Supplemental Data

Neuropsychological Endophenotype Approach to Genome-wide Linkage Analysis Identifies Susceptibility Loci for ADHD on 2q21.1 and 13q12.11

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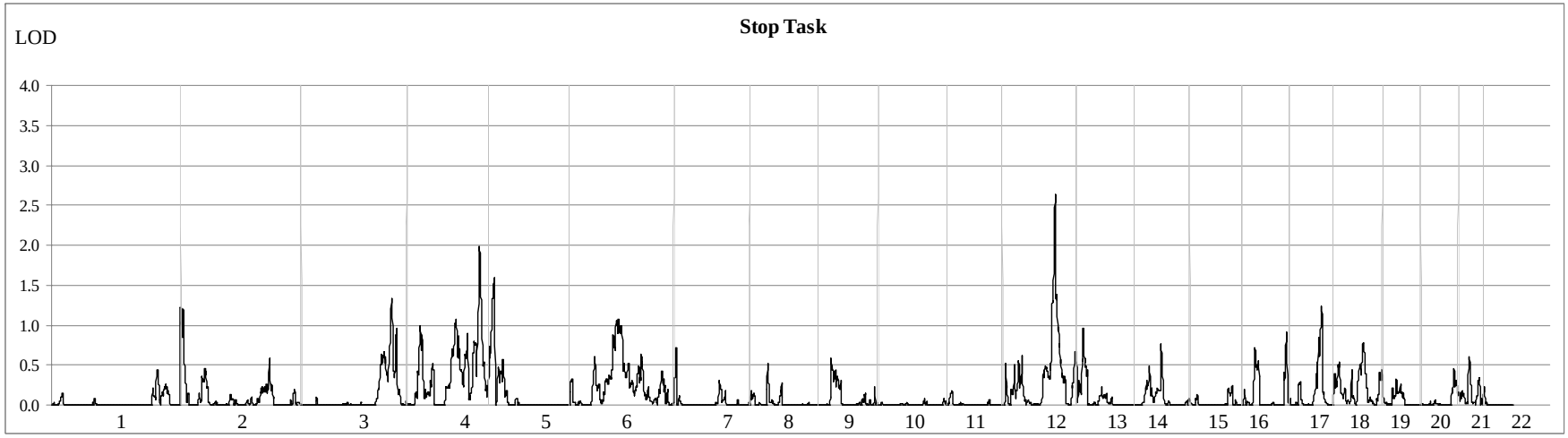
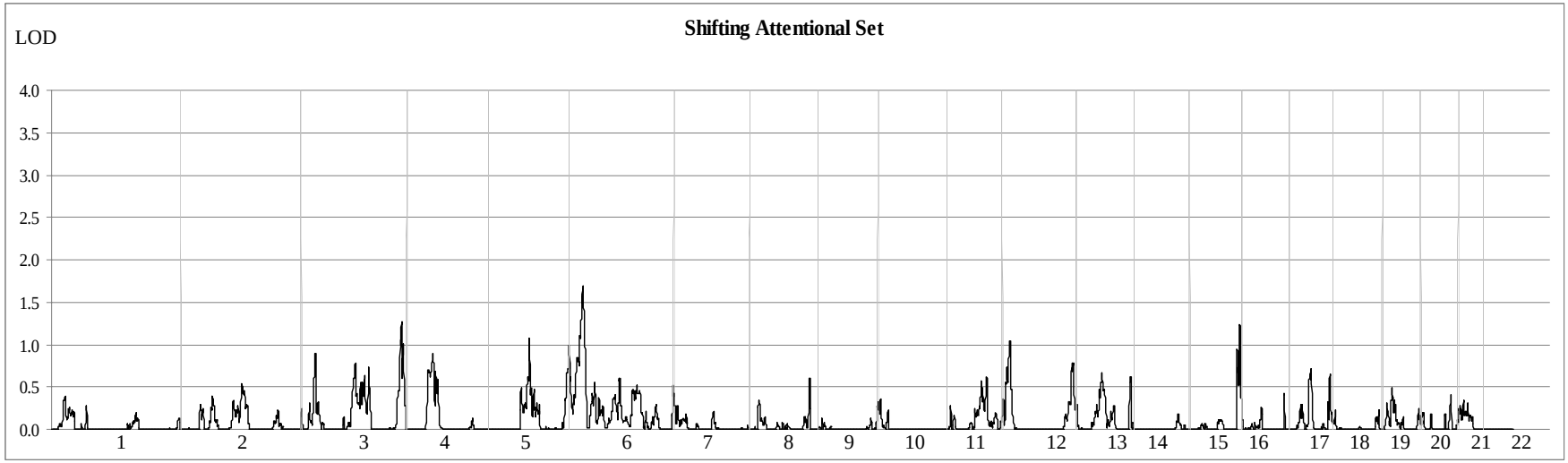
Figure S1. Genome-wide LOD Scores for the Eight Individual Neuropsychological Measures and the Combined Score




