## **ERRATUM**

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## A reverse genetic approach for generating gene replacement mutants in *Ustilago maydis*

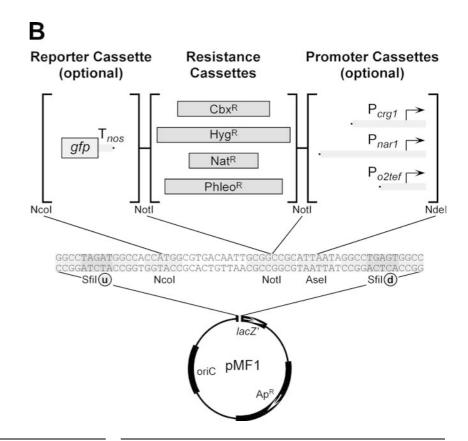
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The plasmid backbone of plasmid pMF1 in Fig.1B was inadvertently depicted the wrong way round. The corrected version is shown below.

Fig. 1 B Schematic representation of the corresponding plasmid system. The optional gfp reporter gene cassette is depicted on the left.  $T_{nos}$  indicates the transcriptional terminator of the nopaline synthase gene from Agrobacterium tumefaciens (Bevan et al. 1983). In the center, mandatory resistance cassettes conferring resistance to carboxin (Cbx<sup>R</sup> hygromycin (Hyg<sup>R</sup> hygromycin (Hyg<sup>k</sup>), nourseothricin (Nat<sup>R</sup>) or phleomycin (Phleo<sup>R</sup>) are shown. The optional promoter cassettes derived from the arabinase gene ( $P_{crg1}$ ), the nitrate reductase gene  $(P_{nar1})$ and the translation elongation factor gene  $(P_{o2tef})$  are indicated on the right. The linker sequence between the two Sfi I sites and the plasmid structure is depicted (lacZ',  $\alpha$  fragment of  $\beta$ -galactosidase; Ap<sup>R</sup>, Ampicillin resistance; oriC, origin of replication) at the bottom



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