RP13	Movement disorder	23	up-regulated	(29) (30)
RP13	Nerve injury	24	down-regulated	(88) (61) (13)
RP13	Neuropathy	16	up-regulated	(91) (88)
RP13	Paralytic syndrome	44	up-regulated	(91)
RP13	Parkinson's disease	21	down-regulated	(30)
RP13	Sleep disorder	29	down-regulated	(56)
RP4	Anxiety disorder	25	down-regulated	(20)
RP4	Autistic disorder	25	down-regulated	(22)
RP4	Cerebral palsy	46	down-regulated	(125)
	Developmental mental			
RP4	disorder	32	down-regulated	(22)
RP4	Disorder of basal ganglia	8	down-regulated	(27) (6)
RP4	Encephalitis	33	down-regulated	(43)
RP4	Encephalomyelopathy	16	up-regulated	(114)
RP4	Huntington's disease	9	down-regulated	(27)
RP4	Meningitis	34	down-regulated	(8)
RP4	Mental retardation	29	down-regulated	(22)
RP4	Mood disorder	36	3.1E-5 p-value	(76)
RP4	Motor neuron disease	3	down-regulated	(62)
RP4	Movement disorder	5	down-regulated	(27) (6)
RP4	Nerve injury	31	down-regulated	(87) (88) (61) (89)
RP4	Neuropathy	27	down-regulated	(46) (88) (89)
RP4	Parkinson's disease	4	up-regulated	(6)
RPL35	Alzheimer's disease	2	up-regulated	(39)
RPL35	Amnestic disorder	20	up-regulated	(92)
RPL35	Autistic disorder	30	up-regulated	(21)
RPL35	Cerebrovascular disease	16	up-regulated	(24)
RPL35	Dementia	2	up-regulated	(39)
RPL35	Disorder of basal ganglia	26	up-regulated	(27) (29) (30)
RPL35	Encephalitis	29	down-regulated	(110)
RPL35	Encephalomyelitis	40	down-regulated	(110)
RPL35	Encephalomyelopathy	6	down-regulated	(110)
RPL35	Huntington's disease	35	up-regulated	(27) (29)
RPL35	Hypoxia of brain	10	up-regulated	(7)
RPL35	Meningitis	87	up-regulated	(33) (8) (34) (35) (45)

RPL35	Mood disorder	4	down-regulated	(3)
RPL35	Motor neuron disease	23	up-regulated	(36)
RPL35	Movement disorder	23	up-regulated	(27) (29) (30)
RPL35	Multiple sclerosis	3	up-regulated	(4)
RPL35	Myoneural disorder	27	up-regulated	(132)
RPL35	Nerve injury	26	up-regulated	(11) (12) (88) (127)
				(11) (90) (132) (88)
RPL35	Neuropathy	28	up-regulated	(127) (47)
RPL35	Parkinson's disease	4	down-regulated	(30)
RPL35	Prion disease	15	down-regulated	(37)
RPL35	Psychotic disorder	1	0.0008 p-value	(38)
RPL35	Schizophrenia	1	0.0008 p-value	(38)
RPL35	Sleep disorder	43	down-regulated	(50) (58)
RPL5	Alzheimer's disease	3	down-regulated	(52)
	Amyotrophic lateral			
RPL5	sclerosis	29	down-regulated	(19)
RPL5	Autistic disorder	23	up-regulated	(21) (71)
RPL5	Cerebrovascular disease	6	up-regulated	(24)
RPL5	Dementia	3	down-regulated	(52)
RPL5	Disorder of basal ganglia	33	up-regulated	(26) (27) (28) (29)
DDLE	Diagraphy of busin	12		(22)
RPL5	Disorder of brain	12	up-regulated	(32)
RPL5	Encephalitis	58	down-regulated	(110) (142) (82)
RPL5	Encephalomyelitis	37	down-regulated	(110)
RPL5	Encephalomyelopathy	2	down-regulated	(110)
RPL5	Huntington's disease	40	up-regulated	(27) (29)
RPL5	Hypoxia of brain	1	up-regulated	(7)
חחוד	Moningitis	F2	down rogulated	(100) (33) (8) (93)
RPL5	Meningitis	52	down-regulated	(45)
RPL5	Motor neuron disease	38	down-regulated	(36) (19)
RPL5	Movement disorder	30	up-regulated	(26) (27) (28) (29)
INI LS	Wovement disorder	150	up regulated	(20) (27) (20) (23)
RPL5	Multiple sclerosis	70	2.5E-6 p-value	(102) (103)
RPL5	Myoneural disorder	17	up-regulated	(111)
RPL5	Nerve injury	22	down-regulated	(12) (134) (19) (95)
	, ,			(90) (134) (128)
RPL5	Neuropathy	7	up-regulated	(111)
RPL5	Paralytic syndrome	17	up-regulated	(91)
RPL5	Parkinson's disease	18	up-regulated	(26) (28)
RPL5	Prion disease	13	down-regulated	(37)
RPL5	Psychotic disorder	54	2.2E-6 p-value	(108) (38)

RPL5	Schizophrenia	55	2.2E-6 p-value	(108) (38)
RPL5	Sleep disorder	24	down-regulated	(56) (50)
RRAGB	Alzheimer's disease	22	down-regulated	(39) (5)
RRAGB	Dementia	21	down-regulated	(39) (5)
RRAGB	Disorder of basal ganglia	36	down-regulated	(10) (26) (27) (28) (6)
RRAGB	Disorder of brain	17	up-regulated	(32)
RRAGB	Encephalitis	27	down-regulated	(82)
RRAGB	Encephalomyelopathy	6	down-regulated	(2)
RRAGB	Huntington's disease	19	down-regulated	(10) (27)
RRAGB	Meningitis	11	up-regulated	(8)
RRAGB	Mood disorder	1	up-regulated	(3)
RRAGB	Motor neuron disease	1	up-regulated	(62)
RRAGB	Movement disorder	33	down-regulated	(10) (26) (27) (28) (6)
RRAGB	Multiple sclerosis	9	down-regulated	(10)
				(60) (112) (87) (88)
RRAGB	Nerve injury	48	down-regulated	(106)
RRAGB	Neuropathy	6	down-regulated	(11) (88) (89)
RRAGB	Parkinson's disease	41	down-regulated	(10) (26) (28) (6)
RRAGB	Psychotic disorder	13	down-regulated	(15) (16)
RRAGB	Schizophrenia	13	down-regulated	(15) (16)
RRAGB	Sleep disorder	18	down-regulated	(58)
RYR3	Alzheimer's disease	26	down-regulated	(18) (40)
RYR3	Anxiety disorder	63	up-regulated	(133) (98)
RYR3	Autistic disorder	21	up-regulated	(1)
RYR3	Cerebral palsy	85	up-regulated	(99)
RYR3	Cerebrovascular disease	65	6.5E-6 p-value	(72)
RYR3	Dementia	25	down-regulated	(18) (40)
	Developmental mental			
RYR3	disorder	36	down-regulated	(22)
RYR3	Disorder of basal ganglia	56	up-regulated	(29) (6)
RYR3	Disorder of brain	49	up-regulated	(32)
RYR3	Encephalitis	50	up-regulated	(110)
RYR3	Encephalomyelitis	61	up-regulated	(110)
RYR3	Encephalomyelopathy	34	up-regulated	(2) (110)
RYR3	Epilepsy	60	0.7E-5 p-value	(130)
RYR3	Huntington's disease	68	up-regulated	(29)
RYR3	Meningitis	57	up-regulated	(33) (35) (45)
RYR3	Mental retardation	34	down-regulated	(22)
RYR3	Mood disorder	57	8.3E-6 p-value	(143) (74)
RYR3	Movement disorder	53	up-regulated	(29) (6)

