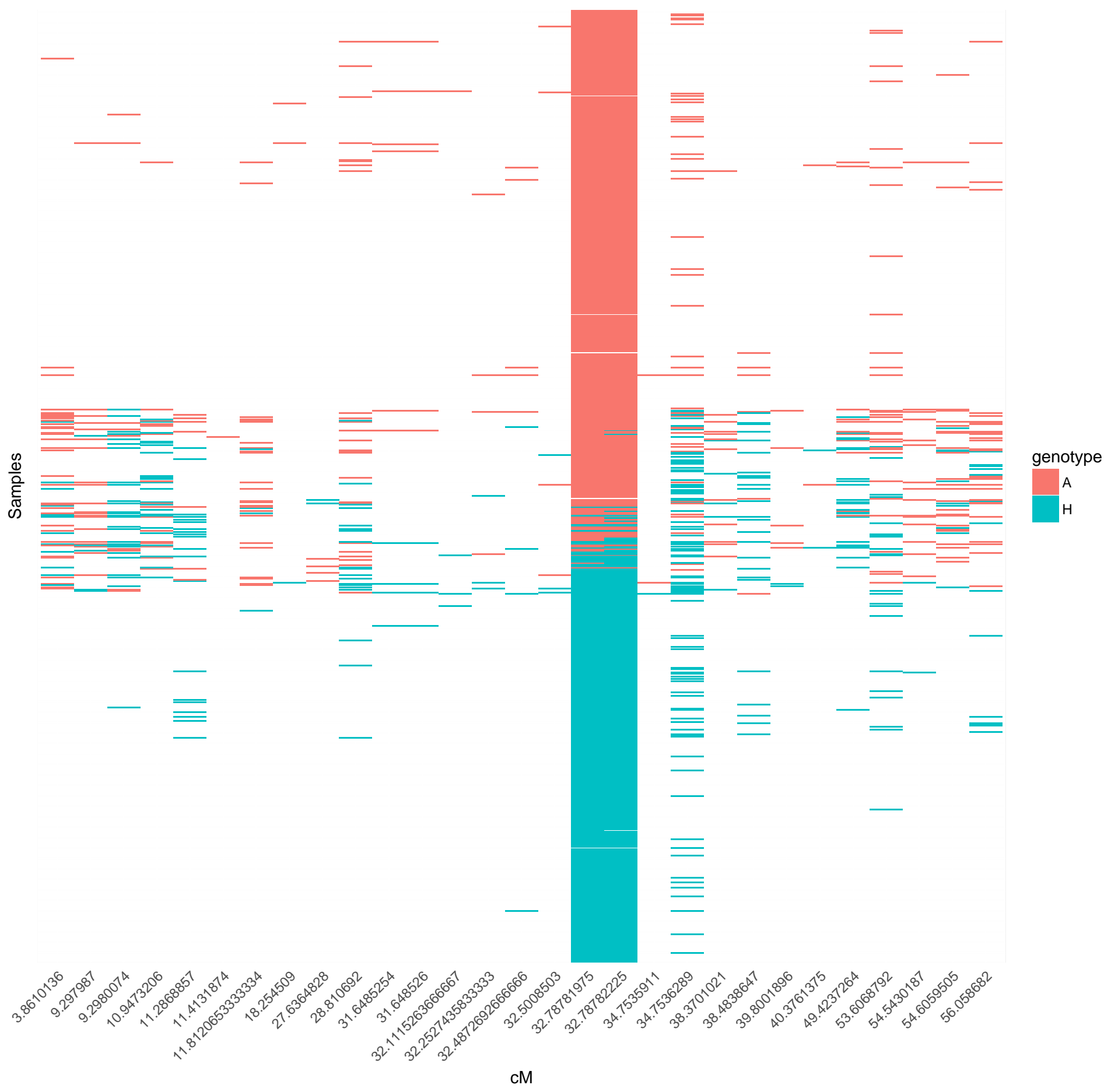


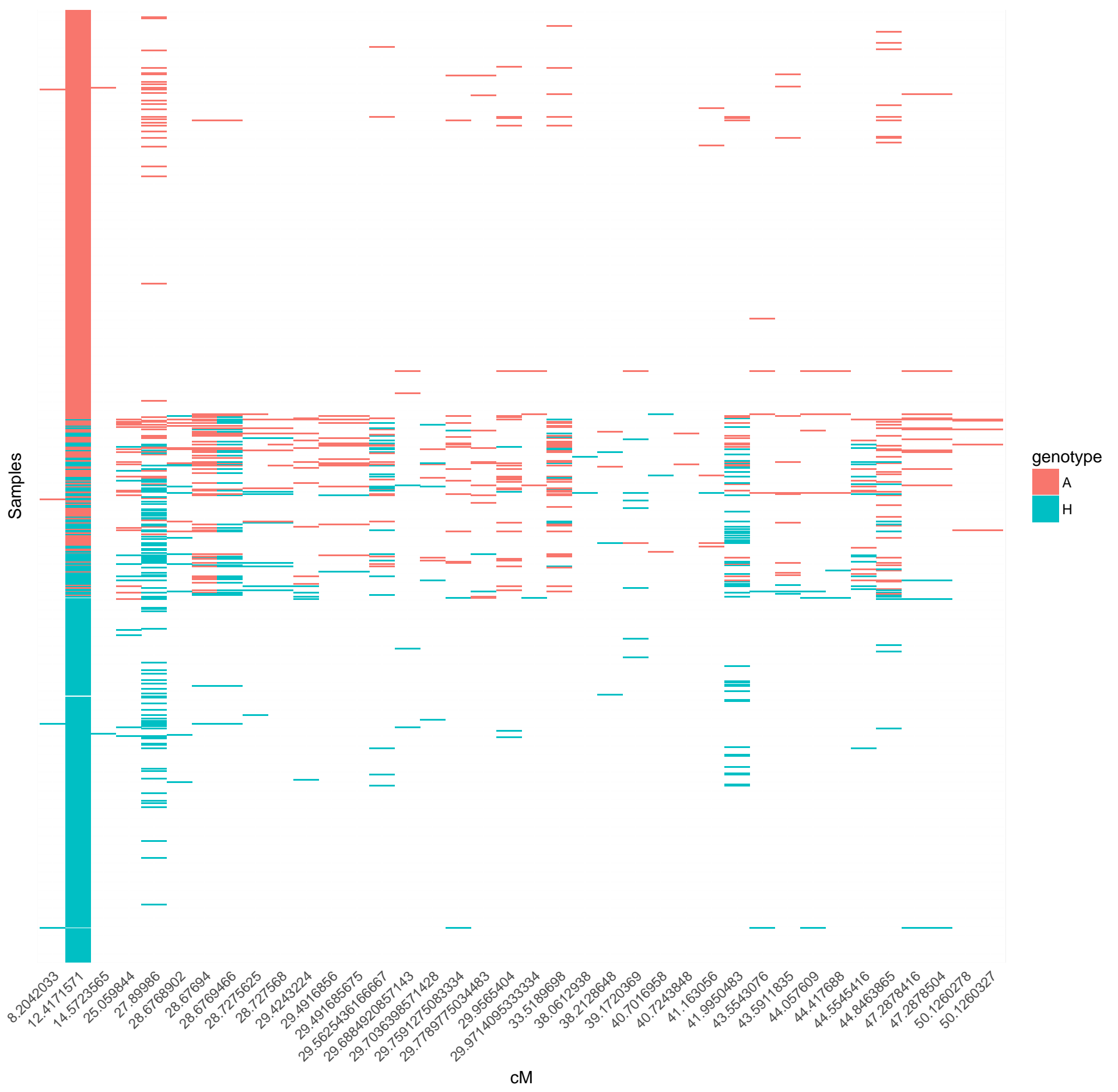
ann2 BC1, missing data, Chr 01



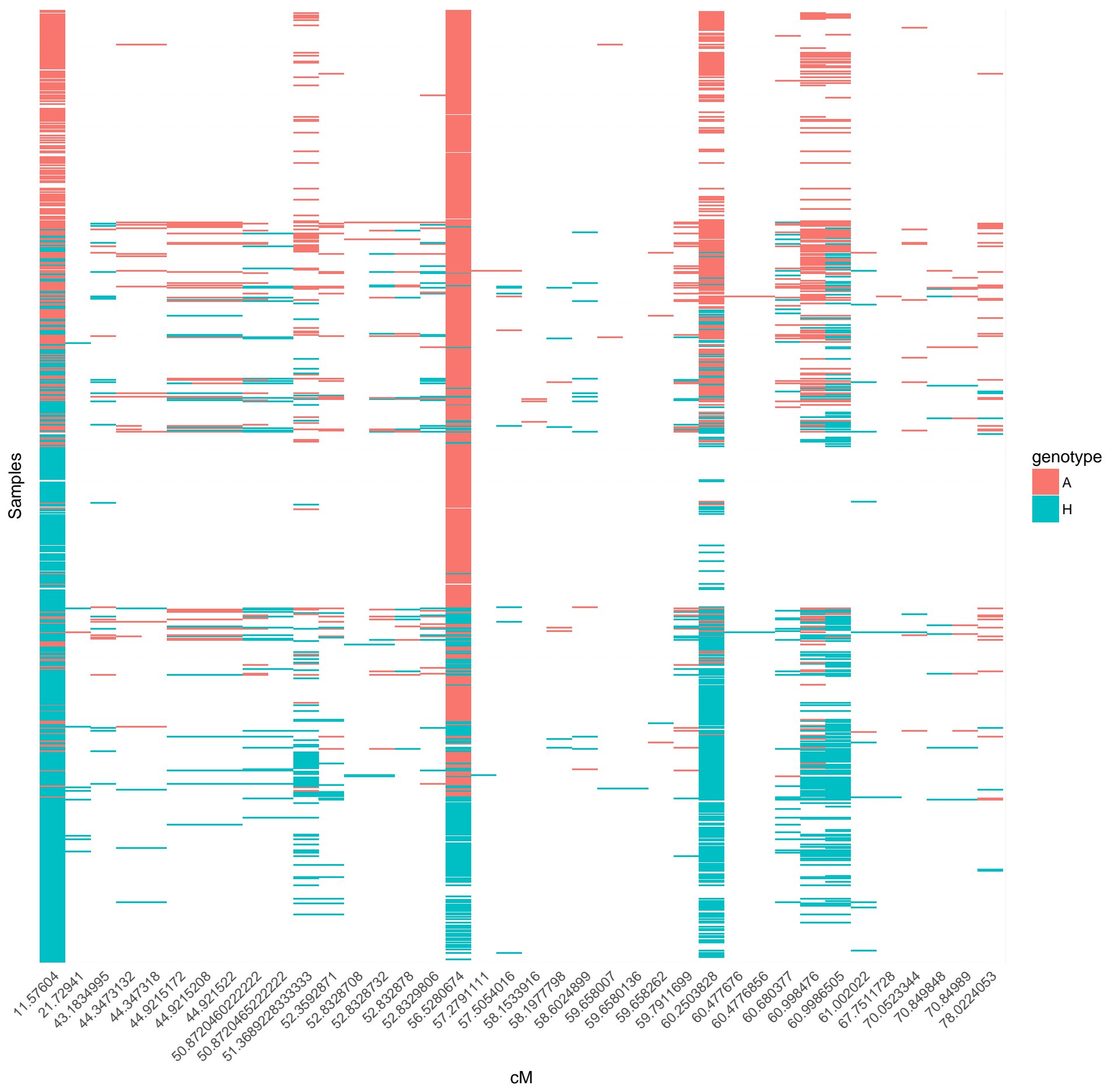
ann2 BC1, missing data, Chr 02



ann2 BC1, missing data, Chr 03



ann2 BC1, missing data, Chr 04



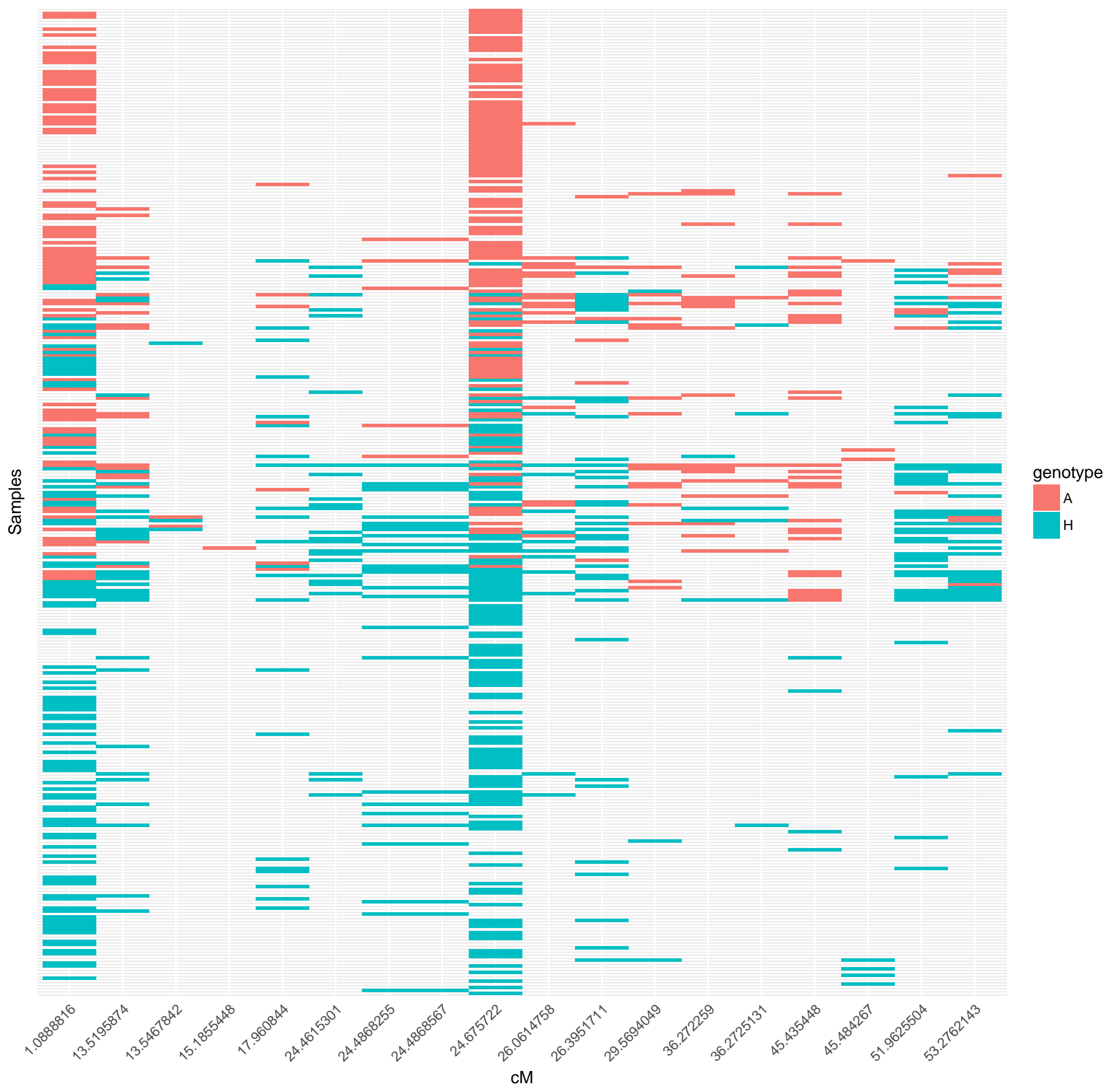
