Table S1. Oligonucleotides used in this work

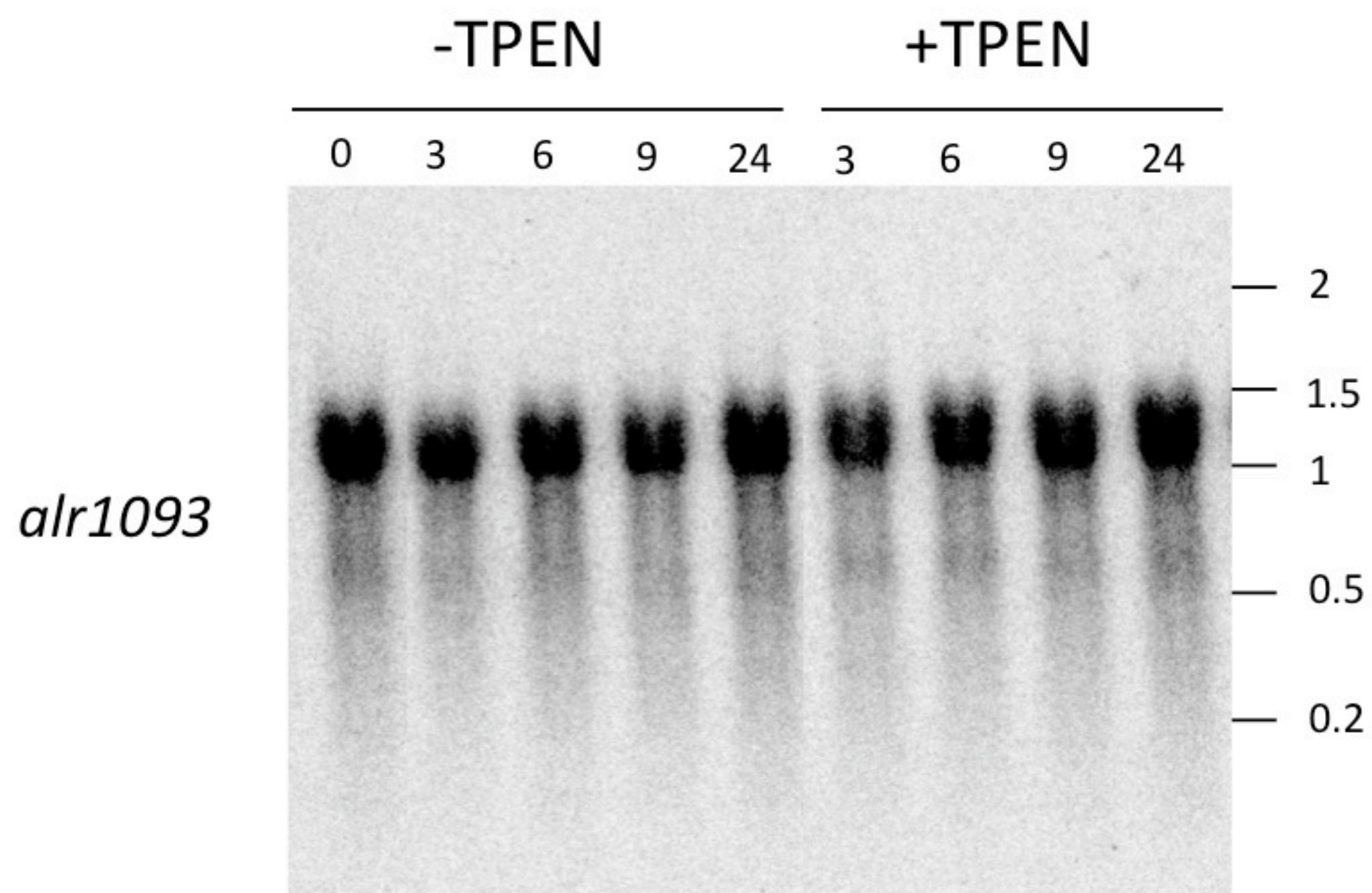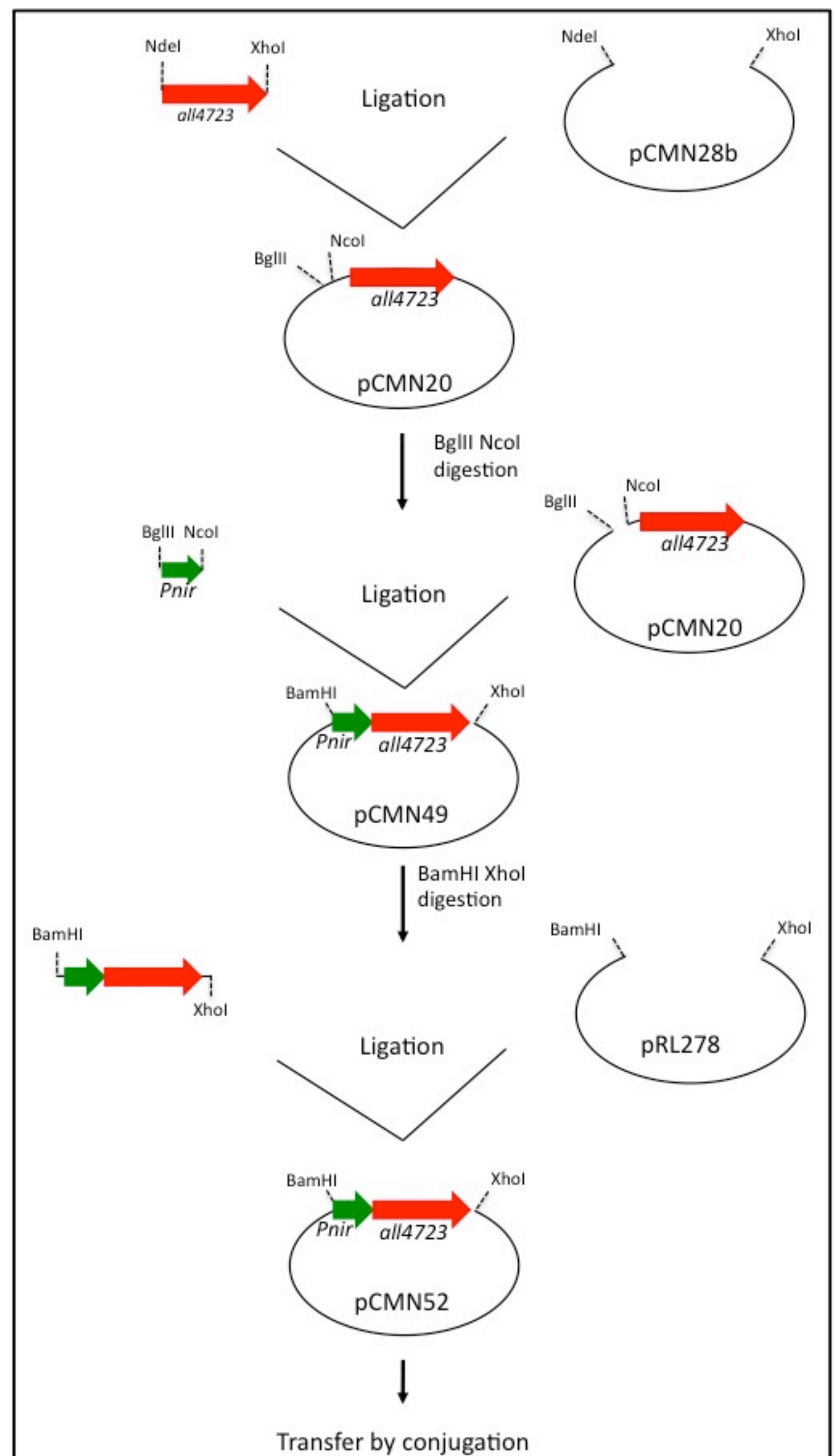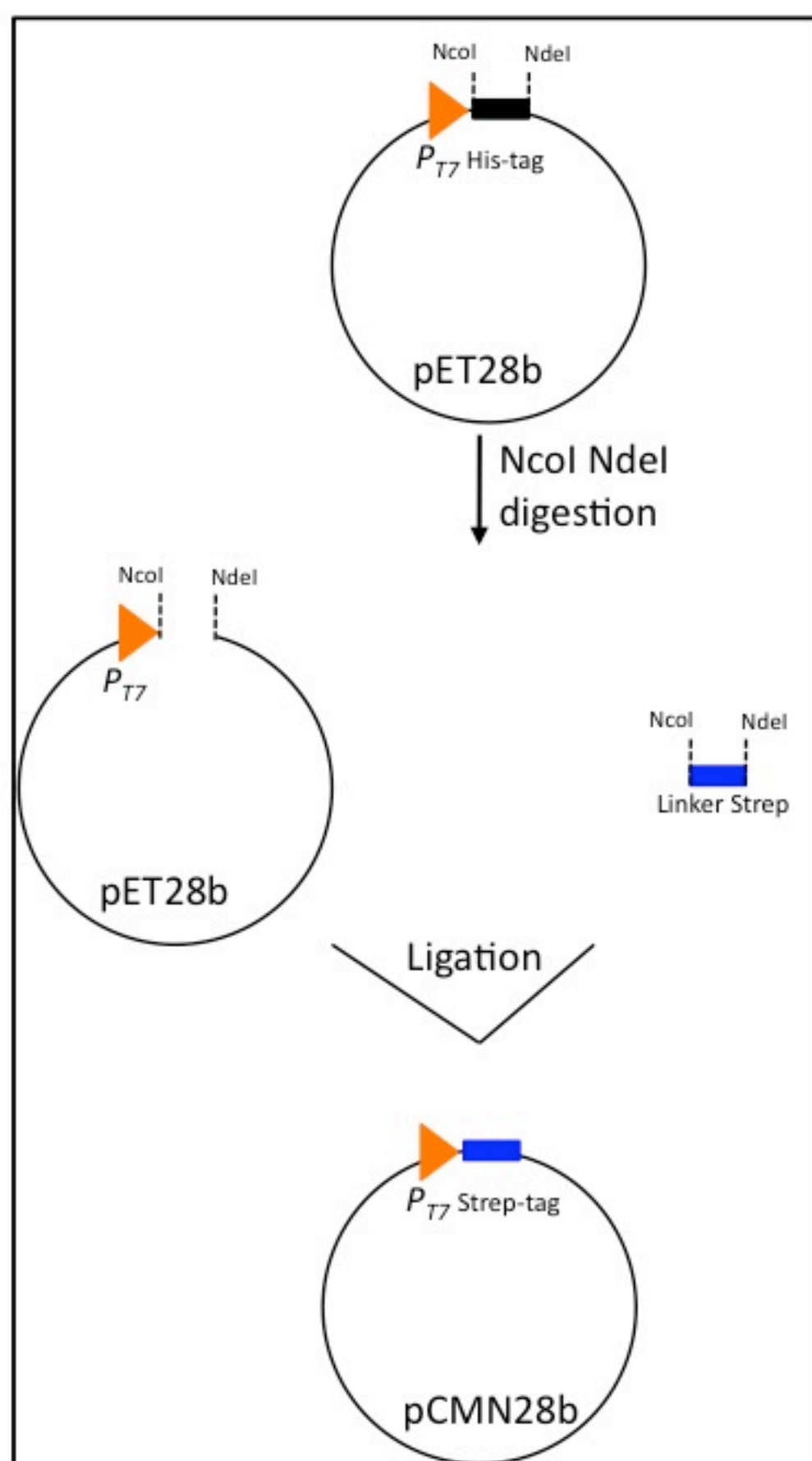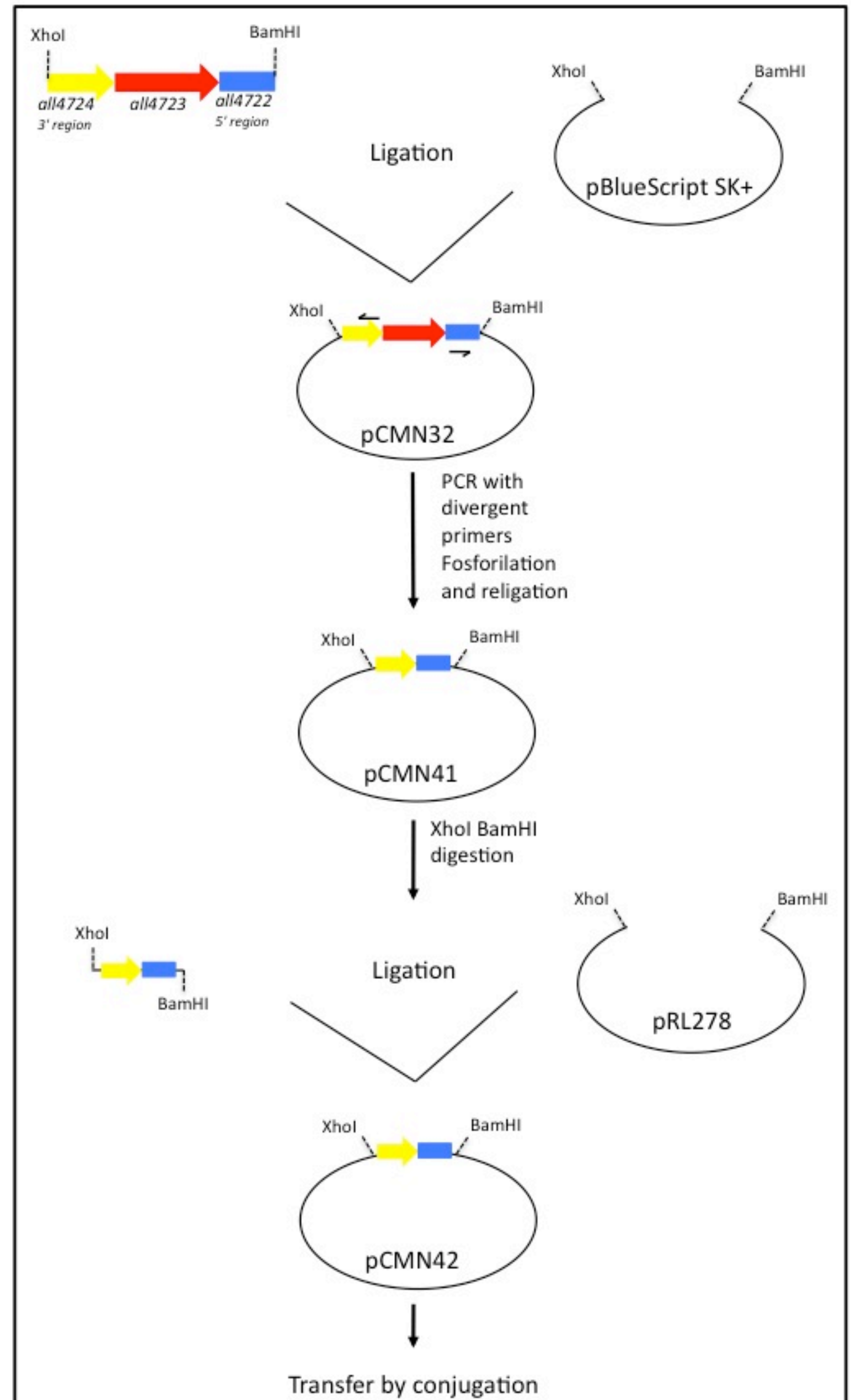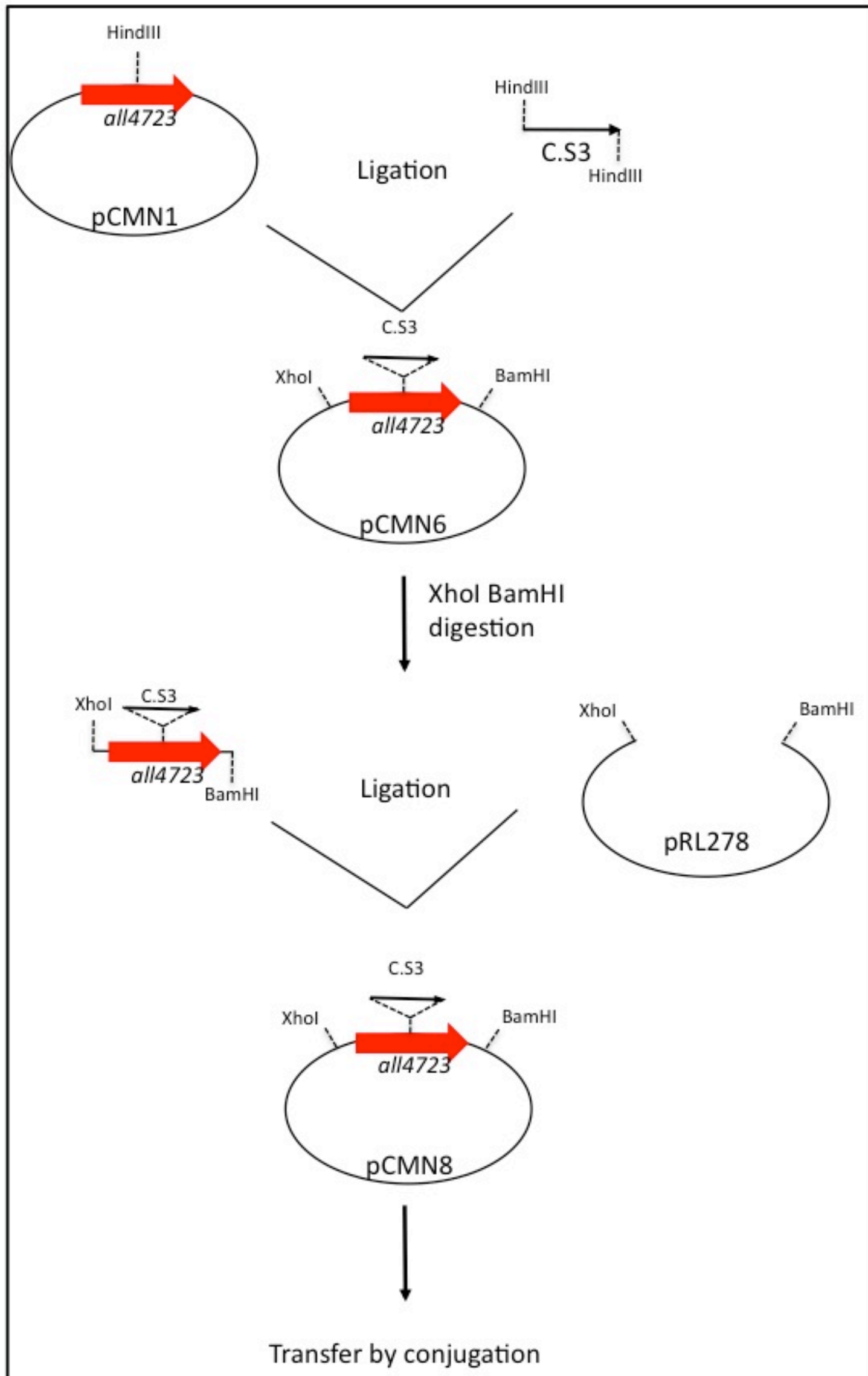| Name	Sequence 5'-3'
A7120 THRS2-1F	TTTATTCATATGGTCAGTTCCTTAAC
A7120 THRS2-1R	CGTATATTCAACTCGAGCAATCTGC
