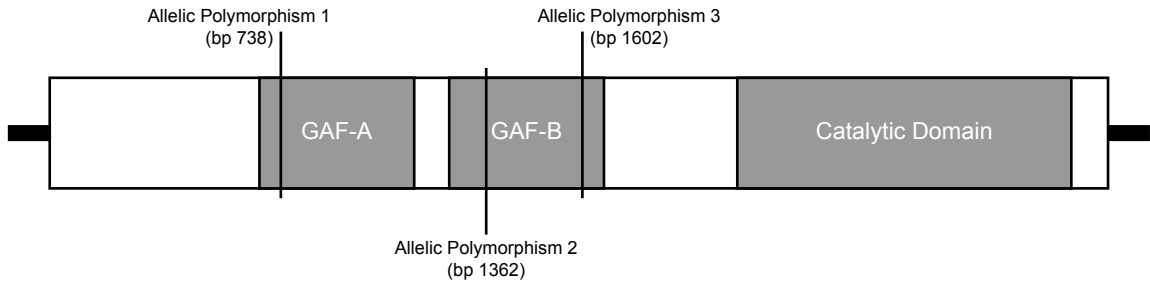


FIG. S1. Schematic diagrams of *TbrPDEB1* and *TbrPDEB2* showing functional domains (grey boxes) and allelic polymorphisms in *T. b. brucei* strain Lister 427 and the Cpd A resistant R0.8 cell line. The ORFs are represented by the largest box (white and grey parts); flanking regions are represented by thick black horizontal lines. Allelic polymorphisms are represented by thin (single nucleotide) or thicker (clusters of multiple nucleotides) vertical lines.

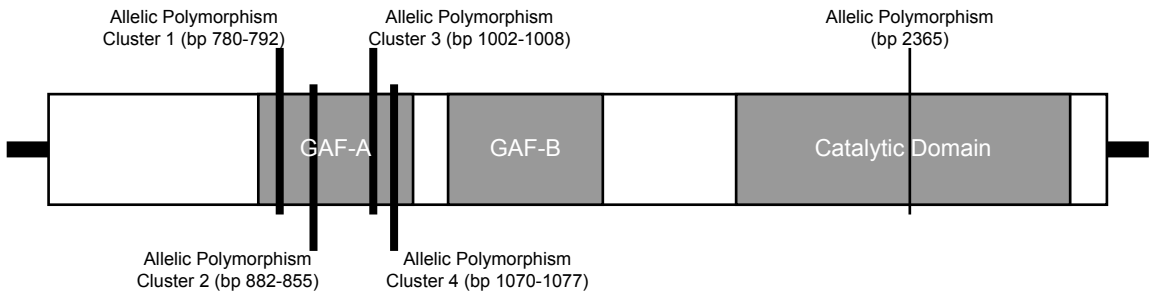
FIG. S2. Western Blot analysis of CARP1 repression (1 μ g/ml tetracycline for 24h, + Tet) in additional *CARP1* RNAi cell lines with *in situ* tagged CARP1 either with an N-terminal 4xTy1 (upper panel) or a C-terminal 3xHA tag (lower panel). CARP1 protein levels were normalized to PFR-A/C detected by the monoclonal antibody L13D6 (36) and set to 100% in the absence of Tet. The relative scan gain in the 800 nm channel was set to 1 for the upper panel and to 3 for the lower panel. Relative expression levels are indicated as percentage of the non-induced cultures.

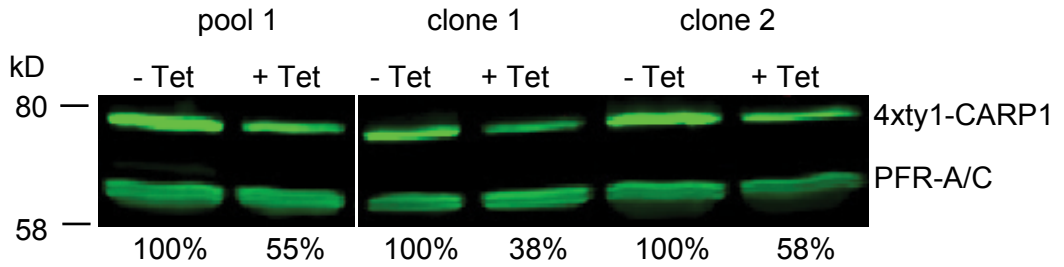
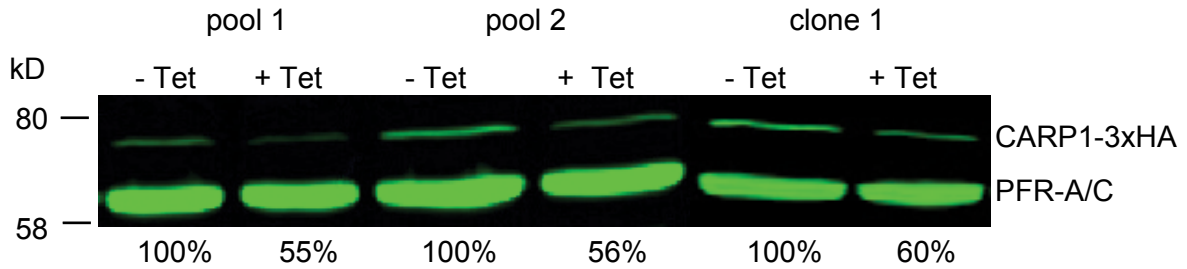
FIG. S3. *CARP1-4* mRNA expression levels in *T. b. brucei* strain Lister 427 wild type (WT) and derived R0.8 cell line in the presence or absence of Cpd A, determined by quantitative real-time PCR. Relative expression normalized to *TERT* as reference gene (37) and the wild type line is given with standard error (SEM) of three biological replicates.

S1 Allelic Polymorphisms of *TbrPDEB1*



Allelic Polymorphisms of *TbrPDEB2*



S2**CARP1 RNAi / p3077 CARP1 (*in situ* N-terminal 4xty1)****CARP1 RNAi / pMOTag2H CARP1 (*in situ* C-terminal 3xHA)**

S3