RYR3	Multiple sclerosis	24	up-regulated	(63) (65)
RYR3	Myoneural disorder	46	up-regulated	(132)
	,		1 0	(112) (12) (94) (105)
RYR3	Nerve injury	70	down-regulated	(106) (107)
				(90) (46) (94) (132)
RYR3	Neuropathy	44	down-regulated	(47)
RYR3	Parkinson's disease	10	up-regulated	(6)
RYR3	Prion disease	47	down-regulated	(37)
				(144) (57) (15) (38)
RYR3	Psychotic disorder	57	up-regulated	(145)
				(144) (57) (15) (38)
RYR3	Schizophrenia	58	up-regulated	(145)
RYR3	Sleep disorder	46	up-regulated	(56) (50)
SCAI	Alzheimer's disease	38	down-regulated	(39) (18) (40)
	Amyotrophic lateral			
SCAI	sclerosis	41	up-regulated	(10)
SCAI	Autistic disorder	16	up-regulated	(21)
SCAI	Cerebrovascular disease	14	down-regulated	(41)
SCAI	Dementia	38	down-regulated	(39) (18) (40)
SCAI	Disorder of basal ganglia	77	down-regulated	(28) (29) (6)
SCAI	Huntington's disease	66	down-regulated	(29)
SCAI	Hypoxia of brain	17	down-regulated	(41)
SCAI	Meningitis	54	down-regulated	(8) (35)
SCAI	Mood disorder	26	down-regulated	(85) (3)
SCAI	Motor neuron disease	38	up-regulated	(10)
SCAI	Movement disorder	74	down-regulated	(28) (29) (6)
SCAI	Multiple sclerosis	3	down-regulated	(9)
SCAI	Nerve injury	41	up-regulated	(60) (12) (61)
SCAI	Neuropathy	14	up-regulated	(47)
SCAI	Parkinson's disease	78	down-regulated	(28) (6)
SCAI	Prion disease	43	up-regulated	(49) (113)
SCAI	Psychotic disorder	35	down-regulated	(57) (15) (38)
SCAI	Schizophrenia	35	down-regulated	(57) (15) (38)
SCAI	Sleep disorder	53	up-regulated	(56) (50) (58)
SEMA3A	Alzheimer's disease	1	5.9E-5 p-value	(40)
SEMA3A	Amnestic disorder	1	down-regulated	(59)
SEMA3A	Autistic disorder	1	down-regulated	(71)
	Childhood disorder of		_	
SEMA3A	conduct and emotion	26	up-regulated	(119) (120)
SEMA3A	Dementia	1	5.9E-5 p-value	(40)
SEMA3A	Disorder of basal ganglia	7	down-regulated	(10) (48) (27) (121)

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SEMA3A	Huntington's disease	17	down-regulated	(10) (27) (121)
SEMA3A	Lissencephaly	100		(146)
SEMA3A	Mood disorder	1	0.0003 p-value	(76)
SEMA3A	Motor neuron disease	1	up-regulated	(36)
SEMA3A	Movement disorder	4	down-regulated	(10) (48) (27) (121)
SEMA3A	Multiple sclerosis	1	up-regulated	(65) (78)
SEMA3A	Nerve injury	8	up-regulated	(11) (12) (88)
SEMA3A	Neuropathy	71	down-regulated	(11) (88) (47) (140)
SEMA3A	Parkinson's disease	1	up-regulated	(48)
SEMA3A	Prion disease	45	2.7E-6 p-value	(147)
SEMA3A	Psychotic disorder	26	down-regulated	(38) (66) (86)
SEMA3A	Schizophrenia	26	down-regulated	(38) (66) (86)
SEMA3A	Sleep disorder	30	up-regulated	(56) (50)
SLC20A2	Amnestic disorder	19	up-regulated	(59)
SLC20A2	Autistic disorder	7	up-regulated	(1)
SLC20A2	Disorder of basal ganglia	28	down-regulated	(10) (29) (6)
SLC20A2	Disorder of brain	26	up-regulated	(31) (32)
SLC20A2	Encephalomyelopathy	14	down-regulated	(2)
SLC20A2	Huntington's disease	29	down-regulated	(10) (29)
SLC20A2	Meningitis	8	up-regulated	(8) (53)
SLC20A2	Mood disorder	19	8.5E-5 p-value	(74)
SLC20A2	Motor neuron disease	5	down-regulated	(62)
SLC20A2	Movement disorder	25	down-regulated	(10) (29) (6)
SLC20A2	Multiple sclerosis	50	up-regulated	(9) (4) (65) (78)
				(112) (12) (87) (88)
SLC20A2	Nerve injury	50	up-regulated	(107) (95)
				(148) (90) (88) (47)
SLC20A2	Neuropathy	28	down-regulated	(14)
SLC20A2	Paralytic syndrome	24	down-regulated	(91)
SLC20A2	Parkinson's disease	24	down-regulated	(6)
SLC20A2	Prion disease	40	up-regulated	(49) (37)
SLC20A2	Psychotic disorder	17	up-regulated	(57) (17)
SLC20A2	Schizophrenia	17	up-regulated	(57) (17)
SLC20A2	Sleep disorder	10	down-regulated	(56)
SLC25A14	Alzheimer's disease	27	down-regulated	(39) (18) (116)
SLC25A14	Autistic disorder	1	down-regulated	(1)
SLC25A14	Cerebral palsy	20	down-regulated	(125)
SLC25A14	Dementia	26	down-regulated	(39) (18) (116)
SLC25A14	Disorder of basal ganglia	45	down-regulated	(10) (26) (42) (29) (6)
SLC25A14	Encephalitis	24	up-regulated	(142)

SLC25A14	Encephalomyelopathy	12	up-regulated	(83) (2)
SLC25A14	Huntington's disease	47	down-regulated	(10) (29)
SLC25A14	Meningitis	16	down-regulated	(8)
SLC25A14	Movement disorder	42	down-regulated	(10) (26) (42) (29) (6)
SLC25A14	Multiple sclerosis	2	down-regulated	(63) (78)
				(12) (87) (94) (88)
SLC25A14	Nerve injury	27	down-regulated	(107)
SLC25A14	Neuropathy	18	down-regulated	(94) (88) (128)
SLC25A14	Parkinson's disease	41	down-regulated	(10) (48) (26) (42) (6)
SLC25A14	Prion disease	29	down-regulated	(37)
SLC25A14	Psychotic disorder	25	up-regulated	(15) (17)
SLC25A14	Schizophrenia	25	up-regulated	(15) (17)
SLC25A14	Spinocerebellar ataxia	14	up-regulated	(83)
SMARCAD1	Alzheimer's disease	19	down-regulated	(39) (118)
SMARCAD1	Amnestic disorder	1	up-regulated	(92)
SMARCAD1	Anxiety disorder	28	up-regulated	(149)
SMARCAD1	Autistic disorder	1	down-regulated	(23) (1)
SMARCAD1	Cerebrovascular disease	11	up-regulated	(24)
SMARCAD1	Dementia	18	down-regulated	(39) (118)
SMARCAD1	Disorder of basal ganglia	1	up-regulated	(117) (27)
SMARCAD1	Encephalomyelopathy	1	down-regulated	(83)
SMARCAD1	Huntington's disease	11	up-regulated	(117) (27)
SMARCAD1	Meningitis	39	down-regulated	(100) (33) (8) (34)
SMARCAD1	Mood disorder	13	up-regulated	(149) (126) (3)
SMARCAD1	Movement disorder	1	up-regulated	(117) (27)
				(112) (11) (94) (88)
SMARCAD1	Nerve injury	17	down-regulated	(95)
SMARCAD1	Neuropathy	14	down-regulated	(11) (94) (88) (47)
SMARCAD1	Paralytic syndrome	11	up-regulated	(91)
SMARCAD1	Prion disease	12	down-regulated	(49)
SMARCAD1	Psychotic disorder	1	0.0002 p-value	(38)
SMARCAD1	Schizophrenia	1	0.0002 p-value	(38)
SMARCAD1	Sleep disorder	26	up-regulated	(56) (50)
SMARCAD1	Spinocerebellar ataxia	8	down-regulated	(83)
	Attention deficit			
SNORA42	hyperactivity disorder	90	4.9E-6 p-value	(150)
SNORA42	Encephalomyelopathy	51	up-regulated	(2)
SNORA42	Neuropathy	52	up-regulated	(140)