Shifting Attentional Set
LOD
4.0
3.5
3.0
2.5
2.0
1.5
1.0
0.5
0.0
1
2
3
4
5
6
7
8
9
10
11
12
13
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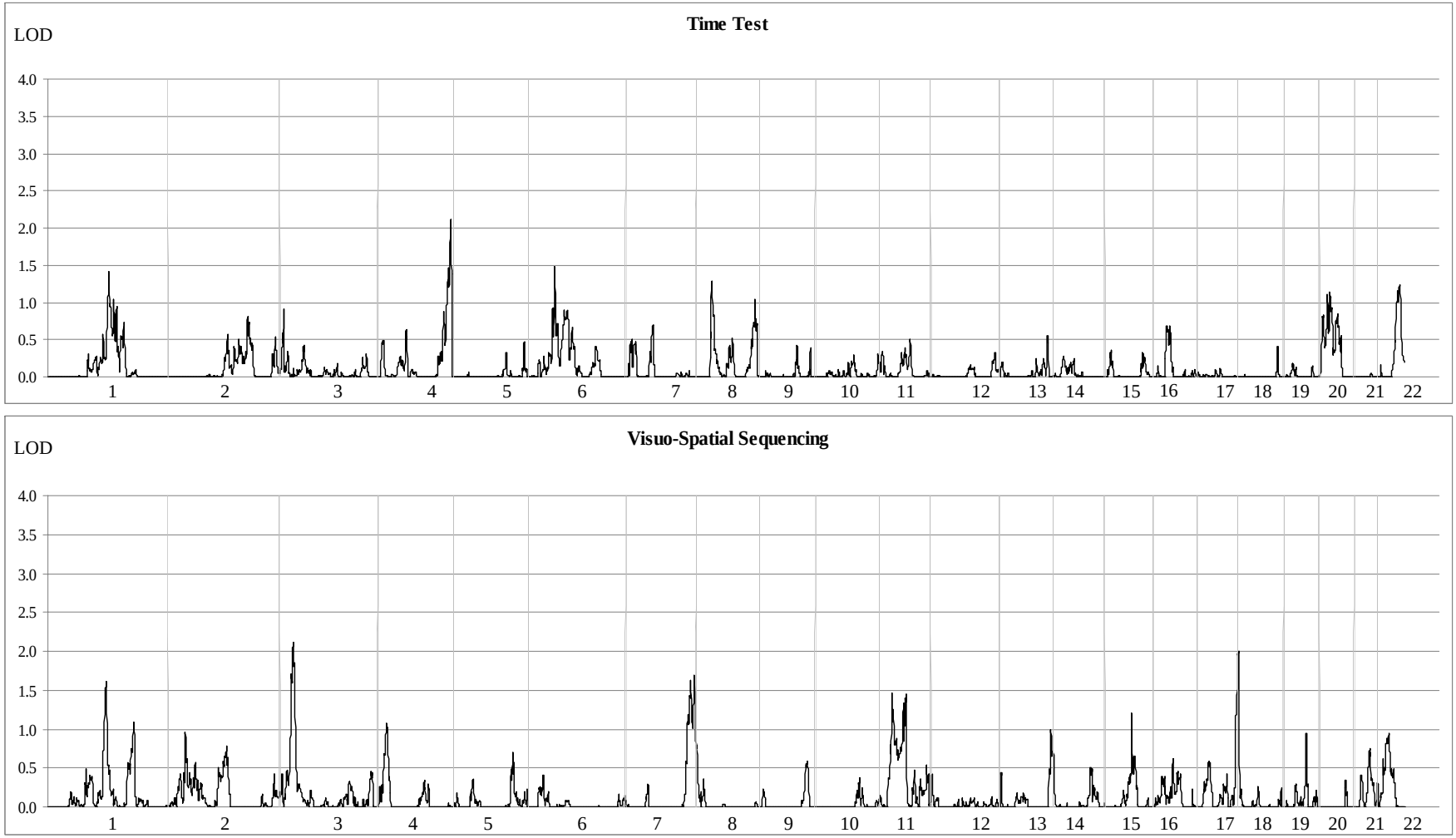
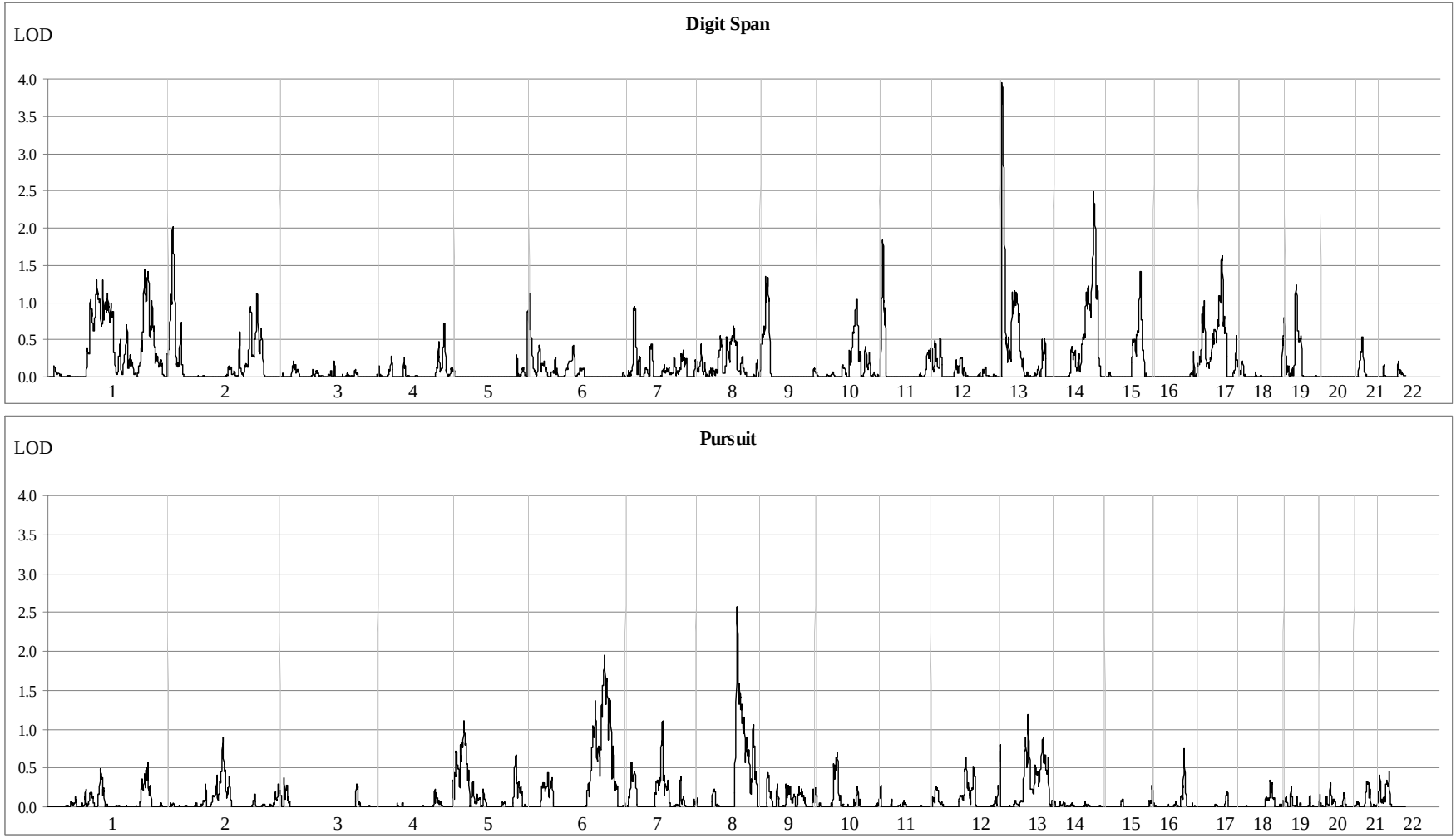
