

**Table S1.** Copy number aberrations (CNA) detected by array comparative genomic hybridization.

Case	Chromosomal location	Loc. Start*	Loc. Stop*	CNA**
Case 1	<b>3p26.3 - q29 (whole chromosome)</b>	<b>93949</b>	<b>197845254</b>	<b>Gain (0.320935)</b>
	4q13.2	69223075	69462438	Homozygous deletion (-1.51678)
	6p22.1	29854870	29896710	Gain (2.735431)
	<b>7p22.3 - q36.3 (whole chromosome)</b>	<b>42976</b>	<b>159118566</b>	<b>Gain (0.565555)</b>
	<b>9p24.3 - p13.1 (whole arm)</b>	<b>204193</b>	<b>38768291</b>	<b>Gain (0.308436)</b>
	9p21.3	21944037	22176231	Homozygous deletion (-1.382063)
	<b>9q21.11 - q34.3 (whole arm)</b>	<b>71151735</b>	<b>141018984</b>	<b>Gain (0.367163)</b>
	10p15.3 - q26.3 (whole chromosome)	136361	135089504	Heterozygous deletion (-0.350396)
	<b>13q12.11 - q34 (whole arm)</b>	<b>19526064</b>	<b>115089535</b>	<b>Gain (0.313476)</b>
	18q11.2 - q12.3	19004109	40832816	Heterozygous deletion (-0.327937)
	<b>18q12.3 - q23</b>	<b>41143464</b>	<b>78010032</b>	<b>Gain (0.317707)</b>
	20p13 - q13.33 (whole chromosome)	121521	62880583	Gain (0.330833)
	21q11.2 - q22.3 (whole arm)	15372467	48090317	Gain (0.558608)
	Case 2	<b>1p36.33 - p36.23</b>	<b>759762</b>	<b>8749497</b>
<b>1p35.3</b>		<b>28266522</b>	<b>29653134</b>	<b>Heterozygous deletion (-0.302864)</b>
<b>1p34.2 - p22.2</b>		<b>40288246</b>	<b>91311655</b>	<b>Heterozygous deletion (-0.279675)</b>
<b>1q32.2 - q42.2</b>		<b>209342761</b>	<b>230753927</b>	<b>Heterozygous deletion (-0.371906)</b>
<b>2p25.3 - p11.2 (whole arm)</b>		<b>42444</b>	<b>89538874</b>	<b>Gain (0.345952)</b>
<b>2q11.1 - q22.2</b>		<b>95529039</b>	<b>142709232</b>	<b>Gain (0.347157)</b>
<b>3q24 - q26.2</b>		<b>145644746</b>	<b>169235435</b>	<b>Gain (0.362882)</b>
<b>4q31.21 - q35.2</b>		<b>142663747</b>	<b>190790881</b>	<b>Heterozygous deletion (-0.380461)</b>
<b>6p25.3 - p11.2</b>		<b>389423</b>	<b>58435710</b>	<b>Heterozygous deletion (-0.317938)</b>
<b>6q11.1 - q22.33</b>		<b>61982931</b>	<b>127407686</b>	<b>Heterozygous deletion (-0.377149)</b>
<b>6q23.3 - q24.1</b>		<b>138476510</b>	<b>142468952</b>	<b>Heterozygous deletion (-0.371592)</b>
<b>7q33 - q36.3</b>		<b>135250552</b>	<b>159118566</b>	<b>Gain (0.593265)</b>
<b>8p23.3 - p11.1 (whole arm)</b>		<b>219683</b>	<b>43396776</b>	<b>Heterozygous deletion (-0.365243)</b>
<b>8q11.21 - q12.3</b>		<b>49091300</b>	<b>62336764</b>	<b>Heterozygous deletion (-0.404825)</b>
<b>8q13.3</b>		<b>71238938</b>	<b>72406658</b>	<b>Heterozygous deletion (-0.42469)</b>
<b>8q13.3 - q24.3</b>		<b>73125197</b>	<b>146085532</b>	<b>Heterozygous deletion (-0.372239)</b>
9p22.1 - p11.2		19564273	26162174	Heterozygous deletion (-0.750144)
9p21.3		21036673	22733971	Homozygous deletion (-1.333769)
9q21.13 - q31.3		79086651	112617941	Gain (0.548115)
<b>11q14.1 - q14.3</b>		<b>81732787</b>	<b>90039085</b>	<b>Gain (0.335935)</b>
<b>13q11 - q34 (whole arm)</b>		<b>19296544</b>	<b>115096466</b>	<b>Heterozygous deletion (-0.374167)</b>
<b>16q22.2 - q23.1</b>		<b>71913790</b>	<b>75360302</b>	<b>Heterozygous deletion (-0.283991)</b>
<b>16q23.2 - q24.3</b>		<b>79225966</b>	<b>89977030</b>	<b>Heterozygous deletion (-0.389361)</b>