SNORA66	Autistic disorder	33	down-regulated	(71)
SNORA66	Multiple sclerosis	100	2.5E-6 p-value	(102) (103)
SNORA66	Psychotic disorder	83	2.2E-6 p-value	(108) (38)
SNORA66	Schizophrenia	83	2.2E-6 p-value	(108) (38)
SNTG1	Alzheimer's disease	1	down-regulated	(18)
SNTG1	Cerebrovascular disease	1	down-regulated	(41)
SNTG1	Dementia	1	down-regulated	(18)
	Developmental mental			
SNTG1	disorder	68	down-regulated	(151)
SNTG1	Disorder of basal ganglia	30	down-regulated	(10) (48) (26) (29)
SNTG1	Huntington's disease	38	down-regulated	(10) (29)
SNTG1	Hypoxia of brain	7	down-regulated	(84) (41)
SNTG1	Meningitis	1	up-regulated	(8)
SNTG1	Mental disorder	100	down-regulated	(151)
SNTG1	Movement disorder	27	down-regulated	(10) (48) (26) (29)
SNTG1	Multiple sclerosis	3	up-regulated	(9) (65)
SNTG1	Neuropathy	1	down-regulated	(47)
SNTG1	Parkinson's disease	13	down-regulated	(10) (48) (26)
SNTG1	Sleep disorder	5	down-regulated	(50)
SNX19	Disorder of basal ganglia	49	down-regulated	(26) (27) (29) (6)
SNX19	Encephalomyelopathy	12	down-regulated	(91)
SNX19	Huntington's disease	55	down-regulated	(27) (29)
SNX19	Meningitis	67	up-regulated	(100) (33) (53)
SNX19	Mood disorder	23	down-regulated	(126)
SNX19	Movement disorder	46	down-regulated	(26) (27) (29) (6)
SNX19	Multiple sclerosis	12	down-regulated	(81)
SNX19	Myoneural disorder	44	down-regulated	(111)
SNX19	Nerve injury	32	down-regulated	(11) (12) (127)
				(11) (90) (46) (91)
SNX19	Neuropathy	43	down-regulated	(127) (111)
				(0.1)
SNX19	Paralytic syndrome	33	down-regulated	(91)
SNX19	Parkinson's disease	38	down-regulated	(26) (6)
SNX19	Prion disease	36	up-regulated	(37)
SNX19	Psychotic disorder	82	down-regulated	(57) (79) (38) (80)
SNX19	Schizophrenia	83	down-regulated	(57) (79) (38) (80)
ZINVIZ	Schizophirenia	63	down-regulated	(37) (73) (30) (00)
SNX19	Sleep disorder	51	up-regulated	(56) (50) (58)
SOD3	Alzheimer's disease	1	down-regulated	(39) (40)

SOD3	Anxiety disorder	1	up-regulated	(152)
SOD3	Cerebrovascular disease	1	down-regulated	(24)
SOD3	Dementia	18	up-regulated	(153) (39) (40)
SOD3	Disorder of basal ganglia	1	up-regulated	(10) (29)
SOD3	Disorder of brain	1	down-regulated	(32) (122)
SOD3	Huntington's disease	1	up-regulated	(10) (29)
SOD3	Meningitis	2	down-regulated	(33) (8) (53)
SOD3	Motor neuron disease	1	down-regulated	(36)
SOD3	Movement disorder	1	up-regulated	(10) (29)
				(112) (12) (87) (88)
SOD3	Nerve injury	20	up-regulated	(107) (95)
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SOD3	Neuropathy	20	up-regulated	(88) (47)
SOD3	Prion disease	32	up-regulated	(49) (37) (113)
SOD3	Psychotic disorder	1	up-regulated	(57)
SOD3	Schizophrenia	1	up-regulated	(57)
SOD3	Sleep disorder	1	up-regulated	(50)
SPATA7	Alzheimer's disease	23	down-regulated	(5)
SPATA7	Autistic disorder	39	down-regulated	(1)
SPATA7	Dementia	23	down-regulated	(5)
SPATA7	Disorder of basal ganglia	71	up-regulated	(10) (117) (26) (29)
SPATA7	Disorder of brain	77	up-regulated	(31) (139)
SPATA7	Encephalomyelopathy	36	up-regulated	(2)
SPATA7	Huntington's disease	81	up-regulated	(117) (29)
SPATA7	Meningitis	54	up-regulated	(8) (35)
SPATA7	Mood disorder	30	down-regulated	(3)
SPATA7	Movement disorder	68	up-regulated	(10) (117) (26) (29)
SPATA7	Nerve injury	76	down-regulated	(87) (88) (61) (107)
SPATA7	Neuropathy	61	down-regulated	(88)
SPATA7	Parkinson's disease	50	down-regulated	(10) (26)
SPATA7	Psychotic disorder	75	down-regulated	(15) (38)
SPATA7	Schizophrenia	76	down-regulated	(15) (38)
SPATA7	Sleep disorder	98	down-regulated	(56) (50)
ST18	Alzheimer's disease	63	down-regulated	(39) (118) (5)
ST18	Amnestic disorder	37	up-regulated	(92)
ST18	Dementia	62	down-regulated	(39) (118) (5)
				(26) (27) (28) (29)
ST18	Disorder of basal ganglia	68	up-regulated	(109) (121)
CT10	Disambar of head	60		(22) (420)
ST18	Disorder of brain	69	up-regulated	(32) (139)
ST18	Epilepsy	58	4.8E-5 p-value	(130)
ST18	Huntington's disease	76	up-regulated	(27) (29) (121)
ST18	Mood disorder	35	down-regulated	(144)

				(26) (27) (28) (29)
ST18	Movement disorder	65	up-regulated	(109) (121)
3110	Wovernent disorder	105	up regulated	(103) (121)
ST18	Multiple sclerosis	53	down-regulated	(63) (65)
ST18	Nerve injury	49	up-regulated	(60) (12) (107)
ST18	Neuropathy	46	down-regulated	(90) (47)
ST18	Parkinson's disease	51	up-regulated	(26) (28) (109)
ST18	Prion disease	49	down-regulated	(113)
ST18	Psychotic disorder	48	up-regulated	(66) (17)
ST18	Schizophrenia	48	up-regulated	(66) (17)
ST18	Sleep disorder	36	down-regulated	(56)
STYK1	Alzheimer's disease	52	down-regulated	(5) (40) (52)
STYK1	Dementia	51	down-regulated	(5) (40) (52)
STYK1	Disorder of basal ganglia	49	down-regulated	(10)
STYK1	Huntington's disease	55	down-regulated	(10)
STYK1	Hypoxia of brain	33	up-regulated	(84)
STYK1	Mood disorder	8	0.0003 p-value	(74)
STYK1	Movement disorder	47	down-regulated	(10)
STYK1	Neural tube defect	100	down-regulated	(54)
STYK1	Neuropathy	7	down-regulated	(60)
STYK1	Parkinson's disease	38	down-regulated	(10)
STYK1	Psychotic disorder	41	down-regulated	(15) (38)
STYK1	Schizophrenia	41	down-regulated	(15) (38)
TMEM135	Cerebral palsy	57	up-regulated	(99)
TMEM135	Dementia	24	down-regulated	(39) (18)
TIVILIVITSS	Dementia	24	uowii-regulateu	(117) (26) (28) (29)
TMEM135	Disorder of basal ganglia	43	down-regulated	(109) (6)
TIVILIVITSS	Disorder of basar gangila	1 73	down regulated	(103) (0)
TMEM135	Disorder of brain	44	up-regulated	(31) (32)
TMEM135	Mood disorder	22	down-regulated	(85)
TMEM135	Paralytic syndrome	62	up-regulated	(91)
11112111133	r drarytic syriai ome	02	ap regulated	(10) (26) (28) (109)
TMEM135	Parkinson's disease	47	down-regulated	(6)
TMEM135	Psychotic disorder	54	up-regulated	(57) (66) (16)
TMEM135	Schizophrenia	54	up-regulated	(57) (66) (16)
TRPS1	Alzheimer's disease	19	up-regulated	(39) (18) (5) (40)
				()(-)(-)
TRPS1	Autistic disorder	1	up-regulated	(21)
TRPS1	Cerebrovascular disease	23	5.0E-5 p-value	(72)
TRPS1	Dementia	18	up-regulated	(39) (18) (5) (40)
				(117) (26) (28) (29)
TRPS1	Disorder of basal ganglia	57	up-regulated	(109) (6)
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TRPS1	Encephalomyelopathy	1	down-regulated	(83)

