

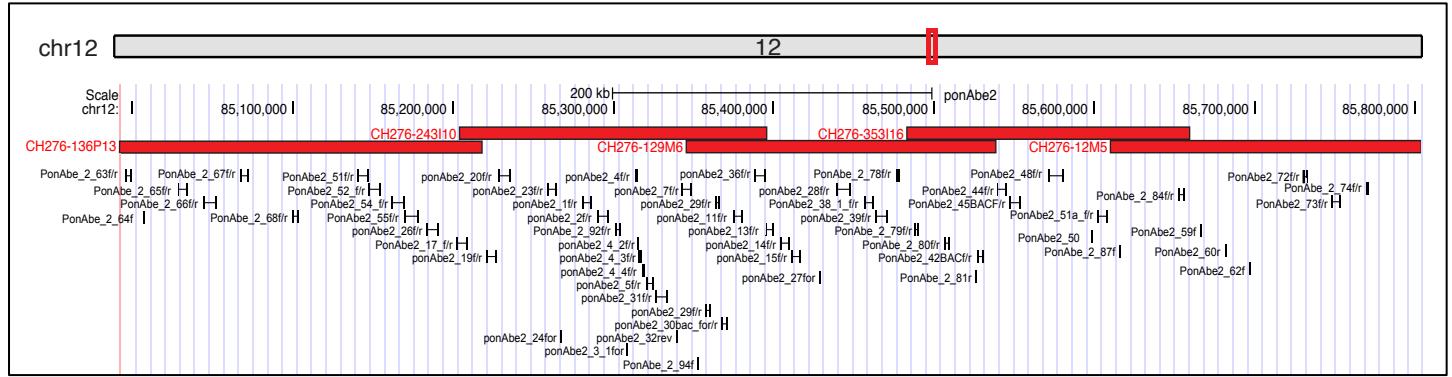
Epigenetic origin of evolutionary novel centromeres

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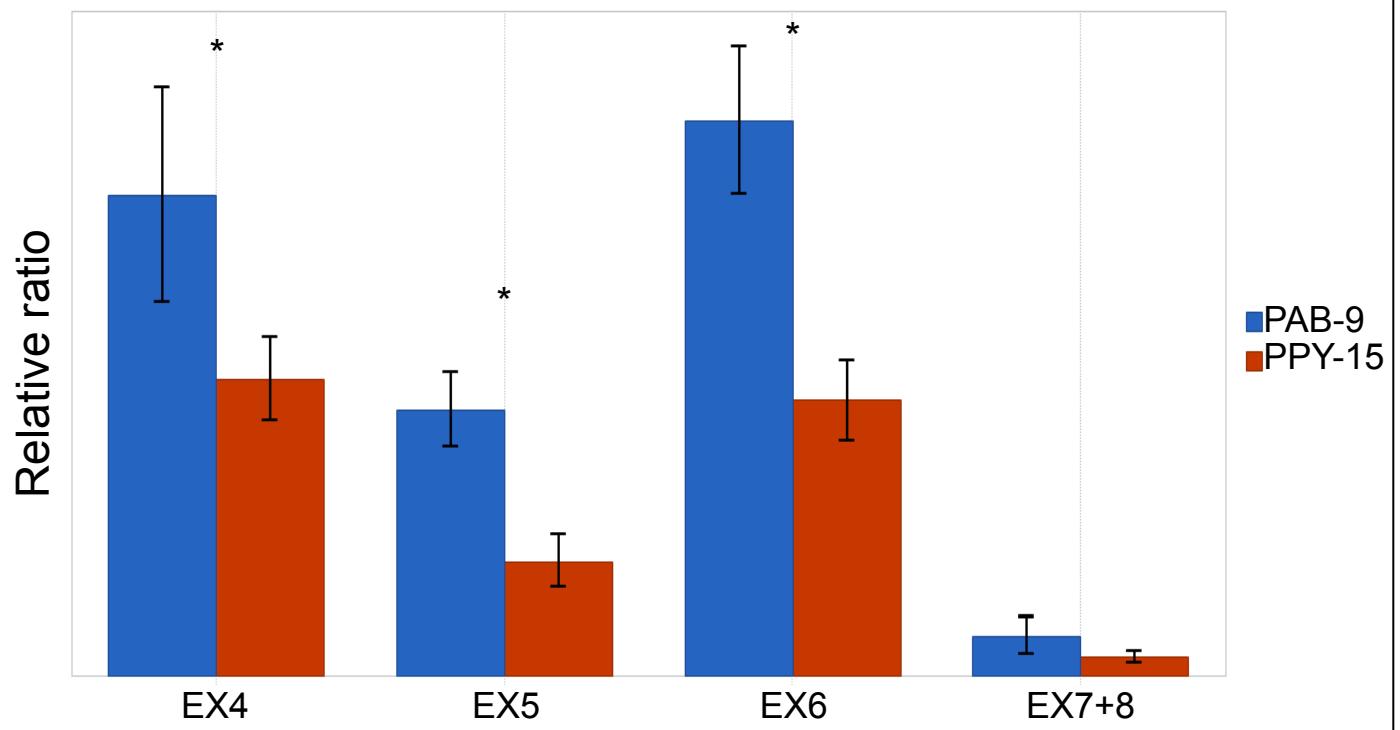
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Supplementary Figure S1. Distribution of Sanger sequenced LR-PCR ends (black vertical bars) with respect to the ponAbe2 sequence of chromosome 12 and to the 5 sequenced BACs of the CH276 library (red rectangles). Some ends are not reported because of their low quality. On the left of each LR-PCR product are reported the corresponding primer names.



Supplementary Figure S2. Exon-specific reverse transcription quantitative PCR analysis of *SLC6A15*, performed in PPY-15 vs PAB-9. The statistical analysis was carried out using the LightCycler® 96 Software 1.1 (Roche). Asterisks indicate statistically significant results ($p < 0.05$).

Supplementary Table S1. Primers used for LR-PCR. Primers designed on

PacBio805 sequence are reported in italic. * = sequence duplicated on PonAbe2 assembly; ✓ = positive result; ✗ = negative result.

primer name	sequence (5'-3')	map position (ponAbe2)
SLC15_4F	TGGAACTCTCTGTGGGTCAA	chr12:85,479,014-85,479,033
SLC15_4R	CCATACACCAATGCTGCCTC	chr12:85,478,984-85,479,003
SLC15_5F	CTCAGTCTTTCAGCAACCCC	chr12:85,478,505-85,478,525
SLC15_5R	TTTCACCAAAGGACACTGATCC	chr12:85,478,476-85,478,497
SLC15_6F	GGGCTTAAACTGGAAGATGACC	chr12:85,476,967-85,476,988
SLC15_6R	CAAGCAAACCATGACCCAGG	chr12:85,476,931-85,476,950
SLC15_7F	TCGCCACATGTTACCCCTA	chr12:85,465,882-85,465,901
SLC15_8R	GCAGCTTCTCTCCAGACCTT	chr12:85,463,057-85,463,076

Supplementary Table S3. Primers used for RT-qPCR of the *SLC6A15* gene.