ann2 BC1, missing data, Chr 05



ann2 BC1, missing data, Chr 06



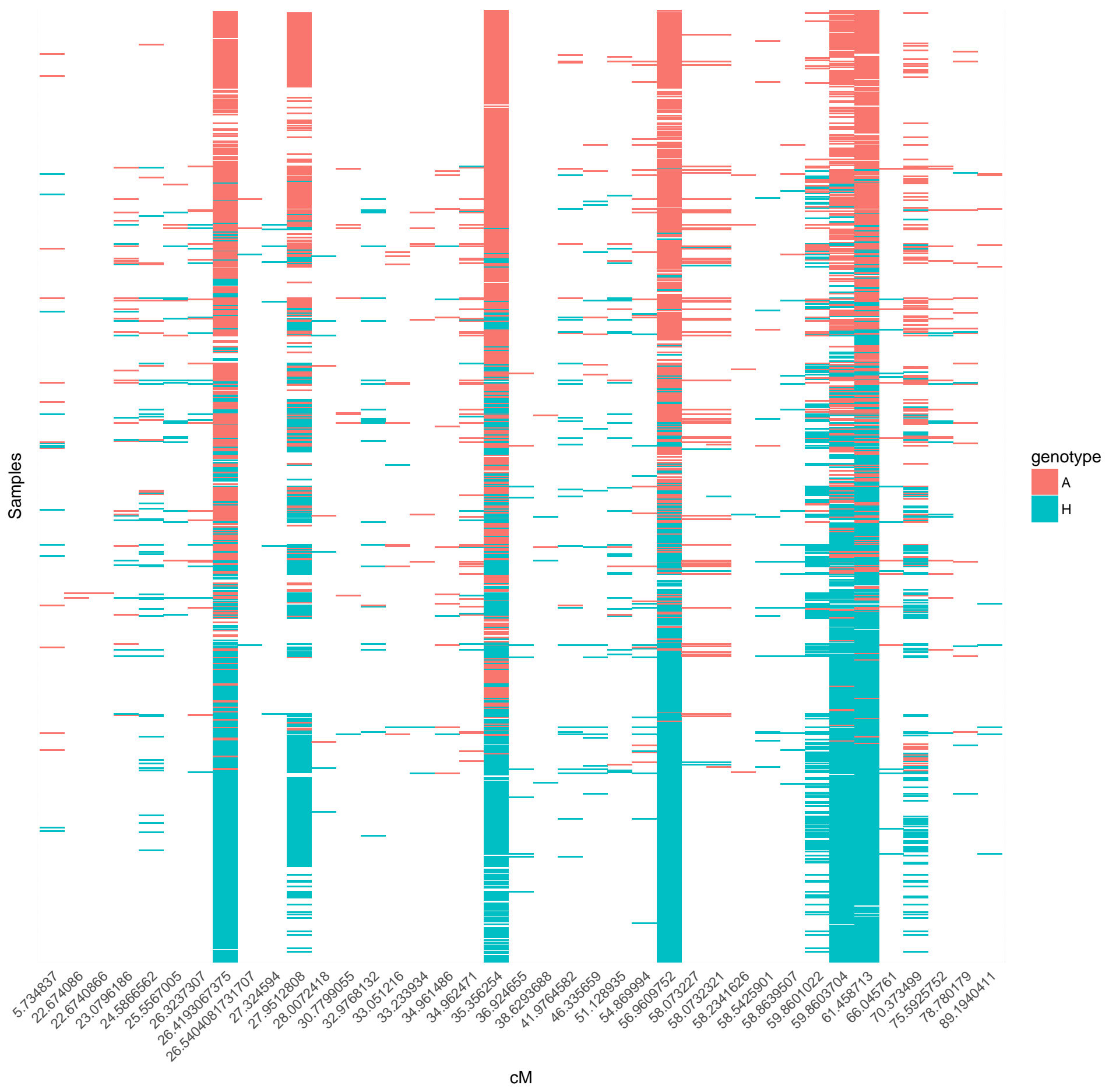
ann2 BC1, missing data, Chr 07



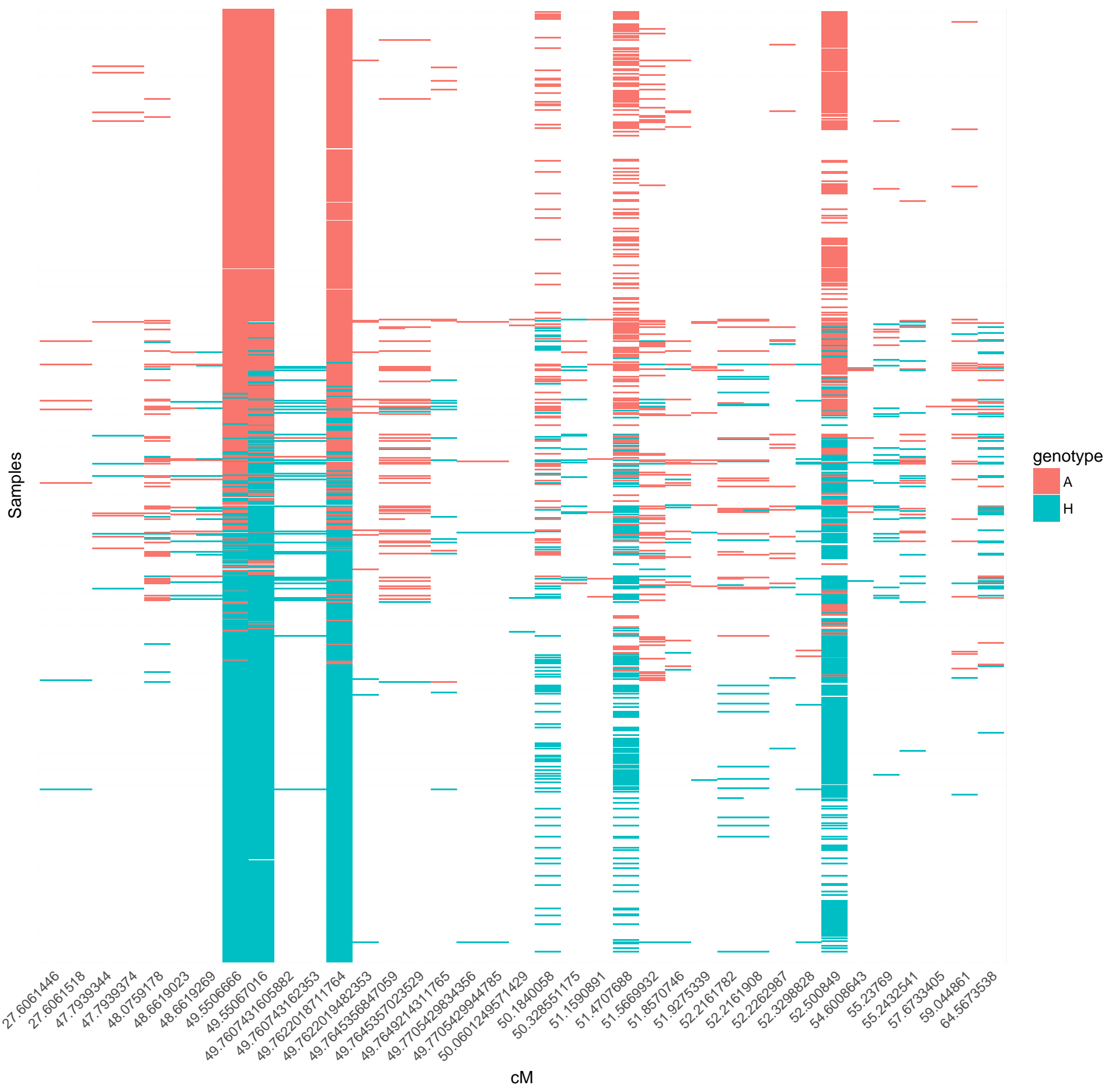
ann2 BC1, missing data, Chr 08



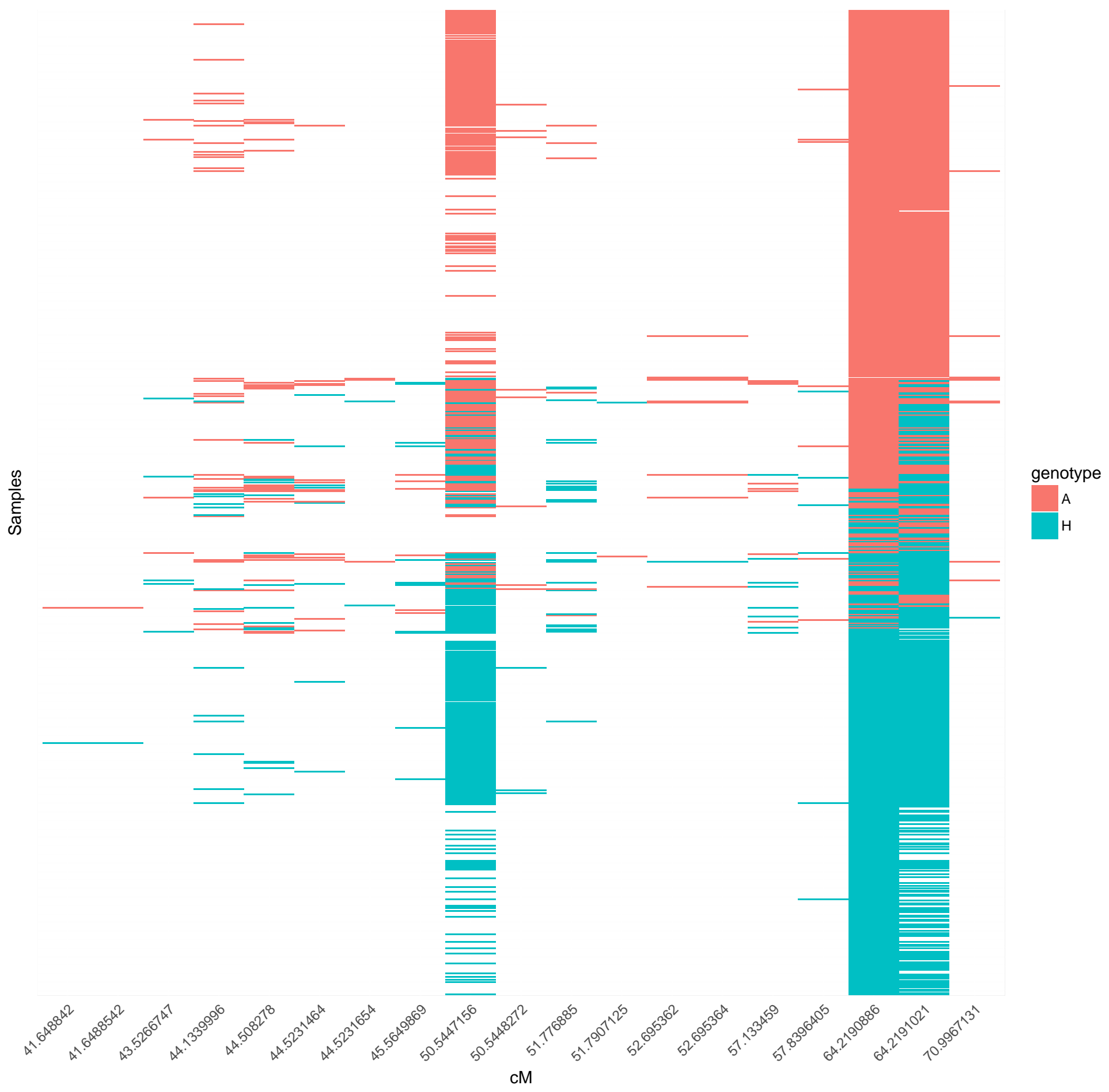
ann2 BC1, missing data, Chr 09



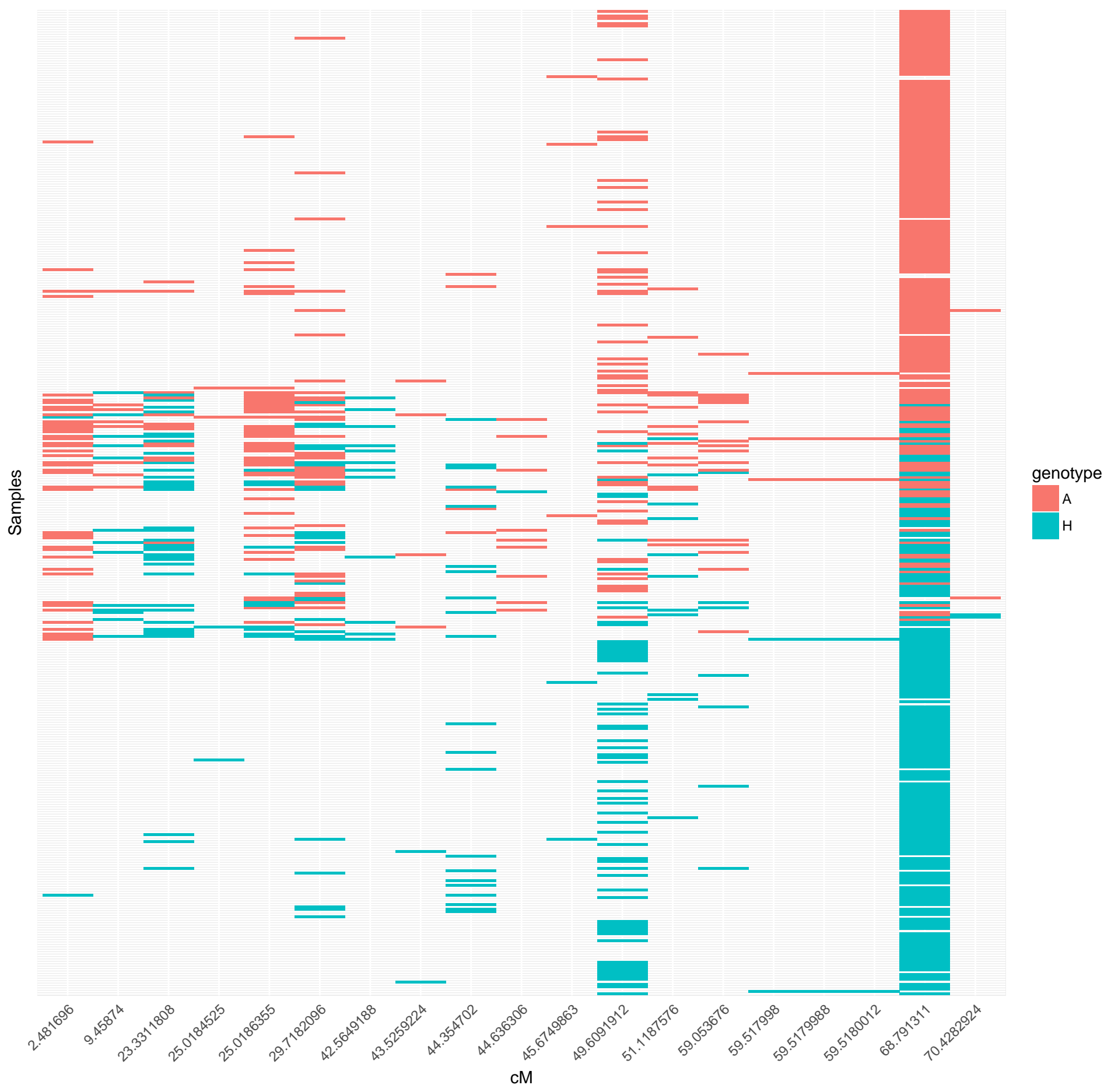