				(10) (117) (27) (29)
TRPS1	Huntington's disease	66	up-regulated	(121)
TRPS1	Hypoxia of brain	14	up-regulated	(84) (7)
TRPS1	Meningitis	51	up-regulated	(8) (35) (45)
TRPS1	Mood disorder	1	0.0004 p-value	(74)
TRPS1	Motor neuron disease	13	down-regulated	(62)
				(117) (26) (28) (29)
TRPS1	Movement disorder	54	up-regulated	(109) (6)
TRPS1	Multiple sclerosis	27	up-regulated	(10) (64) (78)
TRPS1	Nerve injury	27	up-regulated	(112) (11) (95) (89)
				(11) (90) (128) (47)
TRPS1	Neuropathy	29	up-regulated	(14) (89)
TRPS1	Parkinson's disease	36	up-regulated	(26) (28) (109) (6)
TRPS1	Psychotic disorder	18	up-regulated	(15) (38) (66)
TRPS1	Schizophrenia	18	up-regulated	(15) (38) (66)
TRPS1	Sleep disorder	15	down-regulated	(50)
TRPS1	Spinocerebellar ataxia	12	down-regulated	(83)
VANGL1	Autistic disorder	1	down-regulated	(21) (22)
VANGL1	Disorder of basal ganglia	1	up-regulated	(29)
VANGL1	Epilepsy	11	down-regulated	(154)
VANGL1	Huntington's disease	1	up-regulated	(29)
VANGL1	Meningitis	1	up-regulated	(33) (8)
VANGL1	Mood disorder	1	down-regulated	(76) (3)
VANGL1	Neural tube defect	100		(146)
VANGL1	Psychotic disorder	1	down-regulated	(66)
VANGL1	Schizophrenia	1	down-regulated	(66)
VDAC3	Anxiety disorder	27	up-regulated	(98) (20)
VDAC3	Autistic disorder	18	up-regulated	(21) (1)
VDAC3	Dementia	20	down-regulated	(18) (51)
				(10) (26) (42) (28)
VDAC3	Disorder of basal ganglia	48	down-regulated	(29) (6)
VDAC3	Encephalomyelopathy	50	down-regulated	(2) (114)
VDAC3	Meningitis	65	up-regulated	(33) (8) (34) (53)
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VDAC3	Myoneural disorder	56	up-regulated	(111)
VDAC3	Parkinson's disease	53	down-regulated	(10) (26) (42) (28) (6)
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WDR38	Disorder of basal ganglia	41	up-regulated	(29)
WDR38	Huntington's disease	54	up-regulated	(29)
WDR38	Meningitis	38	up-regulated	(8)

	Movement disorder	38	up-regulated	(29)
WDR38	Multiple sclerosis	40	up-regulated	(78)
				(11) (12) (87) (107)
WDR38	Nerve injury	75	up-regulated	(89)
WDR38	Neuropathy	64	up-regulated	(11) (89)
WDR38	Psychotic disorder	54	down-regulated	(38) (66)
WDR38	Schizophrenia	54	down-regulated	(38) (66)
ZC3H14	Alzheimer's disease	9	up-regulated	(18)
	Amyotrophic lateral			
ZC3H14	sclerosis	33	down-regulated	(10)
ZC3H14	Anxiety disorder	43	up-regulated	(98) (20)
ZC3H14	Autistic disorder	16	up-regulated	(21)
ZC3H14	Cerebrovascular disease	29	up-regulated	(24)
ZC3H14	Dementia	8	up-regulated	(18)
				(48) (26) (27) (29)
ZC3H14	Disorder of basal ganglia	59	up-regulated	(109)
ZC3H14	Disorder of brain	16	down-regulated	(32)
ZC3H14	Encephalitis	41	down-regulated	(110)
ZC3H14	Encephalomyelitis	52	down-regulated	(110)
ZC3H14	Encephalomyelopathy	18	down-regulated	(110)
ZC3H14	Huntington's disease	63	up-regulated	(27) (29)
ZC3H14	Meningitis	51	down-regulated	(8) (34)
ZC3H14	Mood disorder	25	down-regulated	(85) (3)
ZC3H14	Motor neuron disease	30	down-regulated	(10)
				(48) (26) (27) (29)
ZC3H14	Movement disorder	56	up-regulated	(109)
ZC3H14	Multiple sclerosis	57	down-regulated	(81) (78)
ZC3H14	Myoneural disorder	49	up-regulated	(111)
ZC3H14	Nerve injury	24	down-regulated	(11) (107) (89)
ZC3H14	Neuropathy	32	down-regulated	(11) (111) (89)
ZC3H14	Paralytic syndrome	41	up-regulated	(91)
ZC3H14	Parkinson's disease	53	up-regulated	(48) (26) (109)
ZC3H14	Prion disease	43	up-regulated	(37) (113)
ZC3H14	Psychotic disorder	37	down-regulated	(57) (38)
ZC3H14	Schizophrenia	38	down-regulated	(57) (38)
ZC3H14	Sleep disorder	68	down-regulated	(56) (50) (58)
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## References

- 1. Ginsberg MR, Rubin RA, Falcone T, Ting AH, Natowicz MR. Brain transcriptional and epigenetic associations with autism. PLoS ONE. 2012;7(9):e44736.
- 2. S N. An induced pluripotent stem cell model for ataxia telangiectasia. NCBI GEO. 2012.
- 3. Belzeaux R, Bergon A, Jeanjean V, Loriod B, Formisano-Treziny C, Verrier L, et al. Responder and nonresponder patients exhibit different peripheral transcriptional signatures during major depressive episode. Transl Psychiatry. 2012;2:e185.
- 4. Gandhi KS, McKay FC, Cox M, Riveros C, Armstrong N, Heard RN, et al. The multiple sclerosis whole blood mRNA transcriptome and genetic associations indicate dysregulation of specific T cell pathways in pathogenesis. Hum Mol Genet. 2010;19(11):2134-43.
- 5. Webster JA, Gibbs JR, Clarke J, Ray M, Zhang W, Holmans P, et al. Genetic control of human brain transcript expression in Alzheimer disease. Am J Hum Genet. 2009;84(4):445-58.
- 6. Zheng B, Liao Z, Locascio JJ, Lesniak KA, Roderick SS, Watt ML, et al. PGC- $1\alpha$ , a potential therapeutic target for early intervention in Parkinson's disease. Sci Transl Med. 2010;2(52):52ra73.
- 7. Zamanian JL, Xu L, Foo LC, Nouri N, Zhou L, Giffard RG, et al. Genomic analysis of reactive astrogliosis. J Neurosci. 2012;32(18):6391-410.
- 8. Djavani MM, Crasta OR, Zapata JC, Fei Z, Folkerts O, Sobral B, et al. Early blood profiles of virus infection in a monkey model for Lassa fever. J Virol. 2007;81(15):7960-73.
- 9. A K, P L, A W, M S, E M, F W, et al. miRNA Profiling from total blood of Multiple Sclerosis and control samples. NCBI GEO. 2009.
- 10. PF D, S F, SN K, TP B, I F, D S, et al. Common neuroinflammatory pathways in neurodegenerative diseases. NCBI GEO. 2011.
- 11. B Y, X G. Gene expression signatures following nerve injury in a rat model (0d-14d). NCBI GEO. 2011.
- 12. A F. Spinal Cord Injury Murine Model. NCBI GEO. 2006.
- 13. Geeven G, Macgillavry HD, Eggers R, Sassen MM, Verhaagen J, Smit AB, et al. LLM3D: a log-linear modeling-based method to predict functional gene regulatory interactions from genome-wide expression data. Nucleic Acids Res. 2011;39(13):5313-27.
- 14. Howell GR, Soto I, Zhu X, Ryan M, Macalinao DG, Sousa GL, et al. Radiation treatment inhibits monocyte entry into the optic nerve head and prevents neuronal damage in a mouse model of glaucoma. J Clin Invest. 2012;122(4):1246-61.
- 15. Maycox PR, Kelly F, Taylor A, Bates S, Reid J, Logendra R, et al. Analysis of gene expression in two large schizophrenia cohorts identifies multiple changes associated with nerve terminal function. Mol Psychiatry. 2009;14(12):1083-94.
- 16. de Jong S, Boks MP, Fuller TF, Strengman E, Janson E, de Kovel CG, et al. A gene co-expression network in whole blood of schizophrenia patients is

