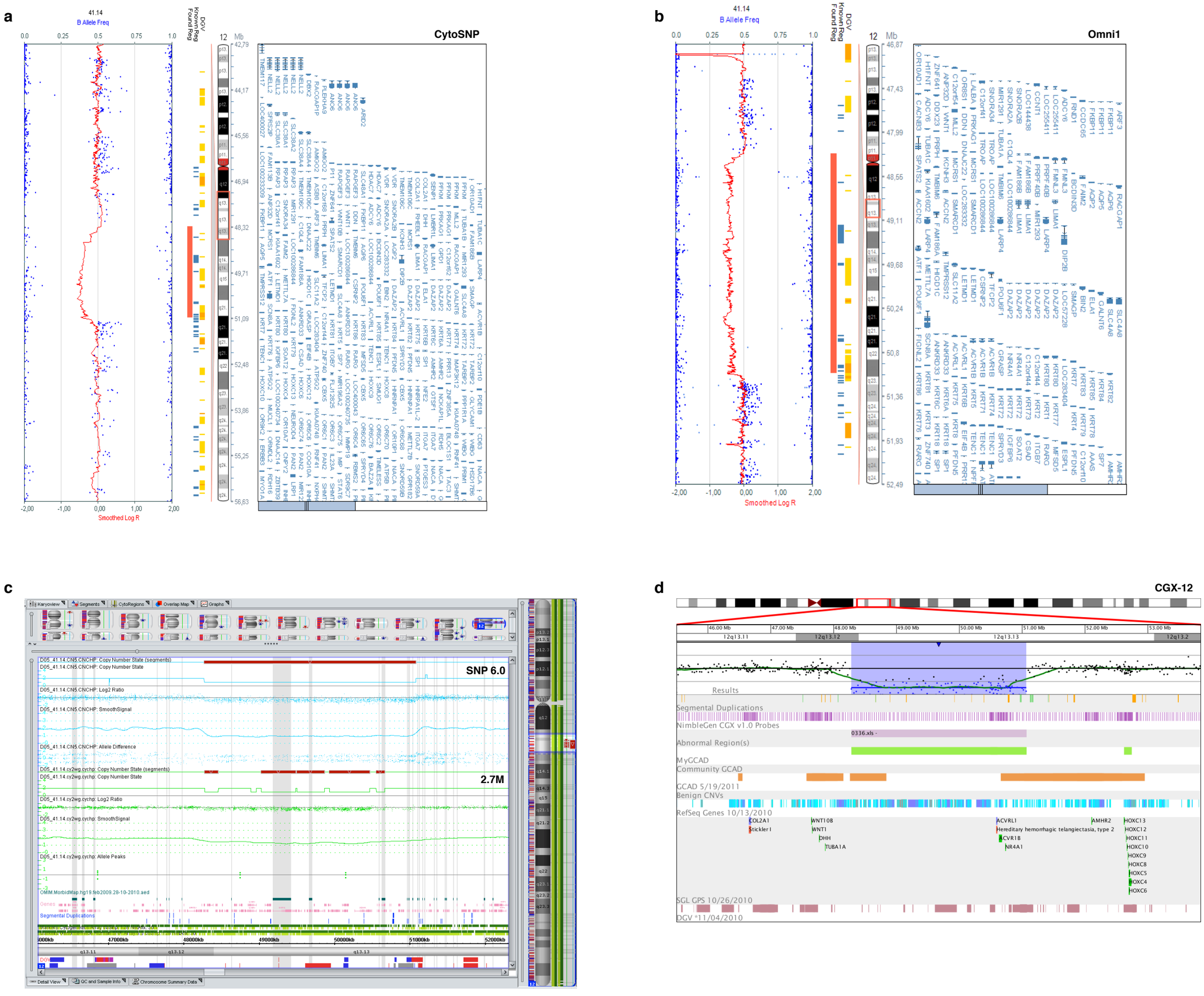
a
41.14
B Allele Freq
DGV
Known Reg
Found Reg
12
Mb
CytoSNP
Smoothed Log R
b
41.14
B Allele Freq
DGV
Known Reg
Found Reg
12
Mb
Omni1
Smoothed Log R
c
SNP 6.0
2.7M
d
CGX-12
Results
Segmental Duplications
NimbleGen CGX v1.0 Probes
Abnormal Region(s)
MyGCAD
Community GCAD
GCAD 5/19/2011
Benign CNVs
RefSeq Genes 10/13/2010
SGL GPS 10/26/2010
DGV *11/04/2010