

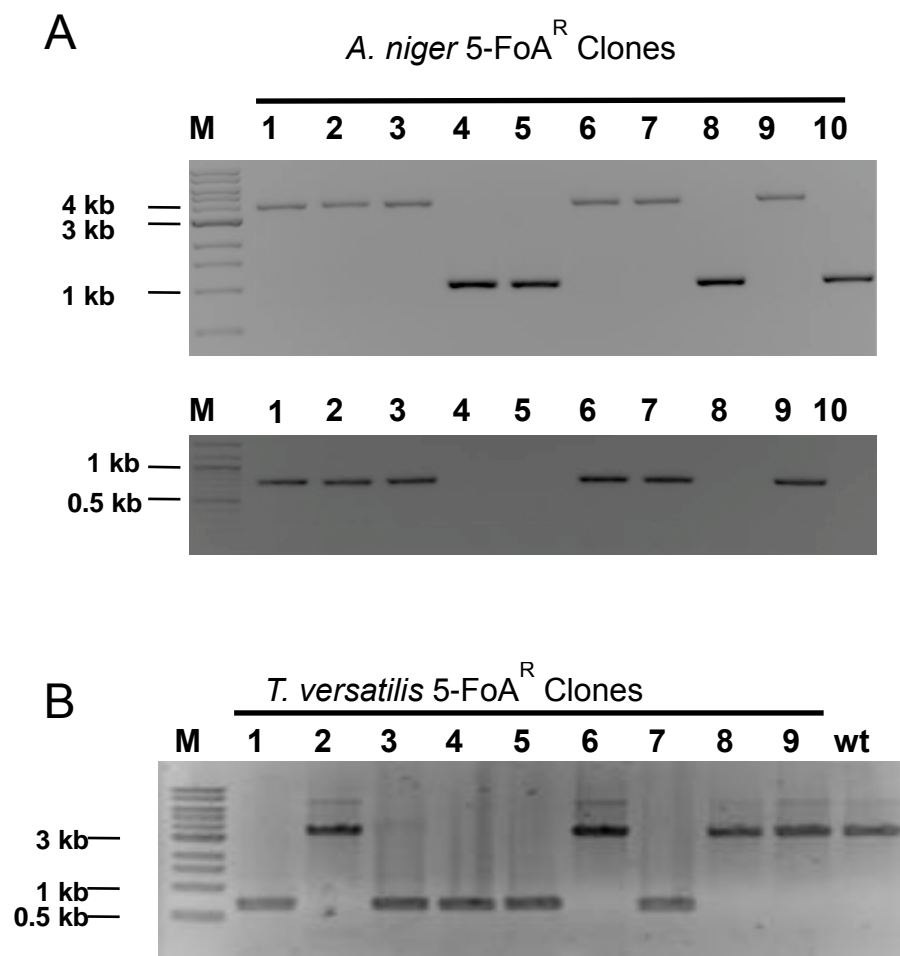
Supplementary Files

Supplementary File 1: Oligonucleotides used in this study

Primer	Sequence (5'-3') ^a	Properties
<i>A. niger</i>		
AnXlnRNotIup	ATGCAT <u>GCGGCCG</u> CTCCCAACTTTATCACTCCC	<i>xlnR</i> deletion upstream external
AnXlnRHindIIIup	ATGCATA <u>AAGCTT</u> GGAATTCGCAAGGAAGTGG	<i>xlnR</i> deletion upstream internal
AnXlnRHindIIIdw	ATGCATA <u>AAGCTT</u> TTTTGCAGTAACACGGCTG	<i>xlnR</i> deletion downstream internal
AnXlnRSpel dw	ATGCATA <u>CTAGT</u> ACTCACGGGATCCACGAAG	<i>xlnR</i> deletion downstream external
AnXlnRScexup	CAGACTGAATCGGCAATGC	Screen <i>xlnR</i> deletion, external
AnXlnRScexdw	GTTTAAAGGAGGGGGTTTGG	Screen <i>xlnR</i> deletion, external
AnXlnRScinup	CACCATCTGAGCTCGCAGCC	Screen <i>xlnR</i> deletion, internal
AnXlnRScindw	GCAGTTTCTGGCATAACG	Screen <i>xlnR</i> deletion, interna
AnXlnRSopr dw	GAGTTTTACCTCTGATGGC	Probe for Southern
AnXlnRSopr dw	ACCGGGTAAAAAAGTCAAG	Probe for Southern
<i>T. versatilis</i>		
TvXlnRSpelup	TTCTCTG <u>CTAGT</u> CGATTCTTCTGG	<i>xlnR</i> deletion upstream external
TvXlnRApaup	TTATAGAAGGGCCCAACCGAAGTCCAC	<i>xlnR</i> deletion upstream internal
TvXlnRApadw	AACGACGGGCCAGGTTCTTGCACTTTATC	<i>xlnR</i> deletion downstream internal
TvXlnRSpedw	AACCTC <u>CTAGT</u> CAGCGTCTTCAGCCG	<i>xlnR</i> deletion downstream external
TvxlnRScexup	CTGTGATTCGGTTGCTTTCTGG	Screen <i>xlnR</i> deletion, external
TvxlnRScexdz	TCGCATCGCTAACAGACAACG	Screen <i>xlnR</i> deletion, external

^a Restriction sites for endonuclease used are underlined.

Supplementary File 2: Screen and confirmation of *xlnR* deletion in *A. niger* and *T. versatilis*. A) *xlnR* deletion in *A. niger*. Ten randomly chosen 5FoA-resistant clones, annotated 1 to 10 were tested by PCR for *xlnR* deletion using external (upper panel) and internal (lower panel) primers to the deletion. “M” line is 1 kb ladder from NEB. Clones 4, 5, 8 and 10 exhibited *xlnR* deletion (i.e. expected 1020 bp band for the PCR with external primer and no band for the PCR with the internal primer). B) *xlnR* deletion strain in *T. versatilis*. Nine randomly chosen 5FoA-resistant clones (annotated 1 to 9) were tested by PCR for *xlnR* deletion using external primers to the deletion. “M” line is 1 kb ladder from NEB and “wt” line control PCR with *T. versatilis* DNA. Clones 1, 3, 4, 5 and 7 exhibited *xlnR* deletion (i.e. expected 673 bp band for the deletion).



Supplementary File 3: Verifying that no extra copies of the plasmid remain in the *xlnR* deleted strain of *A. niger*. Wild-type and *xlnR*-deleted *A. niger* strain genomic DNA was cut with *BglII* and a Southern blot hybridisation performed using the complete plasmid pC3-*An_ΔxlnR* as a probe. Line M represents the 1kb ladder. The expected band for the AB4.1 strain was 6908 bp and for the *xlnR* deletion 4162 bp. Note that a band of ca. 6kb is predicted to be present in both the WT and *xlnR*-deleted *A. niger* strains but was not detected, probably because of the relatively small region of homology to the very large, whole plasmid, probe.