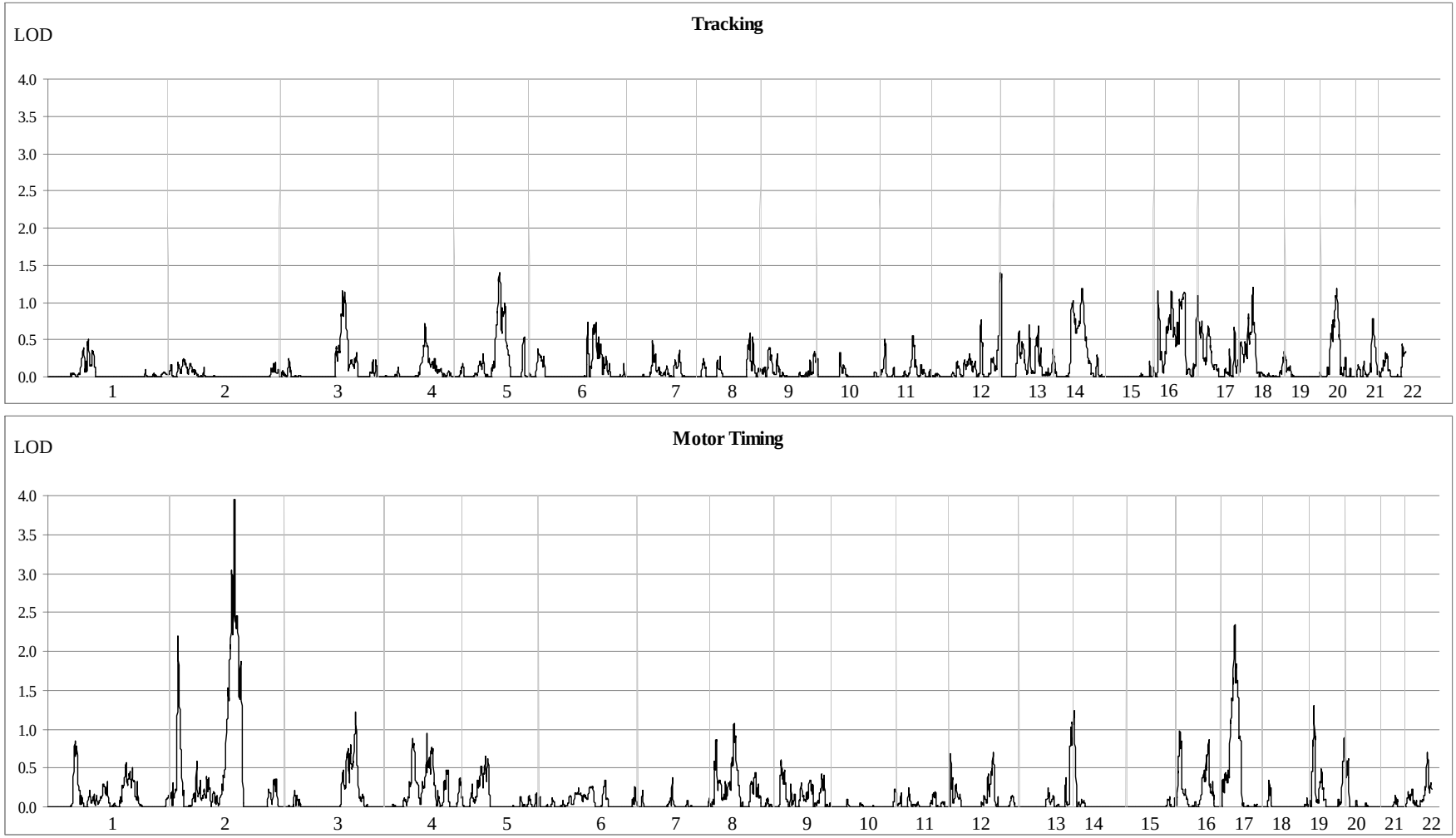
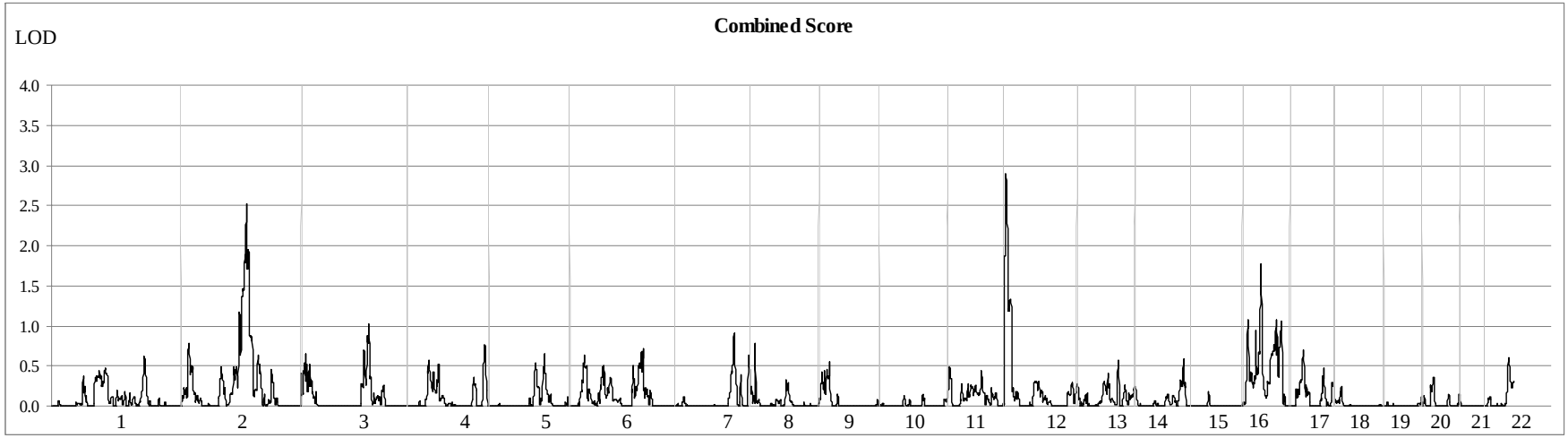
15
16
17
18
19
20
21
22


