

# 4.1 - Segment Graph

Similar to the concept of splicing graphs [Heber 2002], we employ a graph structure for representing the reference transcriptome that is quantified in a non-redundant data structure. Each edge represents a segment of an annotated pre-mRNA molecule by the genomic coordinate of the corresponding 3'-tail and 5'-head position, by the type (*exonic* or *intronic*), and by the set of supporting transcripts (Definition 1).

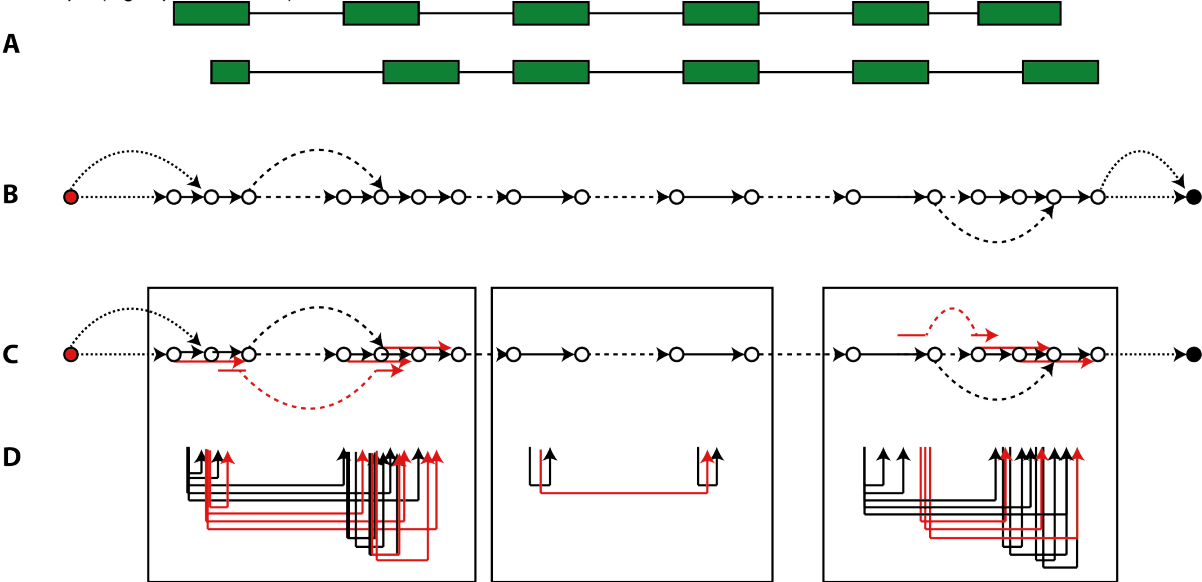
**Definition 1 (Segment Graph Properties):** two adjacent edges in  $G$  are characterized by:

- (i) they share the same intermediary splice site (*adjacency*)

- (ii) they describe the exon-intron structures of all transcripts spanning  $s$  (*completeness*)

- (iii) they either differ in mode or in supporting transcripts (*discrimination*)

To ensure the properties of at the respective transcript edges, all transcription initiation sites are connected to an artificial *source* node, and all cleavage sites are connected to an artificial *sink* node [Sammeth 2008]. Once the segment graph has been constructed for a locus, the edge set  $E$  describes the backbone of exonic segments and introns from the 3'-most transcription start to the 5'-most cleavage site, with additional introns, source and sink links that allow to navigate alternative transcripts (Fig.1, panel A and B).



**Figure 1: segment graph inferred on an alternatively spliced locus.** (A) The exon-intron structure of a locus with two alternative transcripts. (B) Segment graph elements with links by exonic edges shown as solid arrows, links by intronic edges as dashed arrows, and source/sink links as dotted arrows. (C) Expansion of the segment graph by super-edges coalesced from adjacent exon segments or from splice junctions. (D) Super-edges formed by paired-end mappings within the bounds of the three windows marked, to keep (super-) edge combinations within graphical resolution bounds.