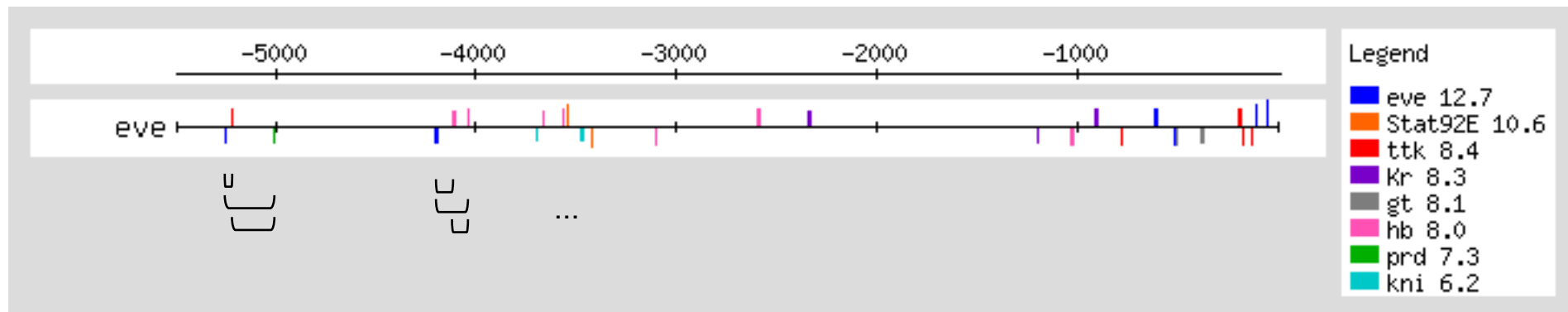


Detecting Cis-Regulatory Modules (CRMs)

Principle of CRM prediction

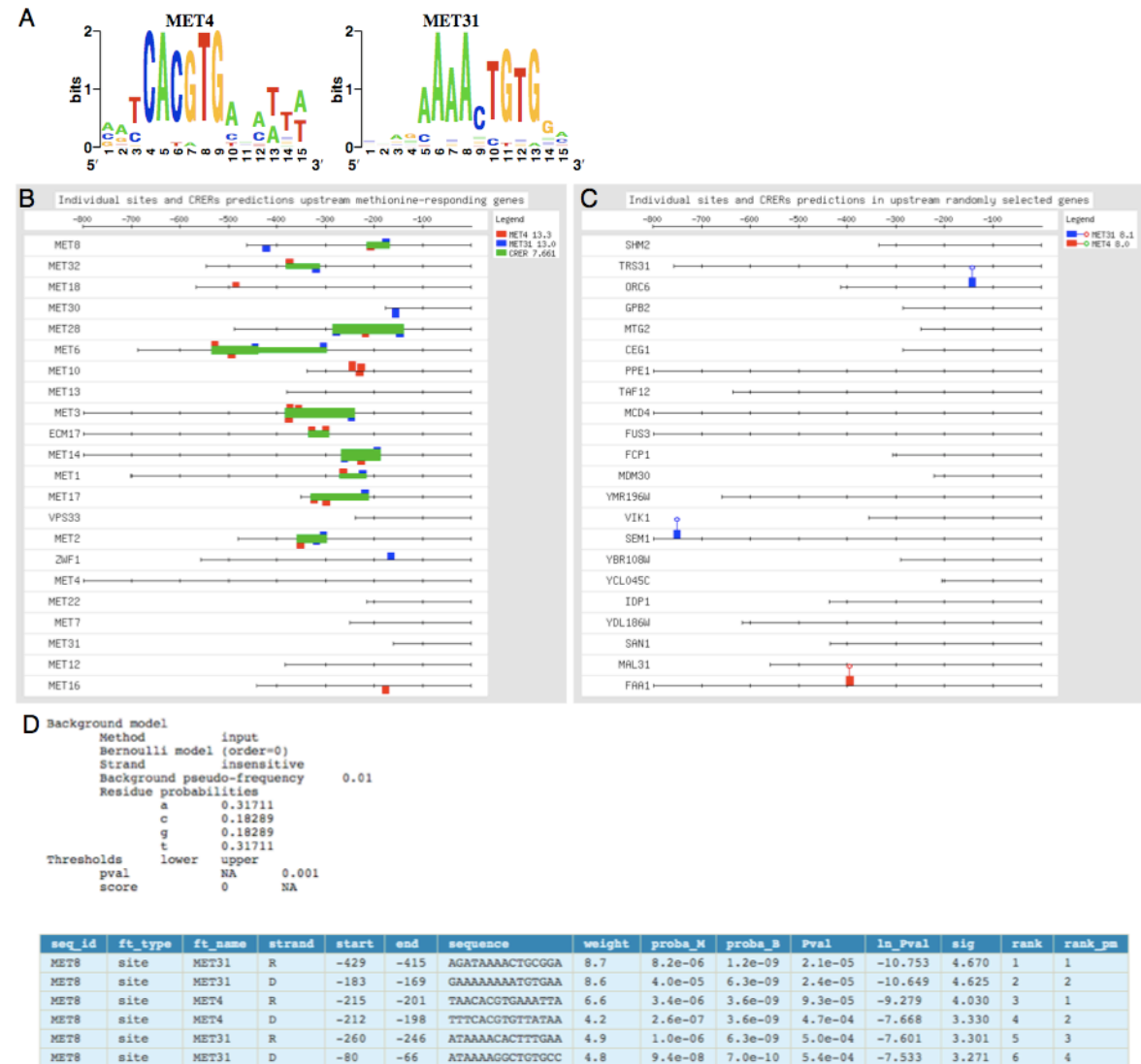
- Detection of regions containing a higher density of predicted TFBS than expected by chance.
- Various programs have been implemented to predict CRMs.
- The new RSAT program *matrix-scan* supports CRM prediction, by detecting regions enriched in cis-regulatory elements.
- Main features
 - Detection of homotypic (single motif) or heterotypic (distinct motifs) models.
 - All possible windows are tested within a user-specified width range (e.g. From 30 to 300).
 - The user has to specify a threshold on the P-value used for individual site predictions.
 - The program predicts all sites passing this threshold for each the input matrices, and detects regions (windows) having significantly more hits than expected by chance.
 - The enrichment is estimated by using the binomial statistics.