- independent of antipsychotic-use and enriched for brain-expressed genes. PLoS ONE. 2012;7(6):e39498.
- 17. Horvath S, Zhang Y, Langfelder P, Kahn RS, Boks MP, van Eijk K, et al. Aging effects on DNA methylation modules in human brain and blood tissue. Genome Biol. 2012;13(10):R97.
- 18. Blalock EM, Geddes JW, Chen KC, Porter NM, Markesbery WR, Landfield PW. Incipient Alzheimer's disease: microarray correlation analyses reveal major transcriptional and tumor suppressor responses. Proc Natl Acad Sci USA. 2004;101(7):2173-8.
- 19. Ticozzi N, LeClerc AL, Keagle PJ, Glass JD, Wills AM, van Blitterswijk M, et al. Paraoxonase gene mutations in amyotrophic lateral sclerosis. Ann Neurol. 2010;68(1):102-7.
- 20. Uddin M, Aiello AE, Wildman DE, Koenen KC, Pawelec G, de Los Santos R, et al. Epigenetic and immune function profiles associated with posttraumatic stress disorder. Proc Natl Acad Sci USA. 2010;107(20):9470-5.
- 21. S K, C C, IA H, LM K, IS K. Blood gene expression signatures distinguish autism spectrum disorders from controls. NCBI GEO. 2009.
- 22. Gregg JP, Lit L, Baron CA, Hertz-Picciotto I, Walker W, Davis RA, et al. Gene expression changes in children with autism. Genomics. 2008;91(1):22-9.
- 23. Chow ML, Winn ME, Li HR, April C, Wynshaw-Boris A, Fan JB, et al. Preprocessing and Quality Control Strategies for Illumina DASL Assay-Based Brain Gene Expression Studies with Semi-Degraded Samples. Front Genet. 2012;3:11.
- 24. Tseveleki V, Rubio R, Vamvakas SS, White J, Taoufik E, Petit E, et al. Comparative gene expression analysis in mouse models for multiple sclerosis, Alzheimer's disease and stroke for identifying commonly regulated and disease-specific gene changes. Genomics. 2010;96(2):82-91.
- 25. Gow JW, Hagan S, Herzyk P, Cannon C, Behan PO, Chaudhuri A. A gene signature for post-infectious chronic fatigue syndrome. BMC Med Genomics. 2009;2:38.
- 26. R R. Human body index transcriptional profiling. NCBI GEO. 2007.
- 27. Borovecki F, Lovrecic L, Zhou J, Jeong H, Then F, Rosas HD, et al. Genomewide expression profiling of human blood reveals biomarkers for Huntington's disease. Proc Natl Acad Sci USA. 2005;102(31):11023-8.
- 28. Moran LB, Duke DC, Deprez M, Dexter DT, Pearce RK, Graeber MB. Whole genome expression profiling of the medial and lateral substantia nigra in Parkinson's disease. Neurogenetics. 2006;7(1):1-11.
- 29. Hodges A, Strand AD, Aragaki AK, Kuhn A, Sengstag T, Hughes G, et al. Regional and cellular gene expression changes in human Huntington's disease brain. Hum Mol Genet. 2006;15(6):965-77.
- 30. Lewandowski NM, Ju S, Verbitsky M, Ross B, Geddie ML, Rockenstein E, et al. Polyamine pathway contributes to the pathogenesis of Parkinson disease. Proc Natl Acad Sci USA. 2010;107(39):16970-5.
- 31. JM S, LA F. Effect of brain death on gene expression in liver from rhesus macaque. NCBI GEO. 2010.
- 32. Chen-Plotkin AS, Geser F, Plotkin JB, Clark CM, Kwong LK, Yuan W, et al. Variations in the progranulin gene affect global gene expression in frontotemporal lobar degeneration. Hum Mol Genet. 2008;17(10):1349-62.

- 33. Pathan N, Hemingway CA, Alizadeh AA, Stephens AC, Boldrick JC, Oragui EE, et al. Role of interleukin 6 in myocardial dysfunction of meningococcal septic shock. Lancet. 2004;363(9404):203-9.
- 34. Djavani M, Crasta OR, Zhang Y, Zapata JC, Sobral B, Lechner MG, et al. Gene expression in primate liver during viral hemorrhagic fever. Virol J. 2009;6:20.
- 35. Johnston RJ, Poholek AC, DiToro D, Yusuf I, Eto D, Barnett B, et al. Bcl6 and Blimp-1 are reciprocal and antagonistic regulators of T follicular helper cell differentiation. Science. 2009;325(5943):1006-10.
- 36. Corti S, Locatelli F, Papadimitriou D, Donadoni C, Del Bo R, Crimi M, et al. Transplanted ALDHhiSSClo neural stem cells generate motor neurons and delay disease progression of nmd mice, an animal model of SMARD1. Hum Mol Genet. 2006;15(2):167-87.
- 37. Xiang W, Hummel M, Mitteregger G, Pace C, Windl O, Mansmann U, et al. Transcriptome analysis reveals altered cholesterol metabolism during the neurodegeneration in mouse scrapie model. J Neurochem. 2007;102(3):834-47.
- 38. Shi J, Levinson DF, Duan J, Sanders AR, Zheng Y, Pe'er I, et al. Common variants on chromosome 6p22.1 are associated with schizophrenia. Nature. 2009;460(7256):753-7.
- 39. H S, A M, J K, H A, S T, H N. The use of non-amplified RNA samples for gene expression analysis by sensitive microarray. NCBI GEO. 2011.
- 40. Harold D, Abraham R, Hollingworth P, Sims R, Gerrish A, Hamshere ML, et al. Genome-wide association study identifies variants at CLU and PICALM associated with Alzheimer's disease. Nat Genet. 2009;41(10):1088-93.
- 41. Stevens SL, Leung PY, Vartanian KB, Gopalan B, Yang T, Simon RP, et al. Multiple preconditioning paradigms converge on interferon regulatory factor-dependent signaling to promote tolerance to ischemic brain injury. J Neurosci. 2011;31(23):8456-63.
- 42. Zhang Y, James M, Middleton FA, Davis RL. Transcriptional analysis of multiple brain regions in Parkinson's disease supports the involvement of specific protein processing, energy metabolism, and signaling pathways, and suggests novel disease mechanisms. Am J Med Genet B Neuropsychiatr Genet. 2005;137B(1):5-16.
- 43. Gersten M, Alirezaei M, Marcondes MC, Flynn C, Ravasi T, Ideker T, et al. An integrated systems analysis implicates EGR1 downregulation in simian immunodeficiency virus encephalitis-induced neural dysfunction. J Neurosci. 2009;29(40):12467-76.
- 44. Zhou T, Chou J, Zhou Y, Simpson DA, Cao F, Bushel PR, et al. Ataxia telangiectasia-mutated dependent DNA damage checkpoint functions regulate gene expression in human fibroblasts. Mol Cancer Res. 2007;5(8):813-22.
- 45. Marshall HD, Chandele A, Jung YW, Meng H, Poholek AC, Parish IA, et al. Differential expression of Ly6C and T-bet distinguish effector and memory Th1 CD4(+) cell properties during viral infection. Immunity. 2011;35(4):633-46.
- 46. Hernandez MR, Agapova OA, Yang P, Salvador-Silva M, Ricard CS, Aoi S. Differential gene expression in astrocytes from human normal and glaucomatous optic nerve head analyzed by cDNA microarray. Glia. 2002;38(1):45-64.
- 47. Howell GR, Macalinao DG, Sousa GL, Walden M, Soto I, Kneeland SC, et al. Molecular clustering identifies complement and endothelin induction as early events in a mouse model of glaucoma. J Clin Invest. 2011;121(4):1429-44.