NIR-2F	ACCAAGATCTTTAGCTACTCATTAG
NIR-1R	GTCCCATGGTTTGTGTGAATCCTTATG
24STREP-1F	GTAGCGCTCGAGGTCAAATC
22DEL-1R	TGGCTAGGATCCTCATGATTATG
22DEL-1F	TTTACAAATACAAAATCCTTAAGCAG
24DEL-1R	ATTAATAAATAATGAGAATCGTTATCAT
25R	AAATCATCCACTGTCAGG
23R	GAAATAACTTCTGTACTGCCA
22R	GTTGTTTCTGCGGTTAG
21R	ATAAAGTCATGTTATTTCG
4722LUX-1F	GTAAGAACTCGAGCAAATGGTG
4722LUX-1R	CTGCTTAAGGATCCTGTATTTG
ALL4721-3F	CATACTCGAGCTAAGAACCAATTGG
ALL4721-4R	GTAGGGATCCTGAGTTAGTTAATAG
ALL4725_3F	GATAGTGAGCTCGGTATTGGC
ALL4725_6R	CGGGATCCCTAGCTATGATAC
ALL4725_BOX35_F	CGTTCCTACCCTGAGAAATG
ALL4725_BOX35_R	CATTTCTCAGGGTAGGAACG
ALL4725_BOX10_F	TTCCATGCGCGCAATGGTTA
ALL4725_BOX10_R	TAACCATTGCGCGCATGGAA
ALL4723_3F	AAACCAAGCAGAGCTCTTTCCC
ALL4723_10R	GTGCGGATCCCTTATCGTTG