ann2 BC1, missing data, Chr 10



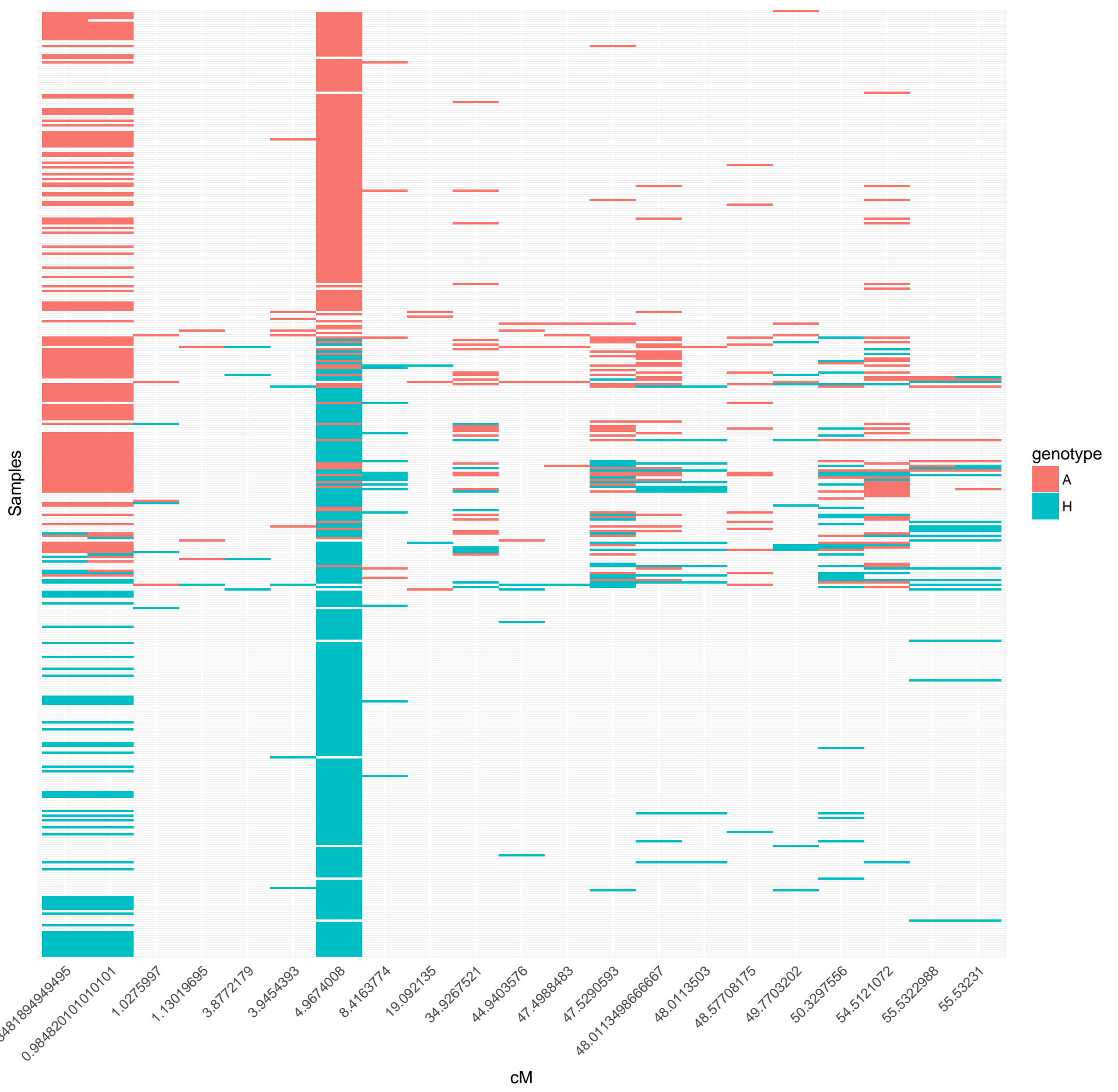
ann2 BC1, missing data, Chr 11



ann2 BC1, missing data, Chr 12



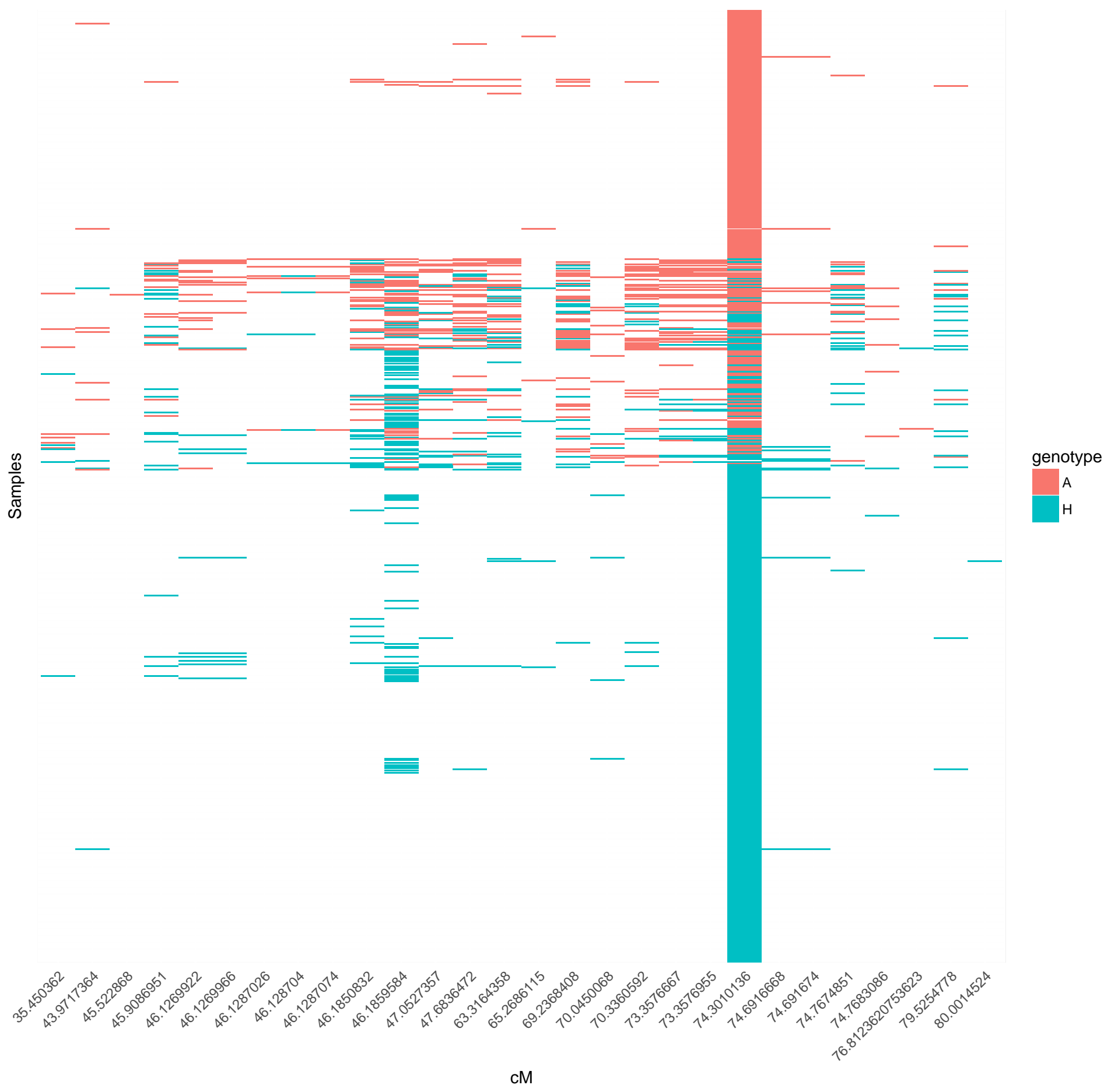
ann2 BC1, missing data, Chr 13



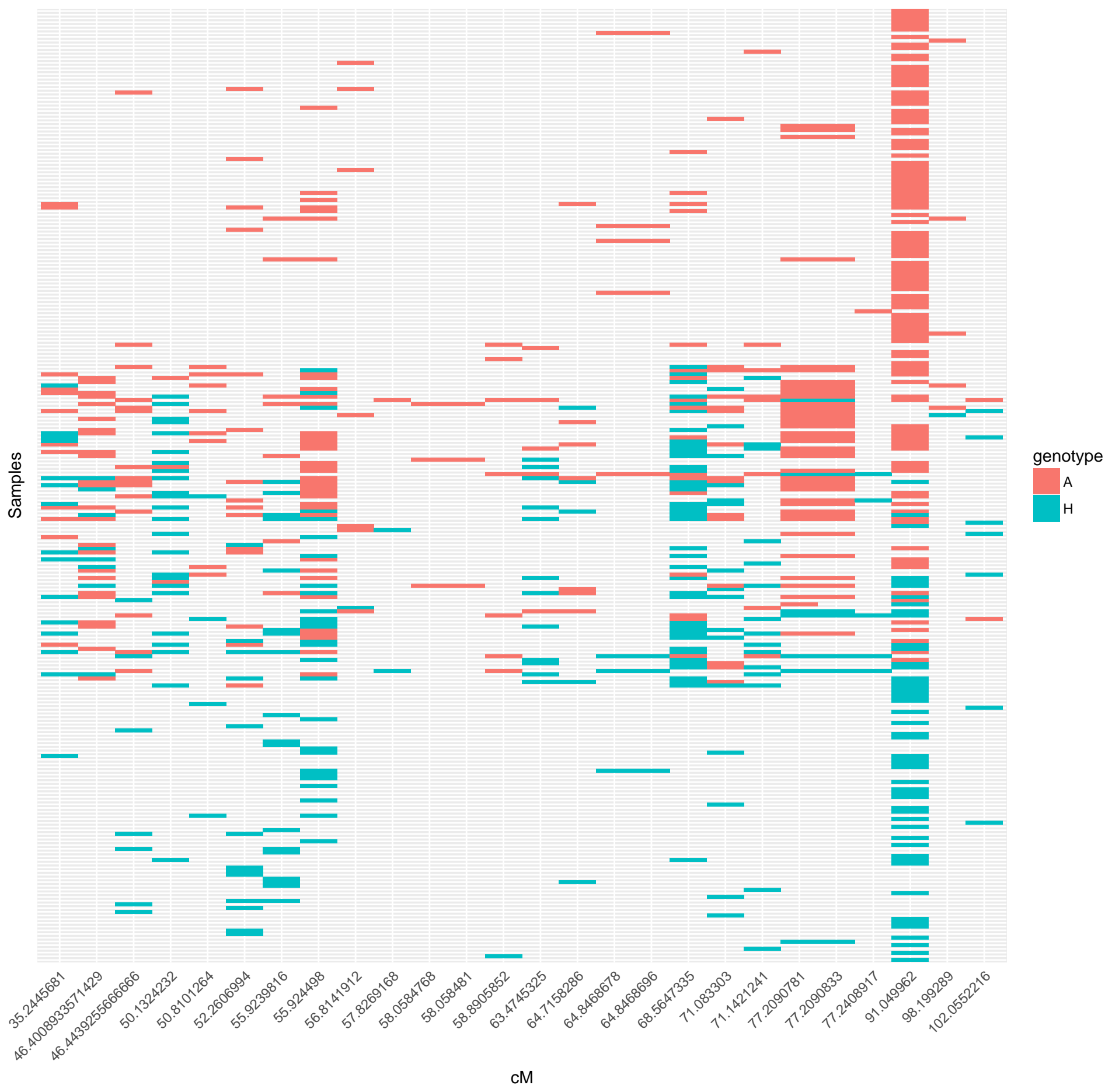
ann2 BC1, missing data, Chr 14



