Table S6. Amplicons used to confirm the molecular organization of the genomic regions corresponding to the breakpoints of the inversion In(2R)51F11-56E2

Amplicon <sup>a</sup>	Primers	Size (nt)	Purpose
A <sup>b</sup>	CB-0236-3up / CB-0236-3down	~700	Verification of flanking sequence at the outer breakpoint in strains $w^{III8}$ and 5-HA-1995
B <sup>b</sup>	5-HA-1995up / 5-HA-1995down	~600	Verification of flanking sequence at the inner breakpoint in strains $w^{1118}$ and CB-0236-3
C <sup>b</sup>	CB-0236-3up / Plac1	~600	Verification of upstream flanking sequence to P{RS3}CB-0236-3 and its derivatives
D <sup>b</sup>	Pry4 / CB-0236-3down	~500	Verification of downstream flanking sequence to P{RS3}CB-0236-3 and its derivatives
E <sup>b</sup>	5-HA-1995up / Plac1	~600	Verification of upstream flanking sequence to P{RS5}5-HA-1995 and its derivatives
F <sup>b</sup>	Pry4 / 5-HA-1995down	~300	Verification of downstream flanking sequence to P{RS5}5-HA-1995 and its derivatives
G <sup>c</sup>	CB-0236-3up / w11678U	~3,400	Diagnostic presence of a reporter gene mini-white with two exons at the outer breakpoint in
			strains CB-0236-3 and REC
G´°	CB-0236-3up / w11678U	~2,100	Diagnostic presence of a reporter gene mini-white with one exon in strains SIM and REV
H <sup>c</sup>	w11678U / 5-HA-1995down	~2,600	Diagnostic presence of a reporter gene mini-white with two exons at the inner breakpoint in
			strains 5-HA-1995, REC, and INV

<sup>&</sup>lt;sup>a</sup> Figure S5 for physical location. <sup>b</sup> Amplified with Takara Taq. <sup>c</sup> Amplified with Takara Ex Taq.