- 48. R K, der Brug M v, J V, J D, S S, A R, et al. Gene expression changes across multiple regions of the Parkinson's disease brain. NCBI GEO. 2011.
- 49. G M, A A. Preclinical CNS changes following peripheral murine RML prion challenge. NCBI GEO. 2007.
- 50. Maret S, Dorsaz S, Gurcel L, Pradervand S, Petit B, Pfister C, et al. Homer1a is a core brain molecular correlate of sleep loss. Proc Natl Acad Sci USA. 2007;104(50):20090-5.
- 51. Williams C, Mehrian Shai R, Wu Y, Hsu YH, Sitzer T, Spann B, et al. Transcriptome analysis of synaptoneurosomes identifies neuroplasticity genes overexpressed in incipient Alzheimer's disease. PLoS ONE. 2009;4(3):e4936.
- 52. Nunez-Iglesias J, Liu CC, Morgan TE, Finch CE, Zhou XJ. Joint genome-wide profiling of miRNA and mRNA expression in Alzheimer's disease cortex reveals altered miRNA regulation. PLoS ONE. 2010;5(2):e8898.
- 53. Kalia V, Sarkar S, Subramaniam S, Haining WN, Smith KA, Ahmed R. Prolonged interleukin-2Ralpha expression on virus-specific CD8+ T cells favors terminal-effector differentiation in vivo. Immunity. 2010;32(1):91-103.
- 54. Nagy GR, Gyorffy B, Galamb O, Molnar B, Nagy B, Papp Z. Use of routinely collected amniotic fluid for whole-genome expression analysis of polygenic disorders. Clin Chem. 2006;52(11):2013-20.
- 55. Steele MR, Inman DM, Calkins DJ, Horner PJ, Vetter ML. Microarray analysis of retinal gene expression in the DBA/2J model of glaucoma. Invest Ophthalmol Vis Sci. 2006;47(3):977-85.
- 56. Mackiewicz M, Shockley KR, Romer MA, Galante RJ, Zimmerman JE, Naidoo N, et al. Macromolecule biosynthesis: a key function of sleep. Physiol Genomics. 2007;31(3):441-57.
- 57. Narayan S, Tang B, Head SR, Gilmartin TJ, Sutcliffe JG, Dean B, et al. Molecular profiles of schizophrenia in the CNS at different stages of illness. Brain Res. 2008;1239:235-48.
- 58. Thompson CL, Wisor JP, Lee CK, Pathak SD, Gerashchenko D, Smith KA, et al. Molecular and anatomical signatures of sleep deprivation in the mouse brain. Front Neurosci. 2010;4:165.
- 59. Pawlowski TL, Bellush LL, Wright AW, Walker JP, Colvin RA, Huentelman MJ. Hippocampal gene expression changes during age-related cognitive decline. Brain Res. 2009;1256:101-10.
- 60. C H. Entrapment Neuropathy Results in Different MicroRNAs Expression Patterns from Denervation Injury in Rats. NCBI GEO. 2010.
- 61. Templeton JP, Nassr M, Vazquez-Chona F, Freeman-Anderson NE, Orr WE, Williams RW, et al. Differential response of C57BL/6J mouse and DBA/2J mouse to optic nerve crush. BMC Neurosci. 2009;10:90.
- 62. Ebert AD, Yu J, Rose FF, Mattis VB, Lorson CL, Thomson JA, et al. Induced pluripotent stem cells from a spinal muscular atrophy patient. Nature. 2009;457(7227):277-80.
- 63. Singh MK, Scott TF, LaFramboise WA, Hu FZ, Post JC, Ehrlich GD. Gene expression changes in peripheral blood mononuclear cells from multiple sclerosis patients undergoing beta-interferon therapy. J Neurol Sci. 2007;258(1-2):52-9.
- 64. Baranzini SE, Wang J, Gibson RA, Galwey N, Naegelin Y, Barkhof F, et al. Genome-wide association analysis of susceptibility and clinical phenotype in multiple sclerosis. Hum Mol Genet. 2009;18(4):767-78.

- 65. Keller A, Leidinger P, Bauer A, Elsharawy A, Haas J, Backes C, et al. Toward the blood-borne miRNome of human diseases. Nat Methods. 2011;8(10):841-3.
- 66. Brennand KJ, Simone A, Jou J, Gelboin-Burkhart C, Tran N, Sangar S, et al. Modelling schizophrenia using human induced pluripotent stem cells. Nature. 2011;473(7346):221-5.
- 67. Lee JH, Cheng R, Barral S, Reitz C, Medrano M, Lantigua R, et al. Identification of novel loci for Alzheimer disease and replication of CLU, PICALM, and BIN1 in Caribbean Hispanic individuals. Arch Neurol. 2011;68(3):320-8.
- 68. Naj AC, Jun G, Beecham GW, Wang LS, Vardarajan BN, Buros J, et al. Common variants at MS4A4/MS4A6E, CD2AP, CD33 and EPHA1 are associated with late-onset Alzheimer's disease. Nat Genet. 2011;43(5):436-41.
- 69. Hollingworth P, Sweet R, Sims R, Harold D, Russo G, Abraham R, et al. Genome-wide association study of Alzheimer's disease with psychotic symptoms. Mol Psychiatry. 2012;17(12):1316-27.
- 70. Lionel AC, Crosbie J, Barbosa N, Goodale T, Thiruvahindrapuram B, Rickaby J, et al. Rare copy number variation discovery and cross-disorder comparisons identify risk genes for ADHD. Sci Transl Med. 2011;3(95):95ra75.
- 71. Sakai Y, Shaw CA, Dawson BC, Dugas DV, Al-Mohtaseb Z, Hill DE, et al. Protein interactome reveals converging molecular pathways among autism disorders. Sci Transl Med. 2011;3(86):86ra49.
- 72. Ikram MA, Seshadri S, Bis JC, Fornage M, DeStefano AL, Aulchenko YS, et al. Genomewide association studies of stroke. N Engl J Med. 2009;360(17):1718-28.
- 73. Simon-Sanchez J, Schulte C, Bras JM, Sharma M, Gibbs JR, Berg D, et al. Genome-wide association study reveals genetic risk underlying Parkinson's disease. Nat Genet. 2009;41(12):1308-12.
- 74. Shyn SI, Shi J, Kraft JB, Potash JB, Knowles JA, Weissman MM, et al. Novel loci for major depression identified by genome-wide association study of Sequenced Treatment Alternatives to Relieve Depression and meta-analysis of three studies. Mol Psychiatry. 2011;16(2):202-15.
- 75. Shi J, Potash JB, Knowles JA, Weissman MM, Coryell W, Scheftner WA, et al. Genome-wide association study of recurrent early-onset major depressive disorder. Mol Psychiatry. 2011;16(2):193-201.
- 76. Terracciano A, Tanaka T, Sutin AR, Sanna S, Deiana B, Lai S, et al. Genomewide association scan of trait depression. Biol Psychiatry. 2010;68(9):811-7.
- 77. Wray NR, Pergadia ML, Blackwood DH, Penninx BW, Gordon SD, Nyholt DR, et al. Genome-wide association study of major depressive disorder: new results, meta-analysis, and lessons learned. Mol Psychiatry. 2012;17(1):36-48.
- 78. Han MH, Lundgren DH, Jaiswal S, Chao M, Graham KL, Garris CS, et al. Janus-like opposing roles of CD47 in autoimmune brain inflammation in humans and mice. J Exp Med. 2012;209(7):1325-34.
- 79. Need AC, Ge D, Weale ME, Maia J, Feng S, Heinzen EL, et al. A genome-wide investigation of SNPs and CNVs in schizophrenia. PLoS Genet. 2009;5(2):e1000373.
- 80. Ripke S, Sanders AR, Kendler KS, Levinson DF, Sklar P, Holmans PA, et al. Genome-wide association study identifies five new schizophrenia loci. Nat Genet. 2011;43(10):969-76.

