If you use this data, please cite:

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Fam0a: a-type VSG

Telomeric VSG from bloodstream expression sites are shown here in yellow boxes. These are largely taken from *T. brucei* 427, and the expression site number is given*. Three VSG from *T. brucei* 927 expression sites are also included (marked with a hash #). Telomeric VSG from metacyclic expression sites are shown in orange and labelled with their database accession number.

It is clear from their distribution that telomeric *VSG* are not a distinct subset of all *VSG*, but are instead a random sample of total diversity.

Similarly, two *VSG* taken from a published mini-chromosome sequence in *T. brucei* 927**, and highlighted in pink, cluster among subtelomeric *VSG*, and so do not appear to be structurally distinct.

NOTES:

Fam0a includes the a-type variant surface glycoprotein genes from *T. brucei*. All full-length genes and pseudogenes (n=427) were selected from vsgDB (www.vsgdb.net).

The Bayesian phylogram was estimated from a multiple nucleotide sequence alignment of 2035 characters, using MrBayes with a GTR+ Γ model and default settings. This included the C-terminal domain. The tree is rooted using transferrin-receptor-like sequences from *T. brucei* and *T. congolense*. Selected nodes are supported by posterior probability values.

* Hertz-Fowler et al. (2008). *PLoS One*, **3**(10): e3527. ** Alsford et al. (2001). *Mol. Biochem. Parasitol.* **113**(1):79-88.