ann2 BC1, missing data, Chr 15



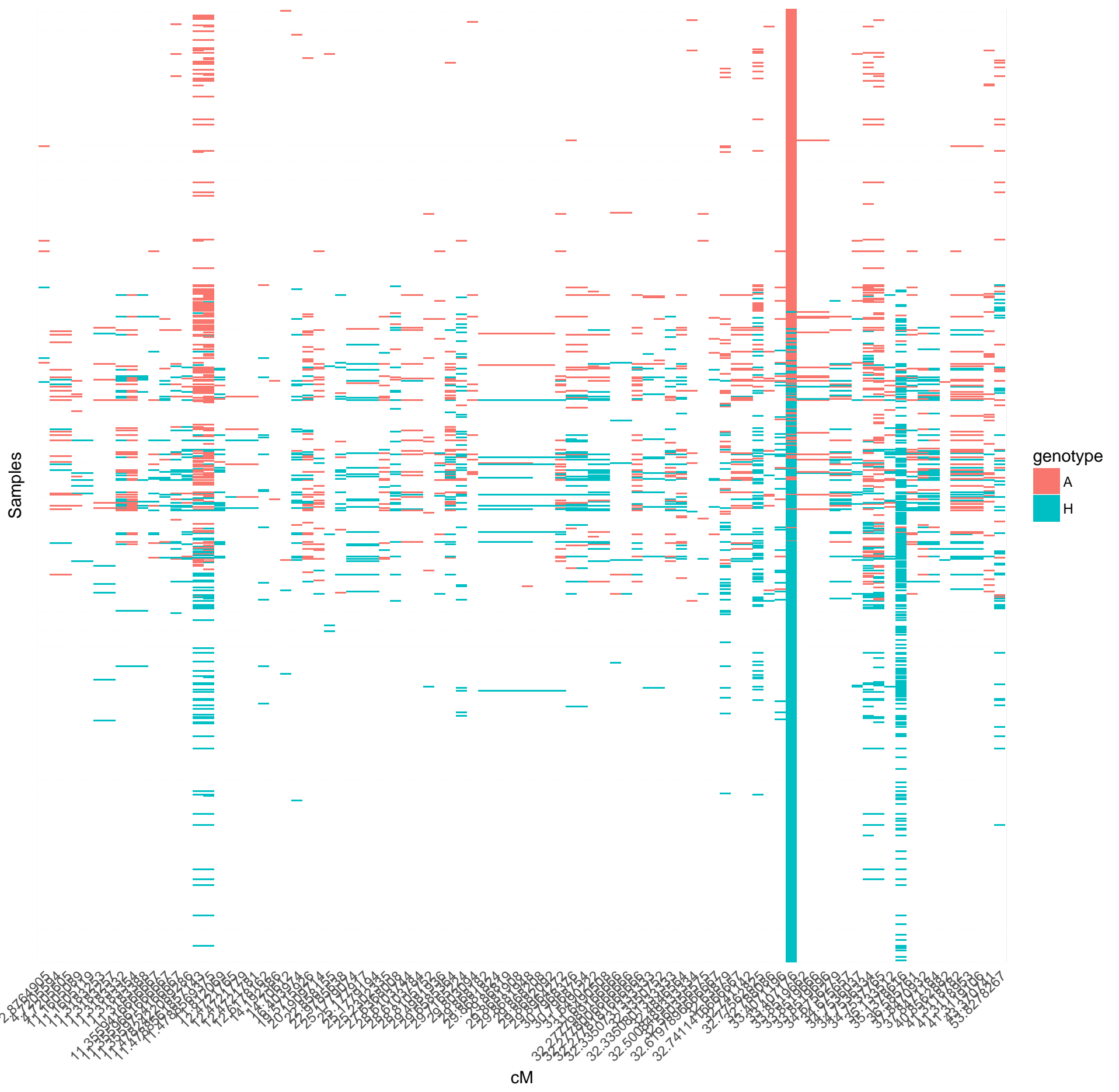
ann2 BC1, missing data, Chr 16



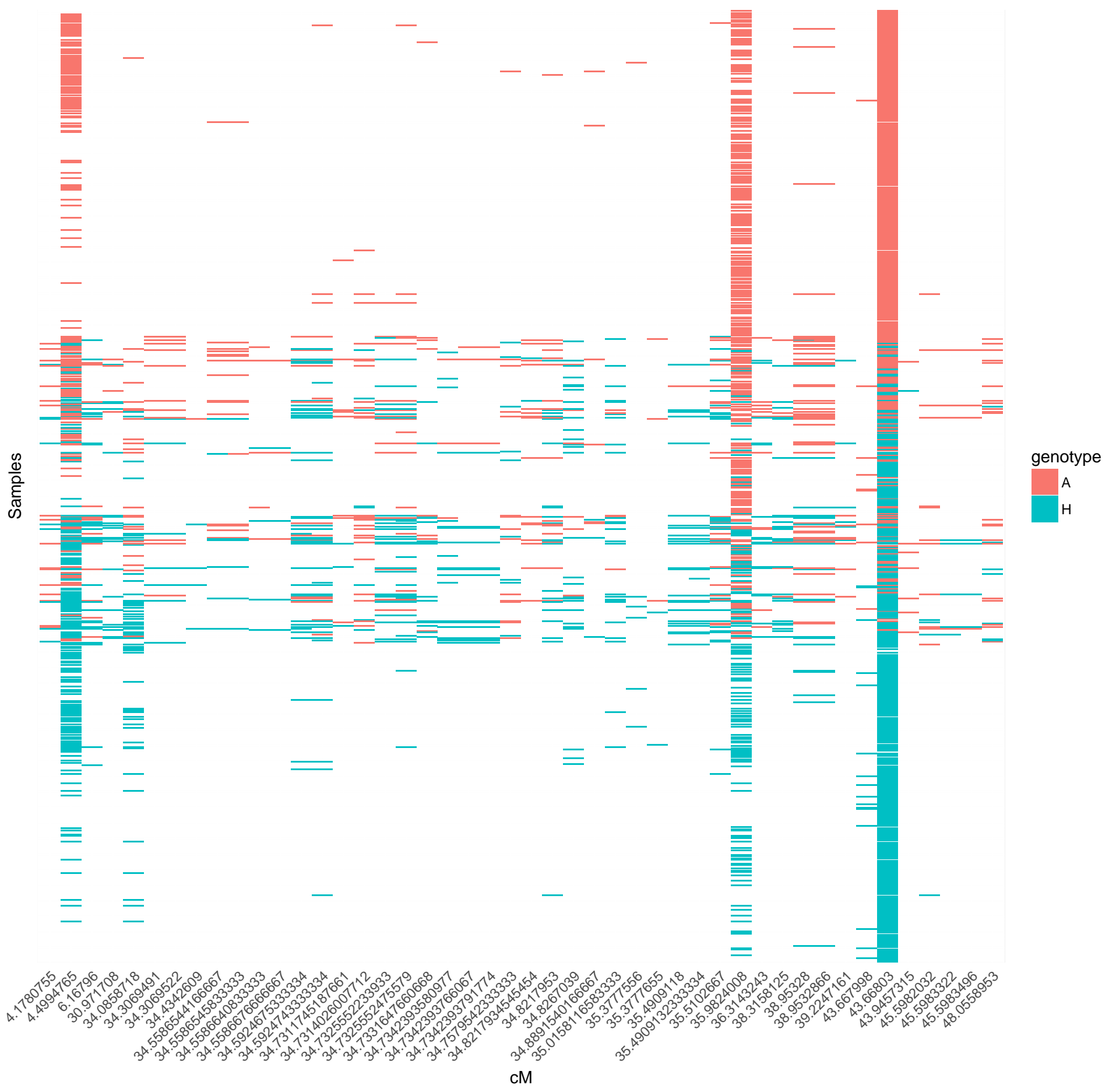
ann2 BC1, missing data, Chr 17



deb BC1, missing data, Chr 01

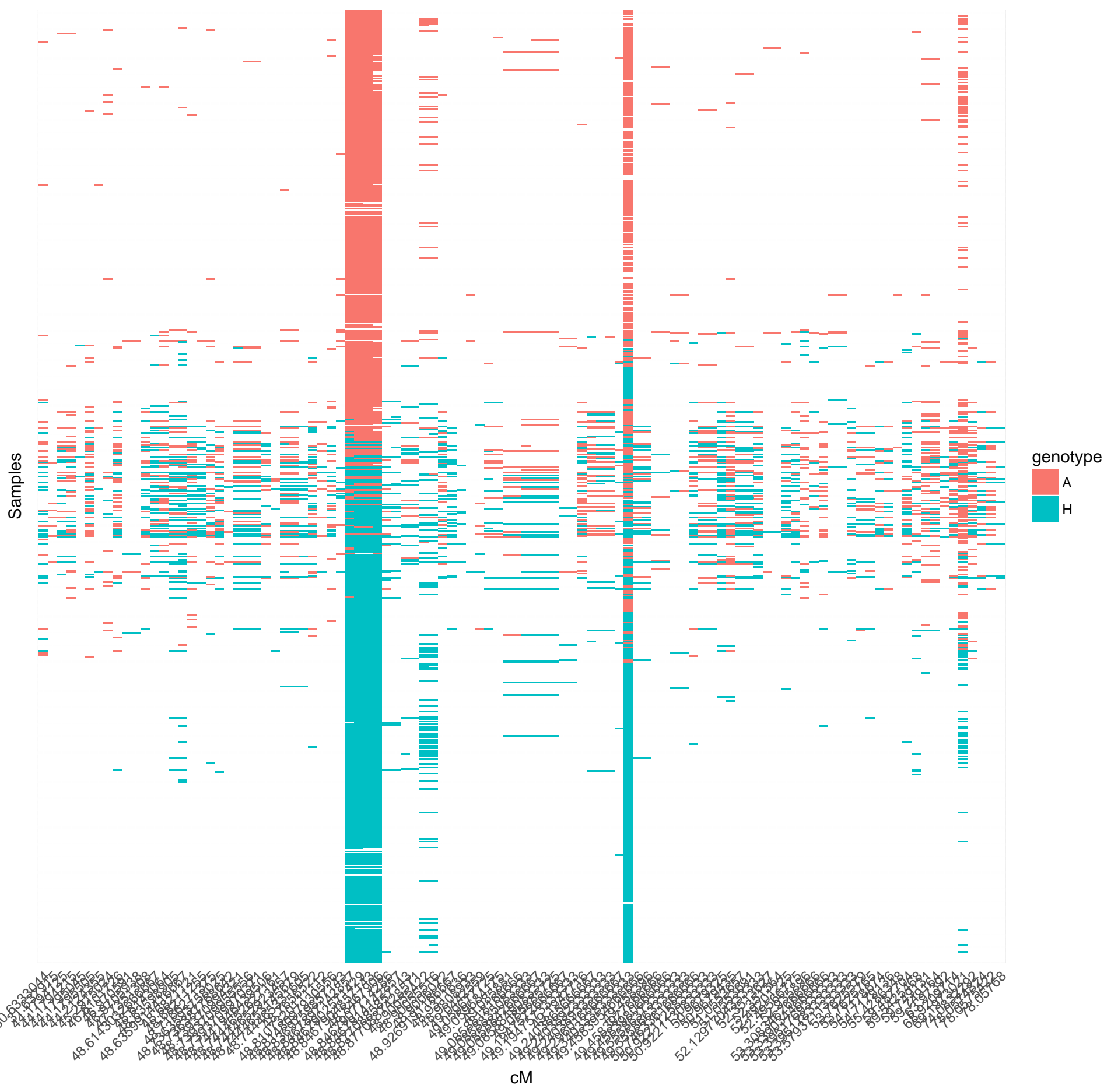


deb BC1, missing data, Chr 02