- 81. Gilli F, Lindberg RL, Valentino P, Marnetto F, Malucchi S, Sala A, et al. Learning from nature: pregnancy changes the expression of inflammation-related genes in patients with multiple sclerosis. PLoS ONE. 2010;5(1):e8962.
- 82. Haque A, Best SE, Ammerdorffer A, Desbarrieres L, de Oca MM, Amante FH, et al. Type I interferons suppress CD4<sup>+</sup> T-cell-dependent parasite control during blood-stage Plasmodium infection. Eur J Immunol. 2011;41(9):2688-98.
- 83. G C, R B, S P, R V, F G, H P, et al. A Gene Expression Phenotype In Lymphocytes From Friedreich's Ataxia Patients. NCBI GEO. 2011.
- 84. Yilmaz G, Alexander JS, Erkuran Yilmaz C, Granger DN. Induction of neuroprotective/regenerative genes in stem cells infiltrating post-ischemic brain tissue. Exp Transl Stroke Med. 2010;2(1):11.
- 85. Ryan MM, Lockstone HE, Huffaker SJ, Wayland MT, Webster MJ, Bahn S. Gene expression analysis of bipolar disorder reveals downregulation of the ubiquitin cycle and alterations in synaptic genes. Mol Psychiatry. 2006;11(10):965-78.
- 86. Chen C, Cheng L, Grennan K, Pibiri F, Zhang C, Badner JA, et al. Two gene co-expression modules differentiate psychotics and controls. Mol Psychiatry. 2012.
- 87. Di Giovanni S, Knoblach SM, Brandoli C, Aden SA, Hoffman EP, Faden AI. Gene profiling in spinal cord injury shows role of cell cycle in neuronal death. Ann Neurol. 2003;53(4):454-68.
- 88. Agudo M, Perez-Marin MC, Lonngren U, Sobrado P, Conesa A, Canovas I, et al. Time course profiling of the retinal transcriptome after optic nerve transection and optic nerve crush. Mol Vis. 2008;14:1050-63.
- 89. Wang Y, Tang X, Yu B, Gu Y, Yuan Y, Yao D, et al. Gene network revealed involvements of Birc2, Birc3 and Tnfrsf1a in anti-apoptosis of injured peripheral nerves. PLoS ONE. 2012;7(9):e43436.
- 90. MG K, DC C, RR DJ, AA TA, RH K, L K, et al. Down-stream effects in demyelinating neuropathies. NCBI GEO. 2007.
- 91. Tezak Z, Hoffman EP, Lutz JL, Fedczyna TO, Stephan D, Bremer EG, et al. Gene expression profiling in DQA1\*0501+ children with untreated dermatomyositis: a novel model of pathogenesis. J Immunol. 2002;168(8):4154-63.
- 92. CA VK, HD V, GV B, CG I, VP K, JM S, et al. Induction of the neuronal major histocompatibility complex class I and paired immunoglobulin-like receptor B pathway in the hippocampus with aging and cognitive decline. NCBI GEO. 2011.
- 93. Wherry EJ, Ha SJ, Kaech SM, Haining WN, Sarkar S, Kalia V, et al. Molecular signature of CD8+ T cell exhaustion during chronic viral infection. Immunity. 2007;27(4):670-84.
- 94. Shetty RS, Bose SC, Nickell MD, McIntyre JC, Hardin DH, Harris AM, et al. Transcriptional changes during neuronal death and replacement in the olfactory epithelium. Mol Cell Neurosci. 2005;30(1):90-107.
- 95. Barrette B, Calvo E, Vallieres N, Lacroix S. Transcriptional profiling of the injured sciatic nerve of mice carrying the Wld(S) mutant gene: identification of genes involved in neuroprotection, neuroinflammation, and nerve regeneration. Brain Behav Immun. 2010;24(8):1254-67.
- 96. Chen J, Lee G, Fanous AH, Zhao Z, Jia P, O'Neill A, et al. Two non-synonymous markers in PTPN21, identified by genome-wide association study

- data-mining and replication, are associated with schizophrenia. Schizophr Res. 2011;131(1-3):43-51.
- 97. Haberman RP, Colantuoni C, Stocker AM, Schmidt AC, Pedersen JT, Gallagher M. Prominent hippocampal CA3 gene expression profile in neurocognitive aging. Neurobiol Aging. 2011;32(9):1678-92.
- 98. Aerssens J, Hillsley K, Peeters PJ, de Hoogt R, Stanisz A, Lin JH, et al. Alterations in the brain-gut axis underlying visceral chemosensitivity in Nippostrongylus brasiliensis-infected mice. Gastroenterology. 2007;132(4):1375-87.
- 99. Smith LR, Chambers HG, Subramaniam S, Lieber RL. Transcriptional abnormalities of hamstring muscle contractures in children with cerebral palsy. PLoS ONE. 2012;7(8):e40686.
- 100. Pereira C. E-MEXP-1035 Transcription profiling of macrophages from healthy human individuals and those with Whipple disease, endocarditis, meningitis or pneumococus infection. ArrayExpress. 2008.
- 101. Savitz J, Frank MB, Victor T, Bebak M, Marino JH, Bellgowan PS, et al. Inflammation and neurological disease-related genes are differentially expressed in depressed patients with mood disorders and correlate with morphometric and functional imaging abnormalities. Brain Behav Immun. 2013;31:161-71.
- 102. Hafler DA, Compston A, Sawcer S, Lander ES, Daly MJ, De Jager PL, et al. Risk alleles for multiple sclerosis identified by a genomewide study. N Engl J Med. 2007;357(9):851-62.
- 103. Bahlo M, Booth DR, Broadley SA, Brown MA, Foote SJ, Griffiths LR, et al. Genome-wide association study identifies new multiple sclerosis susceptibility loci on chromosomes 12 and 20. Nat Genet. 2009;41(7):824-8.
- 104. Sawcer S, Hellenthal G, Pirinen M, Spencer CC, Patsopoulos NA, Moutsianas L, et al. Genetic risk and a primary role for cell-mediated immune mechanisms in multiple sclerosis. Nature. 2011;476(7359):214-9.
- 105. Wienecke J, Westerdahl AC, Hultborn H, Kiehn O, Ryge J. Global gene expression analysis of rodent motor neurons following spinal cord injury associates molecular mechanisms with development of postinjury spasticity. J Neurophysiol. 2010;103(2):761-78.
- 106. Ryge J, Winther O, Wienecke J, Sandelin A, Westerdahl AC, Hultborn H, et al. Transcriptional regulation of gene expression clusters in motor neurons following spinal cord injury. BMC Genomics. 2010;11:365.
- 107. Michaelevski I, Segal-Ruder Y, Rozenbaum M, Medzihradszky KF, Shalem O, Coppola G, et al. Signaling to transcription networks in the neuronal retrograde injury response. Sci Signal. 2010;3(130):ra53.
- 108. O'Donovan MC, Craddock N, Norton N, Williams H, Peirce T, Moskvina V, et al. Identification of loci associated with schizophrenia by genome-wide association and follow-up. Nat Genet. 2008;40(9):1053-5.
- 109. Lesnick TG, Papapetropoulos S, Mash DC, Ffrench-Mullen J, Shehadeh L, de Andrade M, et al. A genomic pathway approach to a complex disease: axon guidance and Parkinson disease. PLoS Genet. 2007;3(6):e98.
- 110. Otaegui D, Mostafavi S, Bernard CC, Lopez de Munain A, Mousavi P, Oksenberg JR, et al. Increased transcriptional activity of milk-related genes following the active phase of experimental autoimmune encephalomyelitis and multiple sclerosis. J Immunol. 2007;179(6):4074-82.

- 111. Eisenberg I, Novershtern N, Itzhaki Z, Becker-Cohen M, Sadeh M, Willems PH, et al. Mitochondrial processes are impaired in hereditary inclusion body myopathy. Hum Mol Genet. 2008;17(23):3663-74.
- 112. X L. Patterns of gene Expression during the Spinal cord Injury/Regeneration. NCBI GEO. 2011.
- 113. Lunnon K, Teeling JL, Tutt AL, Cragg MS, Glennie MJ, Perry VH. Systemic inflammation modulates Fc receptor expression on microglia during chronic neurodegeneration. J Immunol. 2011;186(12):7215-24.
- 114. Crimi M, Bordoni A, Menozzi G, Riva L, Fortunato F, Galbiati S, et al. Skeletal muscle gene expression profiling in mitochondrial disorders. FASEB J. 2005;19(7):866-8.
- 115. Burger C, Lopez MC, Feller JA, Baker HV, Muzyczka N, Mandel RJ. Changes in transcription within the CA1 field of the hippocampus are associated with agerelated spatial learning impairments. Neurobiol Learn Mem. 2007;87(1):21-41.
- 116. Naj AC, Beecham GW, Martin ER, Gallins PJ, Powell EH, Konidari I, et al. Dementia revealed: novel chromosome 6 locus for late-onset Alzheimer disease provides genetic evidence for folate-pathway abnormalities. PLoS Genet. 2010;6(9):e1001130.
- 117. A B, G B. Expression data from human Huntington fibroblasts. NCBI GEO. 2013.
- 118. Abraham R, Moskvina V, Sims R, Hollingworth P, Morgan A, Georgieva L, et al. A genome-wide association study for late-onset Alzheimer's disease using DNA pooling. BMC Med Genomics. 2008;1:44.
- 119. HA D, RJ G. Microarray analysis of male flies selected for increased aggressive behavior. NCBI GEO. 2006.
- 120. Dierick HA, Greenspan RJ. Molecular analysis of flies selected for aggressive behavior. Nat Genet. 2006;38(9):1023-31.
- 121. Hodges A, Hughes G, Brooks S, Elliston L, Holmans P, Dunnett SB, et al. Brain gene expression correlates with changes in behavior in the R6/1 mouse model of Huntington's disease. Genes Brain Behav. 2008;7(3):288-99.
- 122. Hoeger S, Bergstraesser C, Selhorst J, Fontana J, Birck R, Waldherr R, et al. Modulation of brain dead induced inflammation by vagus nerve stimulation. Am J Transplant. 2010;10(3):477-89.
- 123. Borges K, Shaw R, Dingledine R. Gene expression changes after seizure preconditioning in the three major hippocampal cell layers. Neurobiol Dis. 2007;26(1):66-77.
- 124. Uriguen L, Arteta D, Diez-Alarcia R, Ferrer-Alcon M, Diaz A, Pazos A, et al. Gene expression patterns in brain cortex of three different animal models of depression. Genes Brain Behav. 2008;7(6):649-58.
- 125. Smith LR, Ponten E, Hedstrom Y, Ward SR, Chambers HG, Subramaniam S, et al. Novel transcriptional profile in wrist muscles from cerebral palsy patients. BMC Med Genomics. 2009;2:44.
- 126. Beech RD, Lowthert L, Leffert JJ, Mason PN, Taylor MM, Umlauf S, et al. Increased peripheral blood expression of electron transport chain genes in bipolar depression. Bipolar Disord. 2010;12(8):813-24.
- 127. Maratou K, Wallace VC, Hasnie FS, Okuse K, Hosseini R, Jina N, et al. Comparison of dorsal root ganglion gene expression in rat models of traumatic and HIV-associated neuropathic pain. Eur J Pain. 2009;13(4):387-98.

