Table S1. PCR primers used for site directed mutagenesis

Name	Construct	Direction	5′-3′ Sequence[a]
HBx-S25A	S25A	Forward	5′-TCCCGTCAGCGCTGAA**G**CCTGCGGACGATCCCTCTCG-3′
HBx-S25A	S25A	Reverse	5′-CGAGAGGGATCGTCCGCAGG**C**TTCAGCGCTGACGGGA-3′
HBx-S25D	S25D	Forward	5′-TCCCGTCAGCGCTGAA**GA**CTGCGGACGATCCCTCTCG -3′
HBx-S25D	S25D	Reverse	5′-CGAGAGGGATCGTCCGCAG**TC**TTCAGCGCTGACGGGA-3′
HBx-S41A	S41A	Forward	5′-GGGCTGTATCGCCCCCT**G**C**G**CCGTCTGCCGTTCCAGC-3′
HBx-S41A	S41A	Reverse	5′-GCTGGAACGGCAGACGG**C**G**C**AGGGGGCGATACAGCCC-3′
HBx-S41D	S41D	Forward	5′-GGGCTGTATCGCCCCCT**GA**TCCGTCTGCCGTTCCAGC-3′
HBx-S41D	S41D	Reverse	5′-GCTGGAACGGCAGACGGA**TC**AGGGGGCGATACAGCCC-3′
HBx-T81A	T81A	Forward	5′-TCTGCACTCGCATGGAG**G**CCACCGTGACCGCCCCTCG-3′
HBx-T81A	T81A	Reverse	5′-CGAGGGGCGTTCACGGTGG**C**CTCCATGCGACGTGCAGA-3′
HBx-T81D	T81D	Forward	5′-TCTGCACGTCGCATGGAG**GA**CACCGTGAACGCCCCTCG-3′
HBx-T81D	T81D	Reverse	5′-CGAGGGGCGTTCACGGTG**TC**CTCCATCGACGTGCAGA -3′

[a] Bold and underlined, mutagenized nucleotide(s).

Table S2. PCR primers used for cloning in expression vectors

Construct (Vector)	Direction	5′-3′ Sequence[a]	Restriction Site
3xFLAG-HBx (pcDNA3.1-3XFLAG)	Forward	5′-GATCGATCGATC**AAGCTT**ATGGCTGCTCGGTTGTGCTG-3′	HindIII
	Reverse	5′-GATCGATCGATC**GAATTC**TTAGGCAGAGGTGAAAAAGTTG-3′	EcoRI
HBx-GFP (pAcGFP)	Forward	5′-GATCGATCGATC**AAGCTT**ATGGCTGCTCGGTTGTGCTG-3′	HindIII
	Reverse	5′-GATCGATCGATCC**GGTACC**GTGGCAGAGGTGAAAAAGTTGCA-3′	KpnI

[a] Bold and underlined, restriction site.