Time Test
LOD
4.0
3.5
3.0
2.5
2.0
1.5
1.0
0.5
0.0
1 2 3 4 5 6 7 8 9 10 11 12 13 14 15 16 17 18 19 20 21 22
Visuo-Spatial Sequencing
LOD
4.0
3.5
3.0
2.5
2.0
1.5
1.0
0.5
0.0
1 2 3 4 5 6 7 8 9 10 11 12 13 14 15 16 17 18 19 20 21 22


Digit Span
LOD
4.0
3.5
3.0
2.5
2.0
1.5
1.0
0.5
0.0
1 2 3 4 5 6 7 8 9 10 11 12 13 14 15 16 17 18 19 20 21 22
Pursuit
LOD
4.0
3.5
3.0
2.5
2.0
1.5
1.0
0.5
0.0
1 2 3 4 5 6 7 8 9 10 11 12 13 14 15 16 17 18 19 20 21 22


Tracking
LOD
4.0
3.5
3.0
2.5
2.0
1.5
1.0
0.5
0.0
1 2 3 4 5 6 7 8 9 10 11 12 13 14 15 16 17 18 19 20 21 22
Motor Timing
LOD
4.0
3.5
3.0
2.5
2.0
1.5
1.0
0.5
0.0
1 2 3 4 5 6 7 8 9 10 11 12 13 14 15 16 17 18 19 20 21 22


Combined Score
LOD
4.0
3.5
3.0
2.5
2.0
1.5
1.0
0.5
0.0
1
2
3
4
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6
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22

Table S1. Possible Candidate Genes within the Linkage Signals on Chromosomes 2 and 13

Name	Function
Chromosome 2	
ENI	*C*oding for Engrailed homeobox 1, a component of the Wnt signalling pathway involved in pattern formation during development of the central nervous system.
DBI	Encoding the diazepam binding inhibitor, a protein involved in the displacement of benzodiazepines, drugs that modulate signal transduction in brain synapses.
SCTR	Coding for a G protein-coupled receptor possibly involved in autism.
CNTNAP5	Gene product functions as a cell adhesion molecule and receptor in the nervous system.
BIN1	May be involved in synaptic vesicle endocytosis in the central nervous system.
PTPN18	Involved in cell growth and differentiation in the brain.
TUBA3E and *TUBA3D*	Expressed in the brain and possibly their function is related to that of the genes in the *TUBA1* family, which have been associated with abnormalities in the laminar architecture of the hippocampus and cortex accompanied by impaired neuronal migration in mouse mutants and showed association with lissencephaly in humans (Keays et al., 2007).
Chromosome 13	
TUBA3C	*See TUBA3E and TUBA3D.*
FGF9	Encodes a fibroblast growth factor with a possible role in glial cell development in the nervous system.