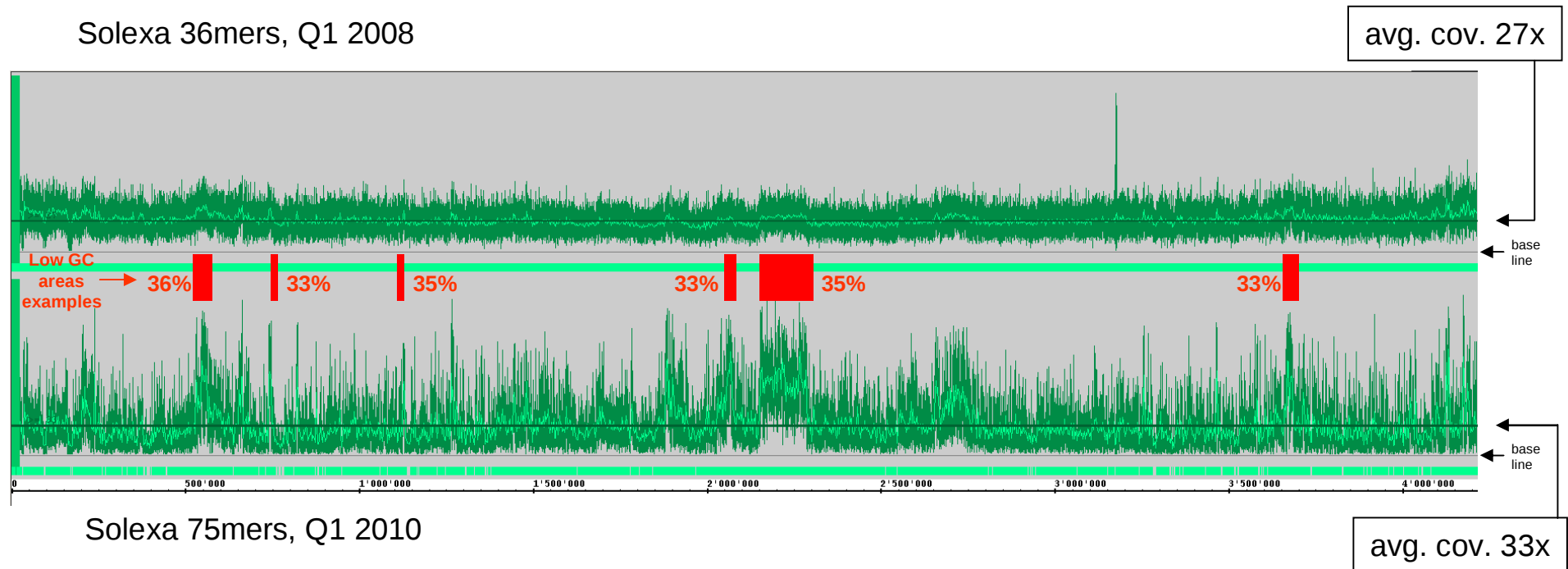


Comparison Solexa sequencing 2008 / 2010
Bacterium, 45%GC average, mapped on perfect reference sequence

Analysis of coverage uniformity



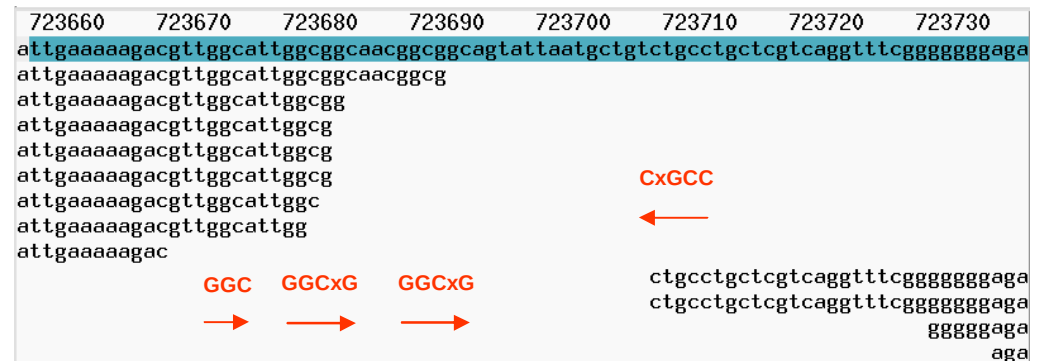
- 2008 coverage uniformity much better than 2010
- bias: higher coverage in low GC regions
 - bias barely noticeable in 2008 data
 - in 2010, **bias strong enough** in low GC regions (red bars in image) **to look like genome duplication!**

Comparison Solexa sequencing 2008 / 2010

Bacterium, 45%GC average, mapped on perfect reference sequence

Analysis of holes in coverage

- 2008 data (27x avg. cov., 36mers):
 - zero unexplained holes in the genome, everything covered
 - perfect
- 2010 data (33x avg. cov., 75mers)
 - 237 unexplained holes in the genome which are not repetitive areas like rRNA etc.
 - many more areas only with coverage of 1 to 5
 - often at sites with GGC or stronger even with GGCxG in 5' to 3' direction
 - worst with multiple local sites, or even when in both directions



→ For 75mers in 2010, avg. coverage of $\geq 70x$ to $80x$ needs to be present get complete coverage (i.e., no holes)