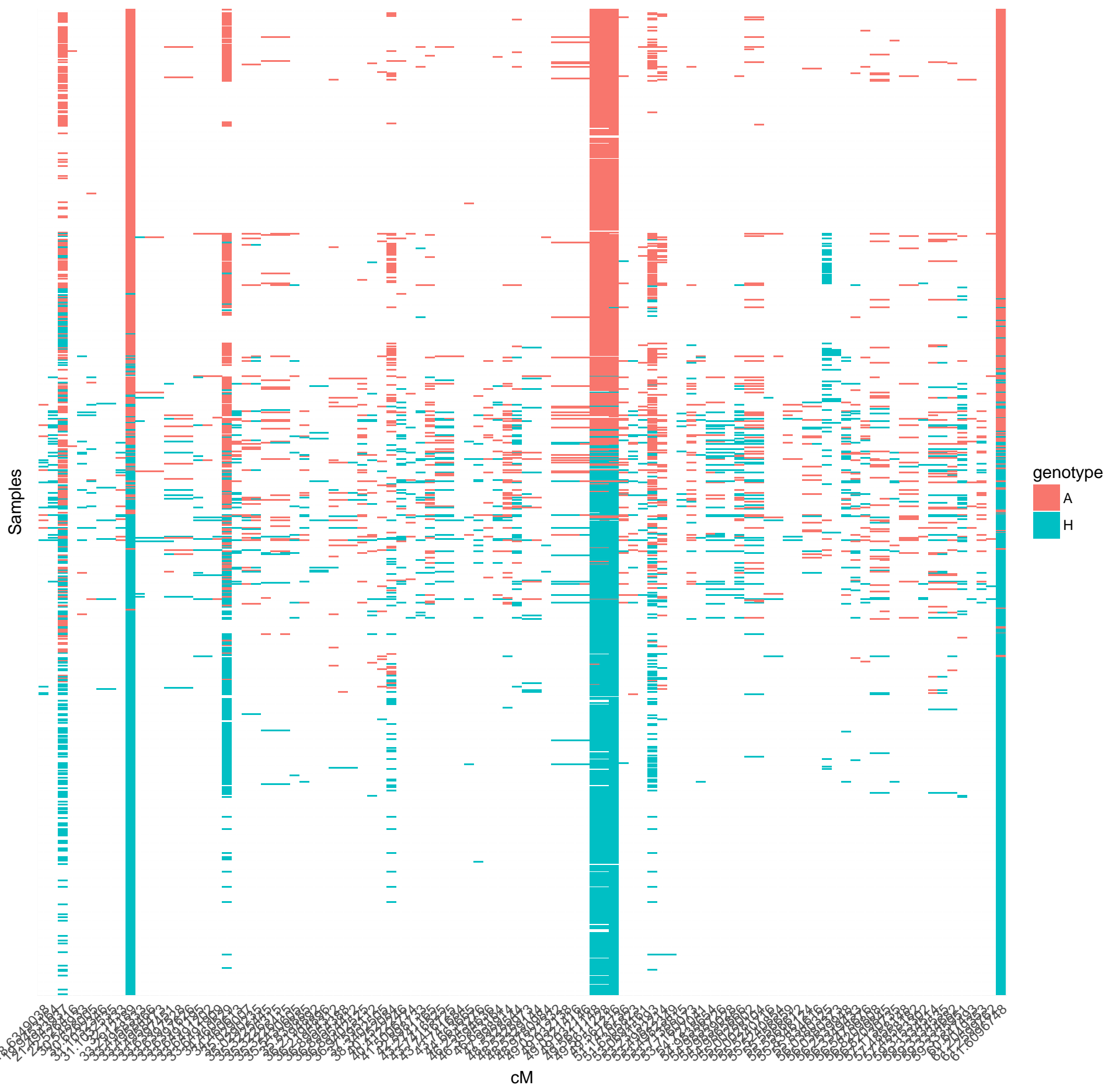


A
H

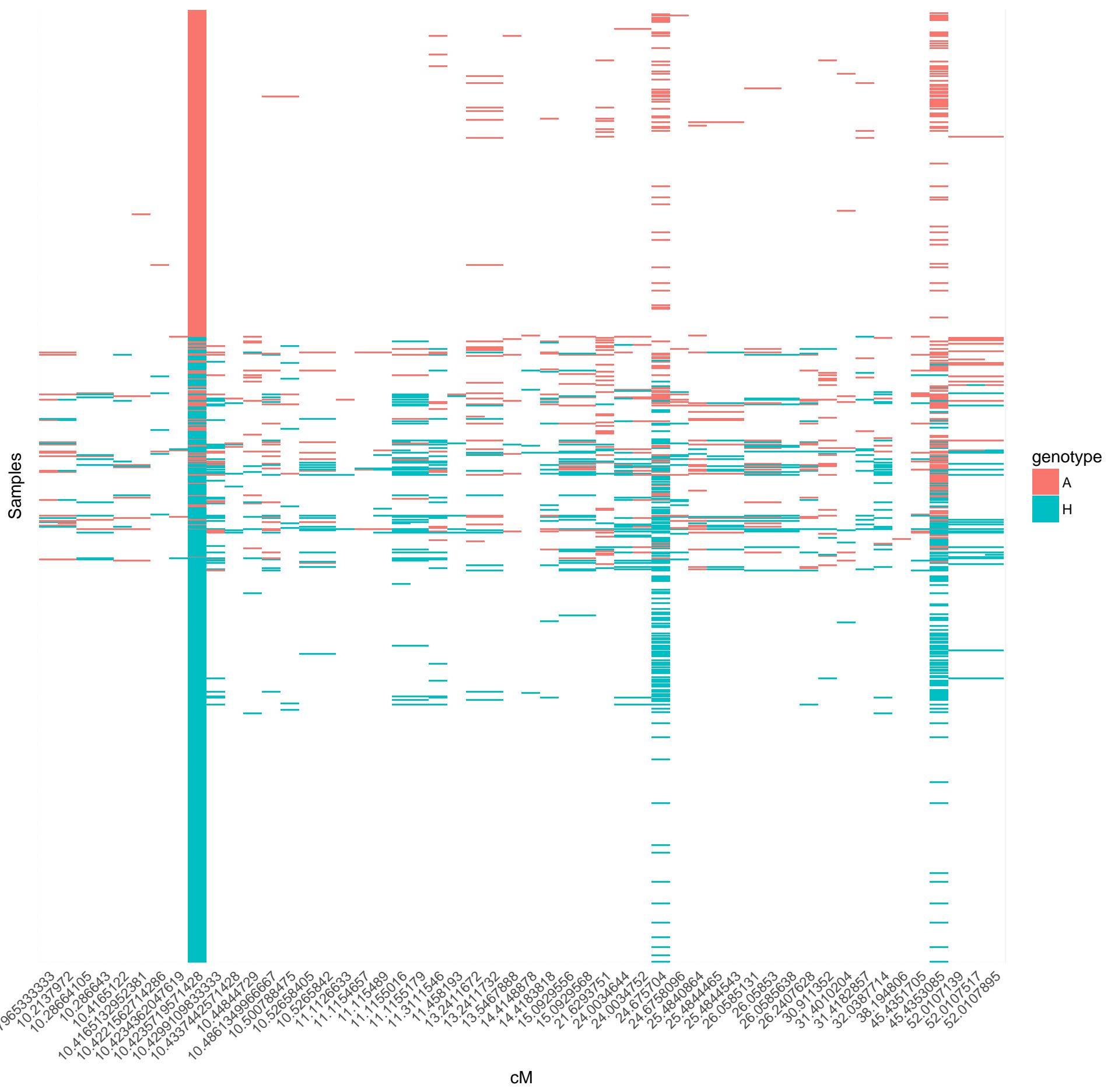
deb BC1, missing data, Chr 05



deb BC1, missing data, Chr 06



deb BC1, missing data, Chr 07



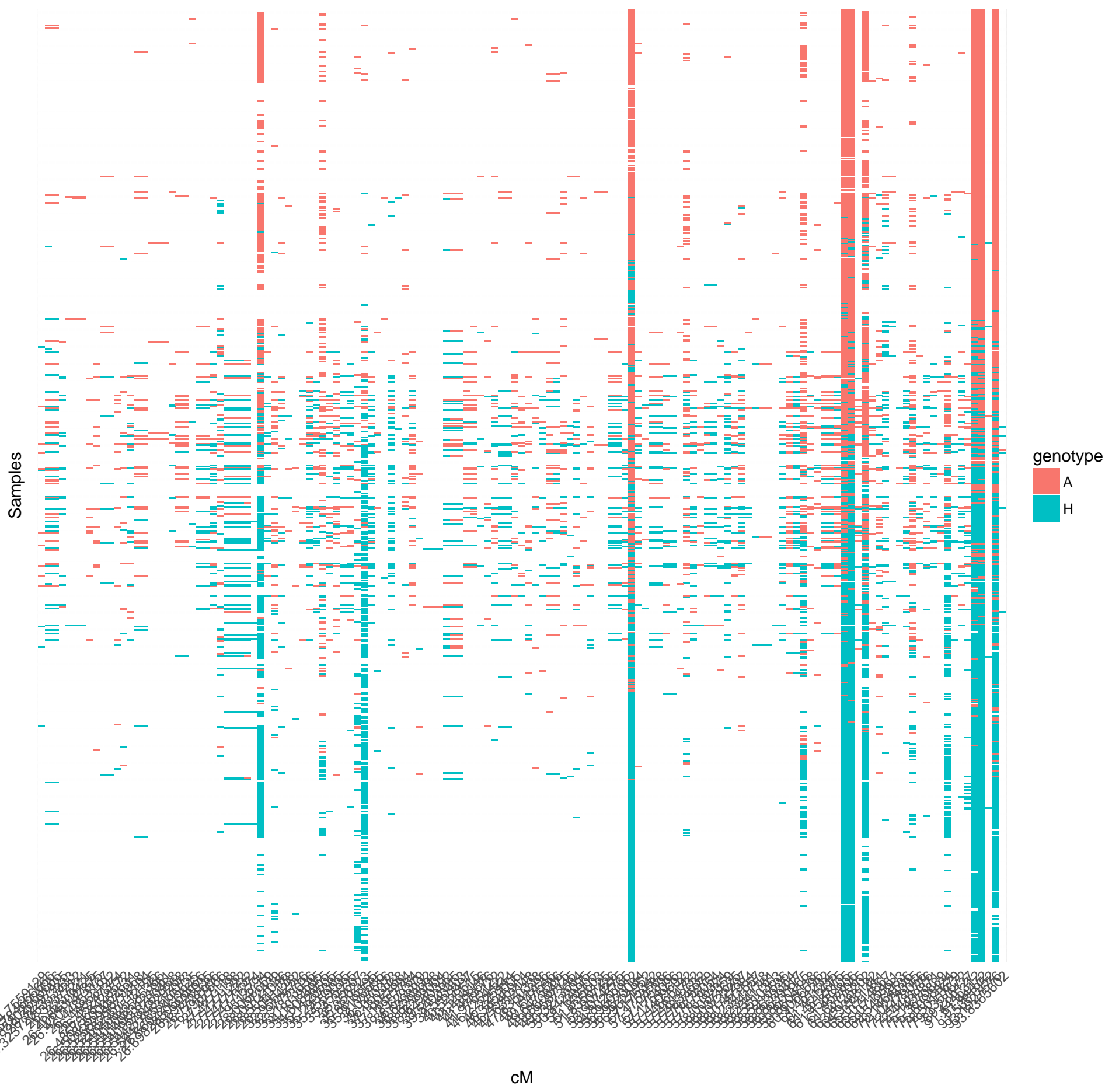
This figure displays a complex genomic visualization, likely a heatmap or a grid of genomic data. The visualization is organized into a grid of columns and rows, with varying lengths and positions of horizontal bars (red and teal) indicating specific genomic features or variants. The overall layout is dense and structured, typical of a genomic data visualization tool like IGV or a similar software.

