A new hobo, Ac, Tam3 transposable element, hopper, from Bactrocera dorsalis is distantly related to hobo and Ac

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Received 24 June 1996; revised 22 August 1996; accepted 23 August 1996; Received by A. Benardi

Abstract

A new transposable element from the hobo, Ac, Tam3 transposon family was isolated as a genomic clone from the oriental fruit fly, Bactrocera dorsalis. It is approximately 3.1 kb in length with 19-bp inverted terminal repeat sequences having a single mismatch. Though sharing several amino acid sequence identities with other hAT elements, it is distantly related to both hobo and Ac. Among hAT elements thus far described in insects, it is apparently the most distantly related to hobo.

Keywords: Transposons; hobo; Ac; Tam3 family; Transposase; Oriental fruit fly; Tephritidae; Gene vector

1. Introduction

Transposable elements appear to be ubiquitous genomic components of most organisms, with many sharing common ancestors (see Berg and Howe, 1989). Determining the organismal range of specific transposon families and elucidating the structural and functional relationships of their constituent elements is important to understanding their evolution, transmission, and regulation. This information is, in turn, important to developing their potential use as gene transfer vectors, among other types of genetic manipulation.

Recently, a structural relationship has been established among transposons found in both plant and insect species. The hobo transposon from Drosophilia melanogaster was shown to have amino acid sequence similarity to the Activator (Ac) transposable element from maize, with a more distant relationship to the Tam3 element from the snapdragon (Calvi et al., 1991). Utilizing some of the common amino acid sequences between Ac and hobo, gene amplification strategies have allowed the subsequent identification of a complete hobo-related element, Hermes, from Musca domestica (Warren et al., 1994) and Hernitz, from Lucilia cuprina (Coates et al., 1996). Discrete hobo-Ac related sequences have been identified as well from Heliothis virescens (DeVault and Narang, 1994), M. vetustissima (Warren et al., 1995), and several tephritid fruit fly species (Handler and Gomez, 1996). These elements are now considered to be members of the hobo, Activator, Tam3 (hAT) transposon family. Of the known complete elements from insects, all exhibit at least a 50% amino acid sequence identity to hobo within their transposase coding regions, while identity to Ac is no greater than 25%. This suggests a clear divergence between these elements in plants and animals. However, translation of the hAT polymerase chain reaction (PCR) sequences from two tephritid fruit flies, Anastrepha suspensa and Bactrocera dorsalis, indicate an equally distant relationship to both hobo and Ac (Handler and Gomez, 1996). To better define the relationships and possible interactions among hAT elements, we have begun to isolate the complete elements from these tephritid species, and here we describe one from B. dorsalis which we designate as hopper.

2. Experimental and discussion

2.1. Cloning and sequencing

Utilizing a hobo-Ac PCR sequence fragment from B. dorsalis (Bd-HRE; Handler and Gomez, 1996) as a probe, we isolated several genomic clones from a λGEM12 (Promega) B. dorsalis Kahuku strain genomic
library. The KahuK strain is a wild strain collected originally in 1986 in Oahu, Hawaii, and is maintained at the University of Hawaii. EcoRI and BamHI restriction fragments of the genomic clones Bd3–2 and Bd6–1 were sequenced after subcloning into pUC19. Sequencing primers were initially derived from the Bd-HRE sequence, with subsequent primer sequencing accomplished in both directions and on both strands. Sequences from the two genomic clones were identical, and the same element was also present within another clone based on adjacent chromosomal DNA sequence. Inverted terminal repeat (ITR) sequences were identified by alignment of complementary distal sequences, which yielded 19-bp inverted repeat sequences (below) containing a single mismatch nucleotide (lower case, bold).

\[ \text{cctgatag/TTGTTGGG} \text{A} \text{ACTATCA} \text{GACAGA} \text{AACTTATTTTACATTCTCATGCTAGT} \text{TTTCAAGAGATACATTCTATTTGTITTT} \text{CTATTCCATGACTTTT} \text{TT} \]

\[ \text{3'--ITR} \text{TTGTTGGG} \text{A} \text{ACTATCA} \text{GACAGA} \text{AACTTATTTTACATTCTCATGCTAGT} \text{TTTCAAGAGATACATTCTATTTGTITTT} \text{CTATTCCATGACTTTT} \text{TT} \]

Interestingly, the chromosomal insertion site (IS) DNA adjacent to the termini of this element does not consist of an 8-bp direct repeat (in lower case), which was verified in another genomic clone only having its termini and insertion sites sequenced. A 50% identity does exist between the 8-bp sequences present.