<b>17p13.3 - p11.2 (whole arm)</b>		<b>76263</b>	<b>22154574</b>	<b>Heterozygous deletion (-0.349955)</b>
<b>17q11.1 - q11.2</b>		<b>25435421</b>	<b>30348467</b>	<b>Heterozygous deletion (-0.324576)</b>
<b>17q21.2 - q21.31</b>		<b>40318960</b>	<b>42940605</b>	<b>Heterozygous deletion (-0.311494)</b>
<b>17q21.31 - q25.1</b>		<b>42962663</b>	<b>72659034</b>	<b>Gain (0.828081)</b>
<b>18p11.32 - q23 (whole chromosome)</b>		<b>162510</b>	<b>78010032</b>	<b>Heterozygous deletion (-0.354022)</b>
<b>22q12.1 - q13.33</b>	<b>27099649</b>	<b>51219009</b>	<b>Gain (0.396596)</b>	
Case 3	<b>6q11.1 - q27</b>	<b>62305272</b>	<b>166262838</b>	<b>Heterozygous deletion (-0.268362)</b>
	<b>7p22.3 - q36.3 (whole chromosome)</b>	<b>65558</b>	<b>159118566</b>	<b>Gain (0.547574)</b>
	8p11.22	39237438	39374789	Homozygous deletion (-1.165137)
	<b>9p22.3</b>	<b>15086757</b>	<b>15920360</b>	<b>Heterozygous deletion (-0.300928)</b>
	<b>9p21.3</b>	<b>21454717</b>	<b>22212612</b>	<b>Homozygous deletion (-1.268826)</b>
	<b>21q11.2 - q22.3 (whole arm)</b>	<b>14420615</b>	<b>48090317</b>	<b>Gain (0.29937)</b>
Case 4	22q11.23	24347959	24390254	Homozygous deletion (-1.52085)
	5p15.33 - q35.3 (whole chromosome)	26142	180684501	Gain (0.429411)
	6p22.1	29854870	29896710	Gain (2.672552)
	<b>7p22.3 - q36.3 (whole chromosome)</b>	<b>42976</b>	<b>159118566</b>	<b>Gain (0.439458)</b>
	8p11.22	39237438	39374789	Homozygous deletion (-1.238193)
	9p21.3	21957735	22017397	Homozygous deletion (-1.342227)
	10p15.3 - p11.1(whole arm)	148206	39076591	Heterozygous deletion (-0.549255)
	<b>10q11.21 - q26.3 (whole arm)</b>	<b>42864620</b>	<b>133956779</b>	<b>Heterozygous deletion (-0.536525)</b>
	13q12.11 - q34 (whole arm)	19526064	115089535	Gain (0.775586)
	<b>17p13.3 - p11.2 (whole arm)</b>	<b>96559</b>	<b>22154574</b>	<b>Gain (0.339977)</b>
17q11.1 - q25.3 (whole arm)	25403446	81062925	Gain (0.420042)	
Case 5	<b>19q13.12 - q13.13</b>	<b>37349894</b>	<b>38698864</b>	<b>Heterozygous deletion (-0.308813)</b>
	20p13 - q13.33 (whole chromosome)	151549	62904501	Gain (0.356953)
	<b>2q22.1</b>	<b>141928411</b>	<b>142036954</b>	<b>Heterozygous deletion (-0.815427)</b>
	<b>3q13.31</b>	<b>115593958</b>	<b>115644733</b>	<b>Homozygous deletion (-1.991462)</b>
	3q26.1	162514534	162619141	Homozygous deletion (-1.849218)
	4q13.2	69392545	69462438	Homozygous deletion (-1.831354)
	<b>4q34.3 - q35.1</b>	<b>182902844</b>	<b>183568174</b>	<b>Heterozygous deletion (-0.61287)</b>
	6p22.1	29854870	29896710	Homozygous deletion (-1.236831)
	8p11.22	39237438	39345479	Gain (0.671442)
	9p21.3	21944037	22176231	Homozygous deletion (-1.62978)
	15q11.1-q11.2	20575646	21933378	Gain (0.937772)
	20p12.1	14903016	15015311	Heterozygous deletion (-0.880196)
	22q11.23	24347959	24390254	Homozygous deletion (-1.449985)

CNA detected only in the epithelioid glioblastoma area are shown in **bold**.

CNA detected both in the epithelioid glioblastoma and lower-grade glioma areas are not in bold.

Loc., location on chromosome.

\*Genome mapping based on genome build hg19.

\*\*The numbers in parentheses are the average log2 ratio of each aberration.