A

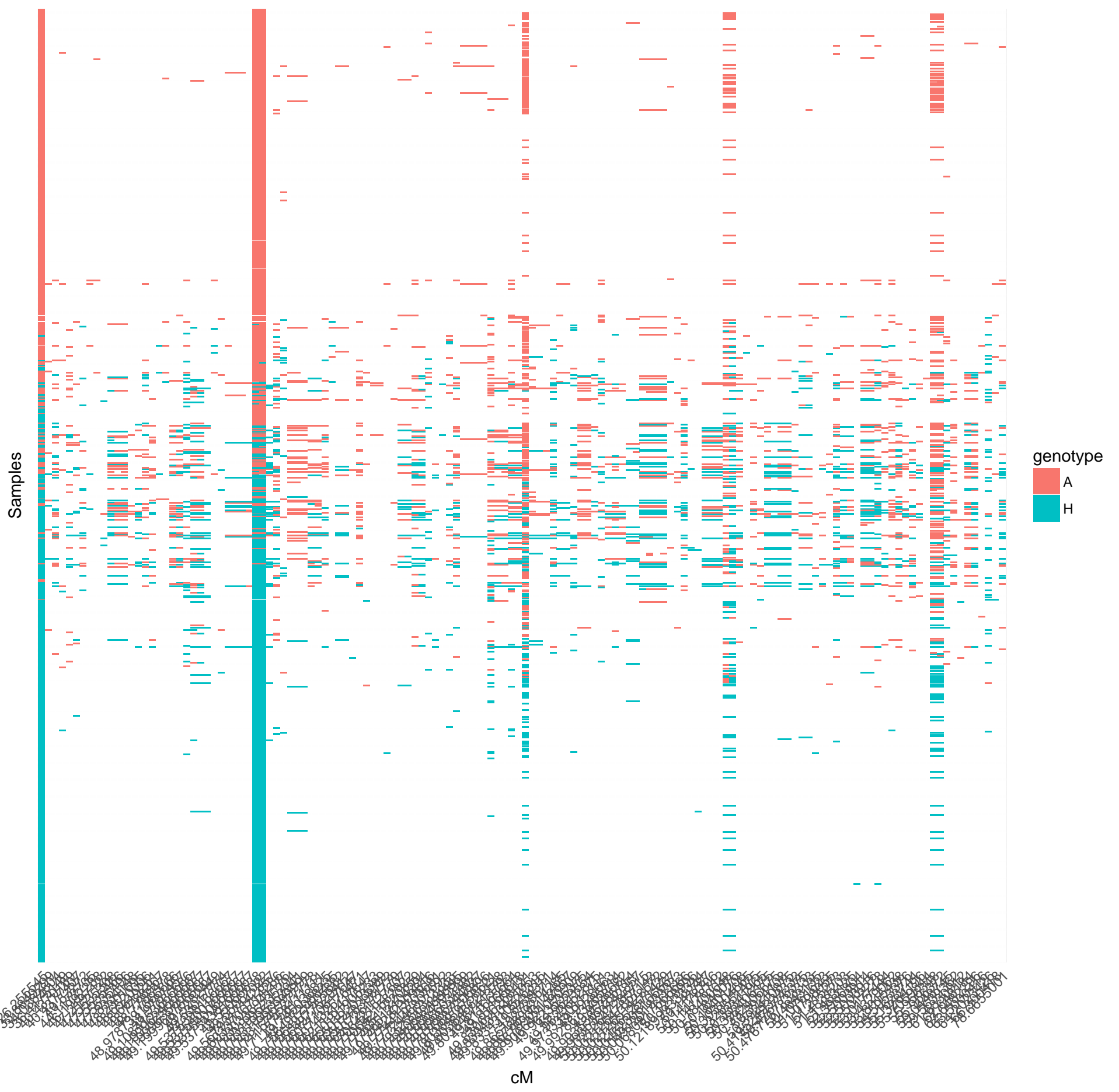
L

cM

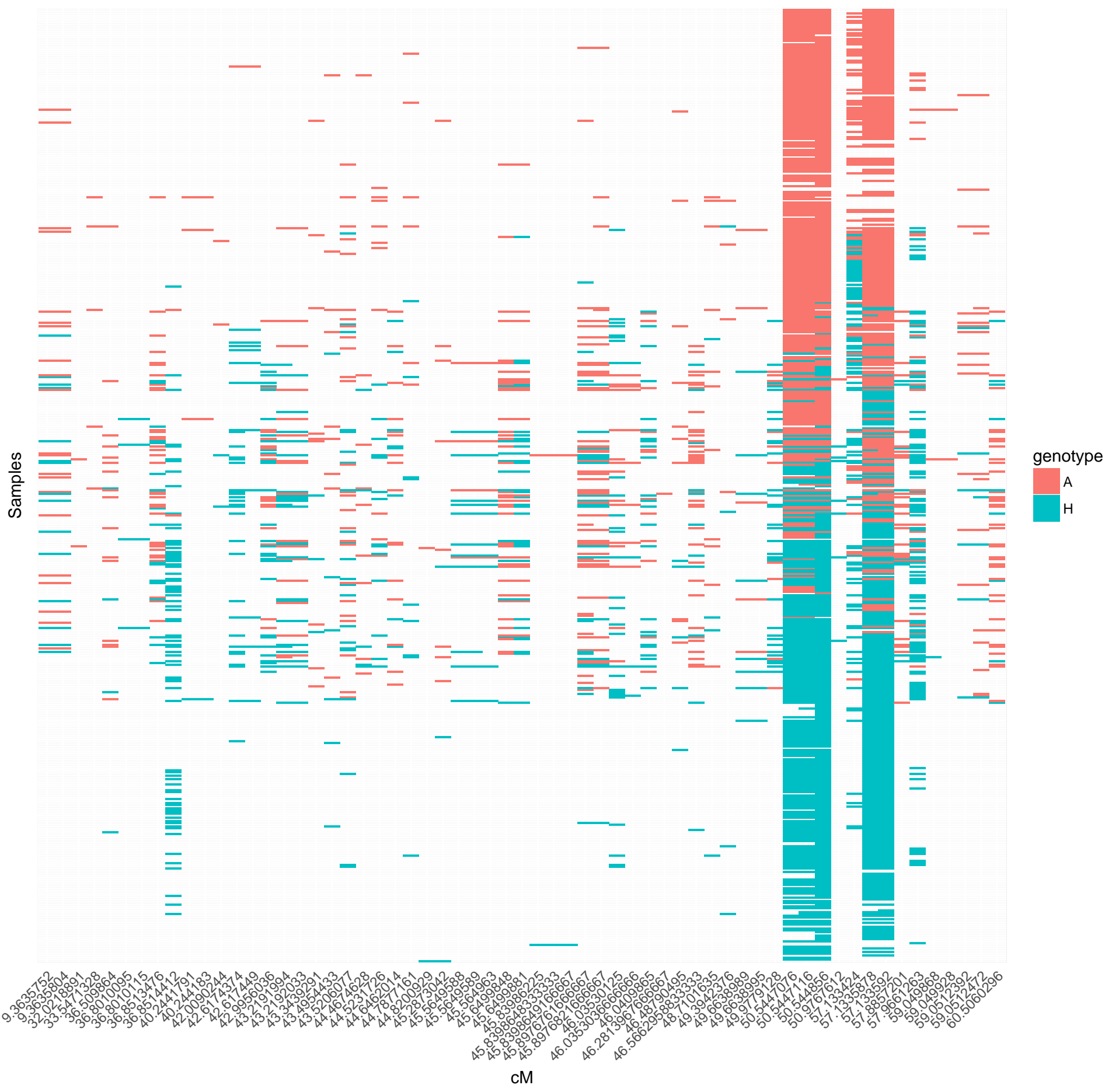
deb BC1, missing data, Chr 09



deb BC1, missing data, Chr 10



deb BC1, missing data, Chr 11



