Identification of a novel retrotransposon-like element in *Xenopus laevis* with a developmentally regulated expression

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A novel cDNA clone named 10A1 was obtained by the differential display PCR (DD-PCR) technique. The structural features of 10A1, as reflected in cDNA and genomic clones, fit into the LTR-retrotransposon properties. It contains long terminal repeats (LTRs), primer binding site and polypurine tract. 10A1 LTRs are bounded by 6 bp inverted repeats, and can be subdivided into U3, R and U3 region. Also multiple copies of 10A1-related element are present in the Xenopus genome, and in vitro synthesized 10A1 complementary RNAs are translated to produce a predicted size of protein. 10A1 open reading frame (ORF) encodes the leucine zipper motif capable of forming the coiled-coil as well as CCHC motif conserved in retroviral gag proteins, raising the possibility that 10A1 may produce ribonucleoprotein particles that can mediate retrotransposition. However, no amino acid homology to usually conserved retroviral pol gene was revealed, suggesting that 10A1 adds to a novel family of LTR-retrotransposon-like element in Xenopus. We also show that zygotically activated 10A1 transcripts are restricted to ventro-posterior specific regions and induced by ventralizing manipulations.