2.2. Transposase coding region analysis

As shown in Fig. 1 the terminal sequences encompass a 3120-bp element which encodes two large open reading frames (ORF) having a 10-bp overlap (starting at the first ATG, extending from base positions 856 to 1572 and 1563 to 2354) as well as additional smaller ORFs. Comparison of all ORFs to GenBank sequences using BLAST (Altschul et al., 1990) identified highest homology to other hAT elements in the first two ORFs as well as three small C-terminal ORFs, two of which overlap. A consensus reading frame containing these ORFs is realized after introducing two framshifts between the first and second, and third and fourth ORFs yielding a 1.9-kb coding sequence with three internal stop signals. The specific positions of the framshifts and stop codons are shown in Fig. 1, as well as the amino acid sequence motifs having highest homology to those in other hAT elements. Utilizing the conceptual translation this consensus ORF for molecular comparisons, GAP analy-
sis (Devereux et al., 1984) yielded almost the same identity of 24% and similarity (identities plus positive substitutions) of 44% to both hobo and Ac. While this level of homology is not a strong indication of relatedness, the BLAST search of the entire GenBank (release 91) indicated highest amino acid sequence similarity to the hobo transposase \( P_{\text{BLAST}} = 7.4e^{-26} \), and interestingly, secondly to a cosmid sequence from the Nematode Sequencing Project \( P_{\text{BLAST}} = 1.3e^{-25} \). The same sequence was recently discovered and defined as a hAT element based on searches with hobo (Bigot et al., 1996). The third closest homology was to Ac \( P_{\text{BLAST}} = 1.2e^{-24} \). Succeeding high scoring homologies were solely to other hAT sequences including Hermes \( P_{\text{BLAST}} = 7.7e^{-21} \), Hermit from Lucilia cuprina \( P_{\text{BLAST}} = 2.1e^{-18} \), Tam3 \( P_{\text{BLAST}} = 1.1e^{-10} \), a Musca vetustissima hobo-related sequence \( P_{\text{BLAST}} = 0.0012 \), and an Ac-related element in pearl millet \( P_{\text{BLAST}} = 0.046 \).

BLAST analysis alignments indicated strongest overall homologies in C-terminal sequences (or region 3; see Calvi et al., 1991) beginning with the ERVFS sequence, though discrete amino acid sequence identities are clearly apparent in regions 1 and 2, including those used for initial PCR screens (TIDMWT and TRWNS sequences; see Handler and Gomez, 1996). The HTAQNIA sequence is identical in C. elegans-hobo, and the GANIKKA sequence is identical in hobo. A specific sequence identity revealed in this analysis has LDPR as a core (Fig. 2), and is thus far observed only in those hAT elements most distantly related to hobo, including the C. elegans hobo, Ac, Tam3, and hopper.

3. Conclusions

We have described a new transposable element in B. dorsalis, which shares common features with other short inverted terminal repeat elements in the hAT transposon family. Its complete length is approximately 3.1 kb, it is bounded by short inverted terminal repeat sequences having the A2;G5 motif implicated in DNA splicing mechanisms (Warren et al., 1994), and it includes putative transposase-encoding ORFs sharing common sequences with those from hobo, Ac, as well as other hAT transposons. Previous studies with a hopper PCR subfragment showed that hopper exists as a repetitive element in the B. dorsalis genome (Handler and Gomez, 1996), and we confirm here that, thus far, it is the most distantly related insect hAT element to hobo. Of interest is the finding that the element described does not contain an 8-bp insertion site direct repeat which is consistently present in all other hAT elements. Taken together with the sequence comparisons, hopper may be an ancient predecessor to other insect hAT elements, perhaps being a link or early branch from the non-insect elements. Sequence changes due to mutations or recombination may have collected in adjacent and internal sequences, and the transposase ORF discontinuity is not supportive of autonomous function. Previous hybridization studies indicated that five to 10 hopper elements exist in the genomes of several B. dorsalis strains. While it is curious that all of our genomic clones contained the same element, eventual isolation of the other elements will further elucidate the structure and function of hopper.

Acknowledgement

Grateful appreciation is extended to Drs. Susan McCombs and Stephen Saul for providing pupae and to the US Department of Agriculture Competitive Grants Program for support.

References


