

## pRPa<sup>TAG</sup>

For inducible expression of N/C-terminal human cMYC- (EQKLISEEDL) or N/C-terminal enhanced GFP-tagged proteins in *T. brucei* from a tetracycline-responsive RRNA promoter.

- Derived from pLEW100 (Wirtz et al, 1999) and p2T7<sup>TAbiue</sup> (Alibu et al., 2005).
- High fidelity polymerase recommended.
- Integrates at the tagged RRNA spacer in 2T1/TAG<sup>PAC</sup> *T. brucei* following digestion with *AscI* (Alsford et al 2005), giving a transformation efficiency of  $\sim 2.5 \times 10^{-6}$  (Alsford & Horn, 2007).

## Cloning

GFP/MYC<sub>X</sub>

To ensure that your gene is in frame with the tag, place the second codon downstream of the *XbaI* site, i.e. TCTAGA:[codon 2]:[codon]<sub>n</sub>:[stop]:GGATCC.

X<sup>GFP</sup>

To ensure that your gene is in frame with the tag, place the last but one codon upstream of the *XbaI* site, i.e. AAGCTT:[start]:[codon]<sub>n-1</sub>:TCTAGA.

X<sup>6MYC</sup><sub>X</sub>

**N-terminal tagging:** clone your gene without a start codon via *AvrII*/*BamHI* digestion (or without start/stop codons via *AvrII*)  
i.e. CCTAGG[codon 2]:[codon]<sub>n</sub>:[stop]:GGATCC.

**C-terminal tagging:** clone your gene without a stop codon via *HindIII*(or *PacI*)/*XbaI* digestion (without start/stop codons via *XbaI*)  
i.e. AAGCTT:[start]:[codon]<sub>n</sub>:TCTAGA.

Use *HindIII* / *BamHI* if you don't want the tag.

There are alternatives if the gene contains *XbaI*, *AvrII* and/or *BamHI*:

<b>Plasmid</b>	<i>XbaI</i> , <i>AvrII</i> <i>BamHI</i>	<b>Insert</b>	<i>AvrII</i> , <i>NheI</i> , <i>SpeI</i> , <i>XbaI</i> <i>BglII</i>
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## Key features

- Complete sequences available.
- Hygromycin for stable selection.
- All vectors allow inducible expression using tetracycline (or analogues).
- Inducible cassette is independent of selectable marker.
- Modular nature allows tag or other components to be exchanged.
- Compatible with *T. brucei* cell lines expressing TetR and containing a tagged RRNA spacer, e.g. 2T1/TAG<sup>PAC</sup> (Alsford et al, 2005).

Upon integration into *T. brucei*, the construct replaces TAG<sup>PAC</sup> and generates a functional *HYG<sup>R</sup>* at the previously tagged RRNA spacer. The operator binds Tet-repressor in the absence of tetracycline so the inducible RRNA promoter is activated and tagged protein is expressed when tetracycline ( $1 \mu\text{g ml}^{-1}$ ) is added to the medium.

## Detection:

cMYC      Mouse anti-cMYC, 9E-10 (Santa Cruz; IFA / western blotting)  
            Mouse anti-cMYC, 4A6 (Upstate Biotechnology; western blotting only, as binds the mitotic spindle in *T. brucei*)

eGFP      Rabbit anti-GFP, IgG fraction (Molecular Probes; IFA and western blotting)

Other questions/comments, contact David Horn ([david.horn@lishtm.ac.uk](mailto:david.horn@lishtm.ac.uk)).