- 128. Lukas TJ, Miao H, Chen L, Riordan SM, Li W, Crabb AM, et al. Susceptibility to glaucoma: differential comparison of the astrocyte transcriptome from glaucomatous African American and Caucasian American donors. Genome Biol. 2008;9(7):R111.
- 129. De Jager PL, Shulman JM, Chibnik LB, Keenan BT, Raj T, Wilson RS, et al. A genome-wide scan for common variants affecting the rate of age-related cognitive decline. Neurobiol Aging. 2012;33(5):1-15.
- 130. Kasperavici?te D, Catarino CB, Heinzen EL, Depondt C, Cavalleri GL, Caboclo LO, et al. Common genetic variation and susceptibility to partial epilepsies: a genome-wide association study. Brain. 2010;133(Pt 7):2136-47.
- 131. Baum AE, Akula N, Cabanero M, Cardona I, Corona W, Klemens B, et al. A genome-wide association study implicates diacylglycerol kinase eta (DGKH) and several other genes in the etiology of bipolar disorder. Mol Psychiatry. 2008;13(2):197-207.
- 132. Zhou Y, Gong B, Lin F, Rother RP, Medof ME, Kaminski HJ. Anti-C5 antibody treatment ameliorates weakness in experimentally acquired myasthenia gravis. J Immunol. 2007;179(12):8562-7.
- 133. W O. Ogle-5P01NS037520-05. NCBI GEO. 2005.
- 134. Gonzalez de Aguilar JL, Niederhauser-Wiederkehr C, Halter B, De Tapia M, Di Scala F, Demougin P, et al. Gene profiling of skeletal muscle in an amyotrophic lateral sclerosis mouse model. Physiol Genomics. 2008;32(2):207-18.
- 135. Hinney A, Scherag A, Jarick I, Albayrak O, Putter C, Pechlivanis S, et al. Genome-wide association study in German patients with attention deficit/hyperactivity disorder. Am J Med Genet B Neuropsychiatr Genet. 2011;156B(8):888-97.
- 136. Hillmeister P, Lehmann KE, Bondke A, Witt H, Duelsner A, Gruber C, et al. Induction of cerebral arteriogenesis leads to early-phase expression of protease inhibitors in growing collaterals of the brain. J Cereb Blood Flow Metab. 2008;28(11):1811-23.
- 137. Guo Y, Baum LW, Sham PC, Wong V, Ng PW, Lui CH, et al. Two-stage genome-wide association study identifies variants in CAMSAP1L1 as susceptibility loci for epilepsy in Chinese. Hum Mol Genet. 2012;21(5):1184-9.
- 138. Yosifova A, Mushiroda T, Kubo M, Takahashi A, Kamatani Y, Kamatani N, et al. Genome-wide association study on bipolar disorder in the Bulgarian population. Genes Brain Behav. 2011;10(7):789-97.
- 139. Almeida S, Zhang Z, Coppola G, Mao W, Futai K, Karydas A, et al. Induced pluripotent stem cell models of progranulin-deficient frontotemporal dementia uncover specific reversible neuronal defects. Cell Rep. 2012;2(4):789-98.
- 140. Jiang Q, Ho YY, Hao L, Nichols Berrios C, Chakravarti A. Copy number variants in candidate genes are genetic modifiers of Hirschsprung disease. PLoS ONE. 2011;6(6):e21219.
- 141. Vissers LE, de Ligt J, Gilissen C, Janssen I, Steehouwer M, de Vries P, et al. A de novo paradigm for mental retardation. Nat Genet. 2010;42(12):1109-12.
- 142. Lovegrove FE, Gharib SA, Patel SN, Hawkes CA, Kain KC, Liles WC. Expression microarray analysis implicates apoptosis and interferon-responsive mechanisms in susceptibility to experimental cerebral malaria. Am J Pathol. 2007;171(6):1894-903.

- 143. Muglia P, Tozzi F, Galwey NW, Francks C, Upmanyu R, Kong XQ, et al. Genome-wide association study of recurrent major depressive disorder in two European case-control cohorts. Mol Psychiatry. 2010;15(6):589-601.
- 144. Iwamoto K, Bundo M, Kato T. Altered expression of mitochondria-related genes in postmortem brains of patients with bipolar disorder or schizophrenia, as revealed by large-scale DNA microarray analysis. Hum Mol Genet. 2005;14(2):241-53.
- 145. Athanasiu L, Mattingsdal M, Kahler AK, Brown A, Gustafsson O, Agartz I, et al. Gene variants associated with schizophrenia in a Norwegian genome-wide study are replicated in a large European cohort. J Psychiatr Res. 2010;44(12):748-53.
- 146. Eppig JT, Blake JA, Bult CJ, Kadin JA, Richardson JE. The mouse genome database (MGD): new features facilitating a model system. Nucleic Acids Res. 2007;35(Database issue):D630-7.
- 147. Mead S, Uphill J, Beck J, Poulter M, Campbell T, Lowe J, et al. Genome-wide association study in multiple human prion diseases suggests genetic risk factors additional to PRNP. Hum Mol Genet. 2012;21(8):1897-906.
- 148. SG G, MH K, RH K, MM H, G A, B J, et al. Neuroinflammation in Advanced Canine Glaucoma. NCBI GEO. 2010.
- 149. Krishnan V, Han MH, Graham DL, Berton O, Renthal W, Russo SJ, et al. Molecular adaptations underlying susceptibility and resistance to social defeat in brain reward regions. Cell. 2007;131(2):391-404.
- 150. Sonuga-Barke EJ, Lasky-Su J, Neale BM, Oades R, Chen W, Franke B, et al. Does parental expressed emotion moderate genetic effects in ADHD? An exploration using a genome wide association scan. Am J Med Genet B Neuropsychiatr Genet. 2008;147B(8):1359-68.
- 151. Whitby H, Tsalenko A, Aston E, Tsang P, Mitchell S, Bayrak-Toydemir P, et al. Benign copy number changes in clinical cytogenetic diagnostics by array CGH. Cytogenet Genome Res. 2008;123(1-4):94-101.
- 152. Segman RH, Shefi N, Goltser-Dubner T, Friedman N, Kaminski N, Shalev AY. Peripheral blood mononuclear cell gene expression profiles identify emergent post-traumatic stress disorder among trauma survivors. Mol Psychiatry. 2005;10(5):500-13.
- 153. AF F, Y A, JI M-S, B B, R S, H T, et al. A DNA Methylation Fingerprint of 1,628 Human Samples. NCBI GEO. 2011.
- 154. McRae AF, Matigian NA, Vadlamudi L, Mulley JC, Mowry B, Martin NG, et al. Replicated effects of sex and genotype on gene expression in human lymphoblastoid cell lines. Hum Mol Genet. 2007;16(4):364-73.