*Bg* transposon transcription from both strands: two products similar to NFI and SET domain proteins may be involved in transcription and chromatin modulation

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PolII Y<mark>SPTSPS</mark>----PPBg4 29 -SP-SPSTS- 35

<u>Bg transposon sequence and probable Bg-encoded proteins</u>. Analysis of the *Bg* transposon sequence suggests that this mobile element encodes several proteins (designated as PPBg1-PPBg3), described previously (MNL 79:32-35; MNL 80, submitted). The analysis also shows certain regions of *Bg* sequence may form Z-DNA and that *Bg*-encoded proteins have Z-DNA binding properties, indicating a possible autoregulation of this transposon at the transcriptional level (MNL 80, submitted). Structure of all above-mentioned proteins was deduced from the strand of *Bg* transposon containing the two longest ORFs. However, some mobile elements (e.g., the maize *MuDR* autonomous element) are transcribed from both strands (Hershberger et al., Genetics 140:1087-1098, 1995). Further analysis of the other sstrand for *Bg* indicates this mobile element may encode 2 further transcription and chromatin modulation proteins.

An 87 amino acid protein encoded by the second strand of Bg

<u>regulators</u>. The ORF of the second strand of *Bg*, from position 724 to 460 (positions for both strands according to the first strand, the sequence of GenBank accession X56877.1), encodes an 87 amino acid protein, designated hereafter as PPBg4 (Fig. 1a). It is unusually rich in tryptophan (7 residues) and has several PS dipeptide residues.

This protein shows significant similarity with the transcription regulators of the nuclear factor I (NFI) family (Fig. 1a), using CLUSTALW analysis at the European Bioinformatics Institute (http://www.ebi.ac.uk) using default parameters. In the human genome, the promoter sites of NFI and Z-DNA forming regions (ZDRs) are near transcriptional start sites (Champ et al., Nucl. Acids Res. 32:6501-6510, 2004). In the case of the *PPBg4* gene, possible ZDRs are located just downstream of the *PPBg4* gene at positions 120 and 402. In addition, a perfect canonical NFI binding site (5'-TGG(N)<sub>6</sub>GCCAA-3'; Zorbas et al., J. Biol. Chem. 267:8478-8484) is present at position 1775 of the *Bg* sequence; i.e., at -1051 bp upstream on the opposite strand in relation to the translation start site of *PPBg4*. The SP-rich stretch S29-S35 of PPBg4 (SPSPSTS, Fig. 1b) is similar to the SPTSPSYSP motif contained in the NFI transcriptional activation domain (Wendler et al., Nucl.

element is similar to Nuclear Factor I family of transcriptional In the NFI transcriptional activation do
a) PPBg4_1_MRQQLQW <b>S</b> -CAAWRQQPHL <mark>PWRRTCWFWLSPSPSTS</mark> 35 NFI_109_MEEDVDTSPGGDYYTSPNSPTSSSRNWTEDIEGGISSPVKKTEMDKSPENSPSQDSPRLS169
PPBg4_36CCSRGLA-TPRGTPQTDLHVNEVAVSWSLPEPSSTLI47 NFI170 SFTQHHRPVIAVHSGIARSPHPTSALHFFATPLPQTASTYFPHTAIRYPPHLNPQDPLKD 230
PPBg4 48 IWEMELLSRRVADGDG 87 NFI 231 IVSLACD ATQQPGPSWYLG 250
b) NFI - <mark>SPTSPSYS</mark>

Figure 1. (a) Alignment of PPBg4 with the transcription factor NF I of *Mus musculus* (GenBank accession AAK21332.1). The T23-S35 sequence similar to the P-4 peptide (Fujii et al., 2003) is underlined. (b) Similarity of the S29-S35 region of PPBg78 with the SPTSPSYSP motif of NFI transcription factor (NFI) and with repeat YSPTSPS of the RNA polymerase II (PoIII). Identical residues are shown in a black background, similar ones are in a grey background.

PPBg5	1	MSCTGLPCNVWIQS <mark>NELSTCLLIVGPIICNNLNDILHPNLINNHLSN</mark> T	48
CAG25109.1	3427	NNMNNMNNTMNNIMNNMNNTMNIMN	3459
PPBg5	49	IINKFCILNTTSCIYRLVKKHPSTAIYHEINNAHHGRT 86	
CAG25109.1	3460	INNNNIFNNDVSNNVDMQHKSDQTCIFNS-NNIH 3492	

Figure 2. Similarity of PPBg5 with a SET-domain protein of *Plasmodium falciparum* (GenBank accession CAG25109.1). Identical residues are shown in a black background, similar ones are in a grey background.