ALL4723_BOX35_F	ATAGTCATCCTGAGTCCGCC
ALL4723_BOX35_R	GGCGGACTCAGGATGACTAT
ALL4723_BOX10_F	TTTTTGCGCGCGCGCGATAA
ALL4723_BOX10_R	TTATCGCGCGCGCGCAAAAA
ALL4722_6F	GAAATAGAGCTCAGTGGCGAG
ALL4722_8R	GGGAATTTCAGGAATTATG
ALL4722_BOX35_F	AATGATCCTGAGAATCACTTAC
ALL4722_BOX35_R	GTAAGTGATTCTCAGGATCATT
ALL4722_BOX10_F	GAGAAACGCGCGCTGCCAAG
ALL4722_BOX10_R	CTTGGCAGCGCGCGTTTCTC
ALL4721_4F	GTGATAGTGAGCTCCGCCAT
ALL4721_5R	CATCCAAGGATCCATTCTATCC
ALL4721_BOX35_F	CAAATTTACCTGAGACTTAAC
ALL4721_BOX35_R	GTTAAGTCTCAGGTAAATTTG
ALL4721_BOX10_F	TGTTTGGGCGCGCTAAGAAG
ALL4721_BOX10_R	CTTCTTAGCGCGCCCAAACA
CK3-1	CAGAGCAGCCGATTGTCTGTTG
CK3-2	CTGATAAGTGAGCTATTCAC

Fig. S1. Expression of the *alr1093* gene in the absence or presence of TPEN. A Northern blot assay was carried out with RNA from cells cultured in the absence or presence of 20 μM TPEN for the time indicated in hours at the top and an internal DNA probe of the *alr1093* gene. The positions of molecular weight markers are indicated at the right (in Kb).

Fig. S2. Description of the cloning steps followed for the construction of the plasmids used in this work