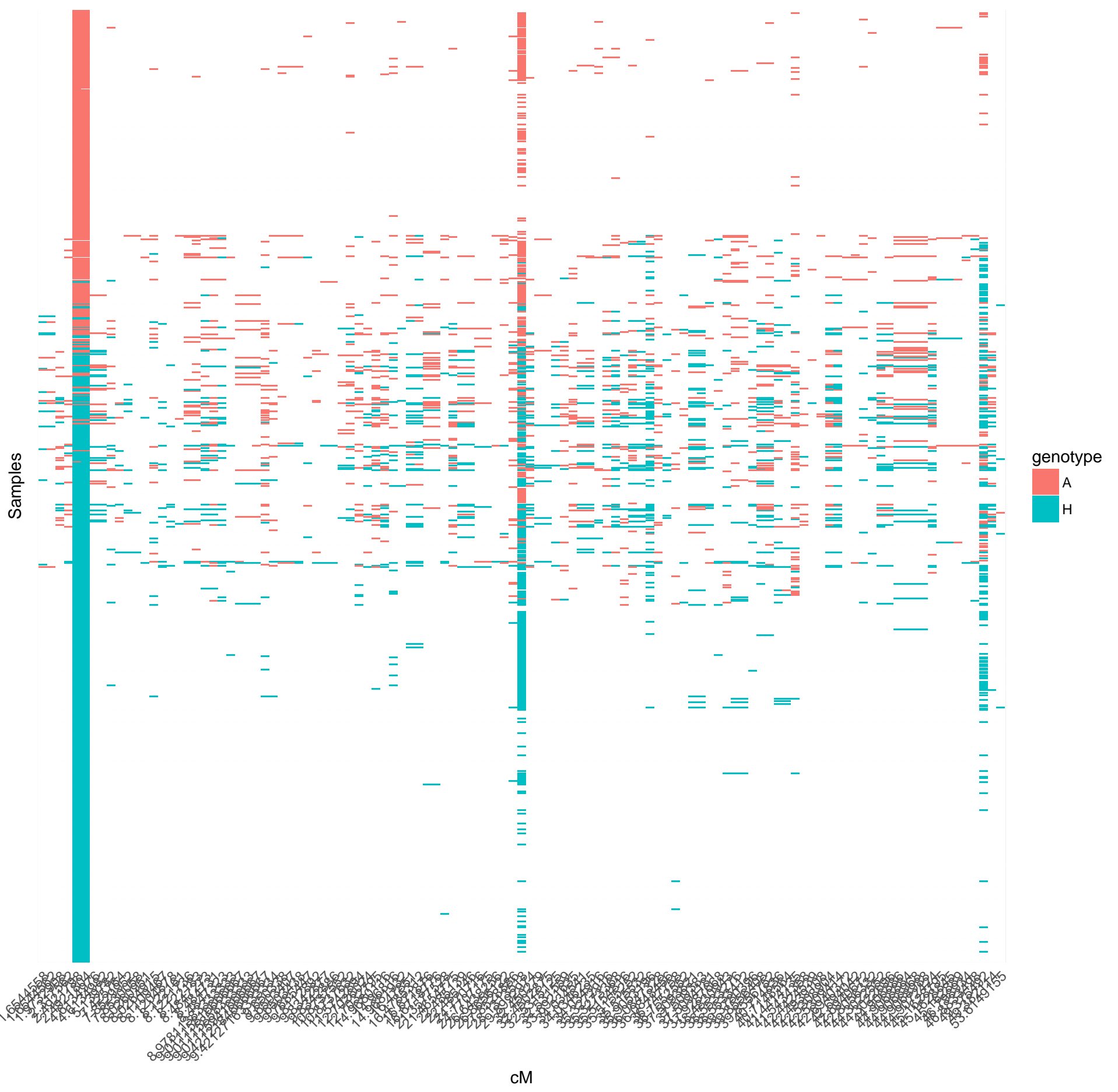
A
H

genotype

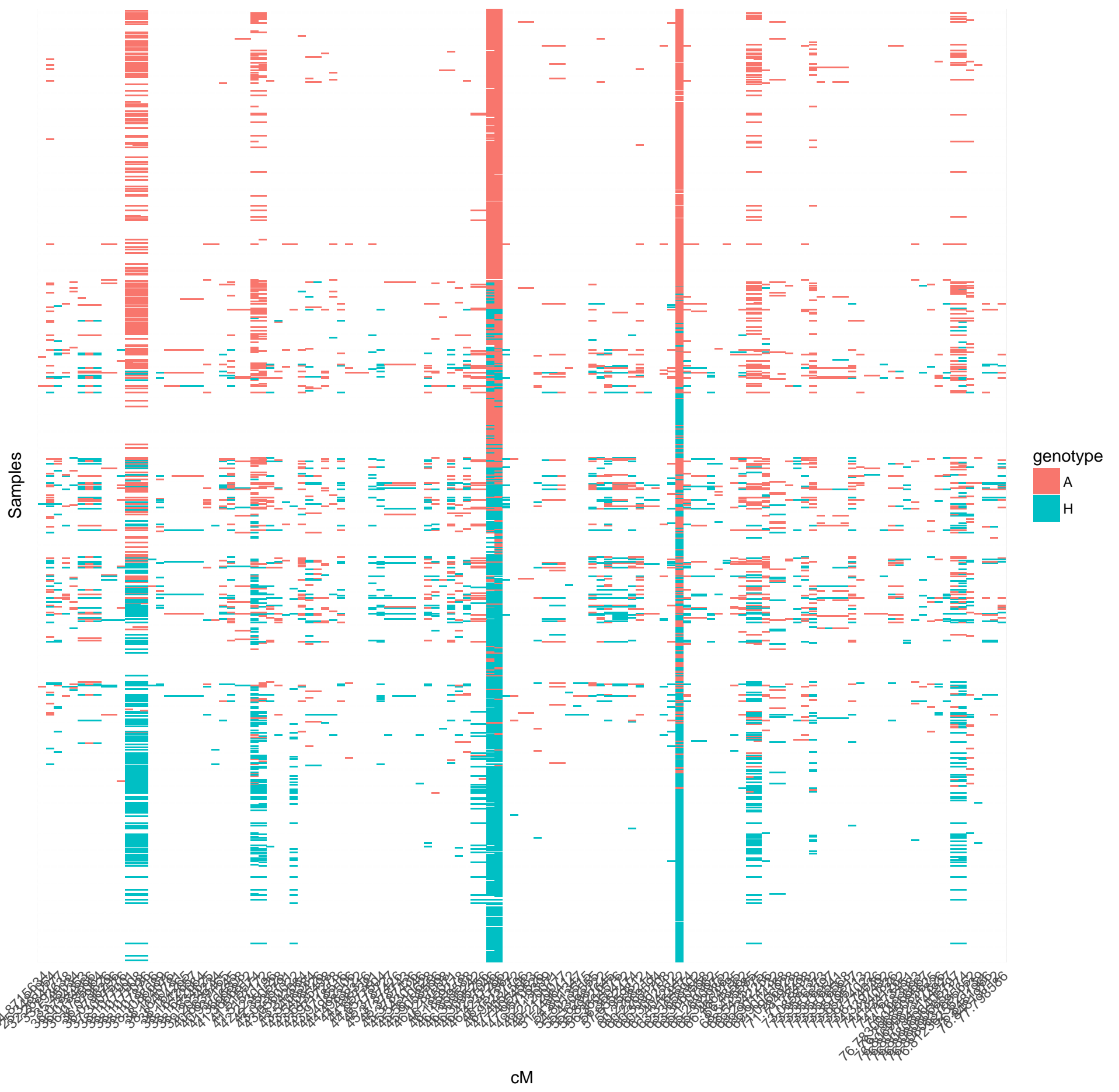
| |
|---|
| A |
| H |

cM

deb BC1, missing data, Chr 14



deb BC1, missing data, Chr 15



A
H

H

100%

cM

A
H