Turatsinze, J. V., Thomas-Chollier, M., Defrance, M. and van Helden, J. Using RSAT to scan genome sequences for transcription factor binding sites and cis-regulatory modules Nature Protocols accepted, (2008).

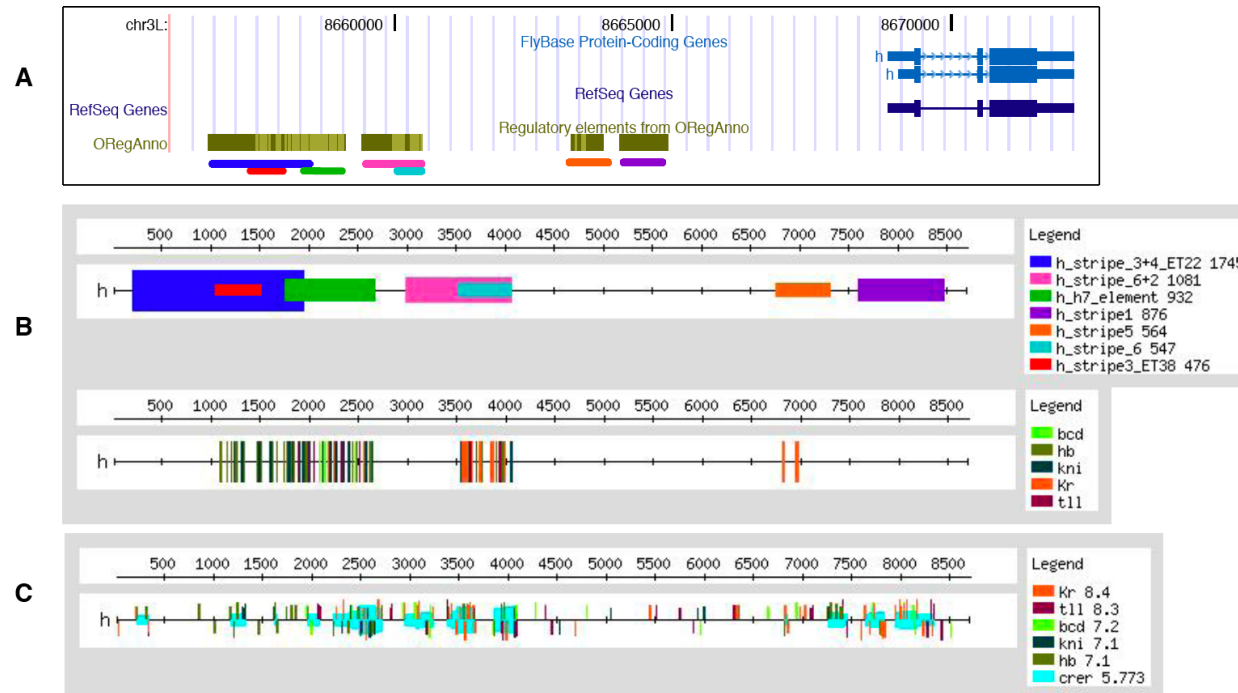
Cis-regulatory element enriched regions (CRERs) as putative cis-regulatory modules (CRMs)

- Example of CRER detection.
- Detection of methionine-responding genes in the yeast *Saccharomyces cerevisiae*.
- **A: matrices**
 - MET4: motif bound by the complex Met4p/Cbf1P/Met28p.
 - MET31: motif bound by either Met31p or Met32p (two homologous transcription factors).
- **B: predicted sites and CRERs in upstream non-coding sequences of MET genes.**
- **C: predicted sites and CRERs in random selections of yeast genes.**
- **D: examples of sites reported by matrix-scan.**



Using matrix-scan to detect Transcription Factor Binding Sites (TFBS) and Cis-regulatory Element Enriched Regions (CRERs)

- Example of CRER detection: upstream region of the *Drosophila* gene *hairy*



A

- Annotations in the ORegAnno database (displayed with the UCSC genome browser)

B

- Cis-regulatory modules annotated in FlyReg (top)
- individual sites annotated in ORegAnno (bottom).

C

- Scanning with 5 PSSM to detect individual binding sites + CRERs

Multi-genome CRER detection

- Morgane Thomas-Chollier (PhD thesis VUB/ULB, May 2008)
- Detection of CRERs with two matrices (HoxB1/Pbx and Prep/Meis) in the introns of the gene HoxA2.

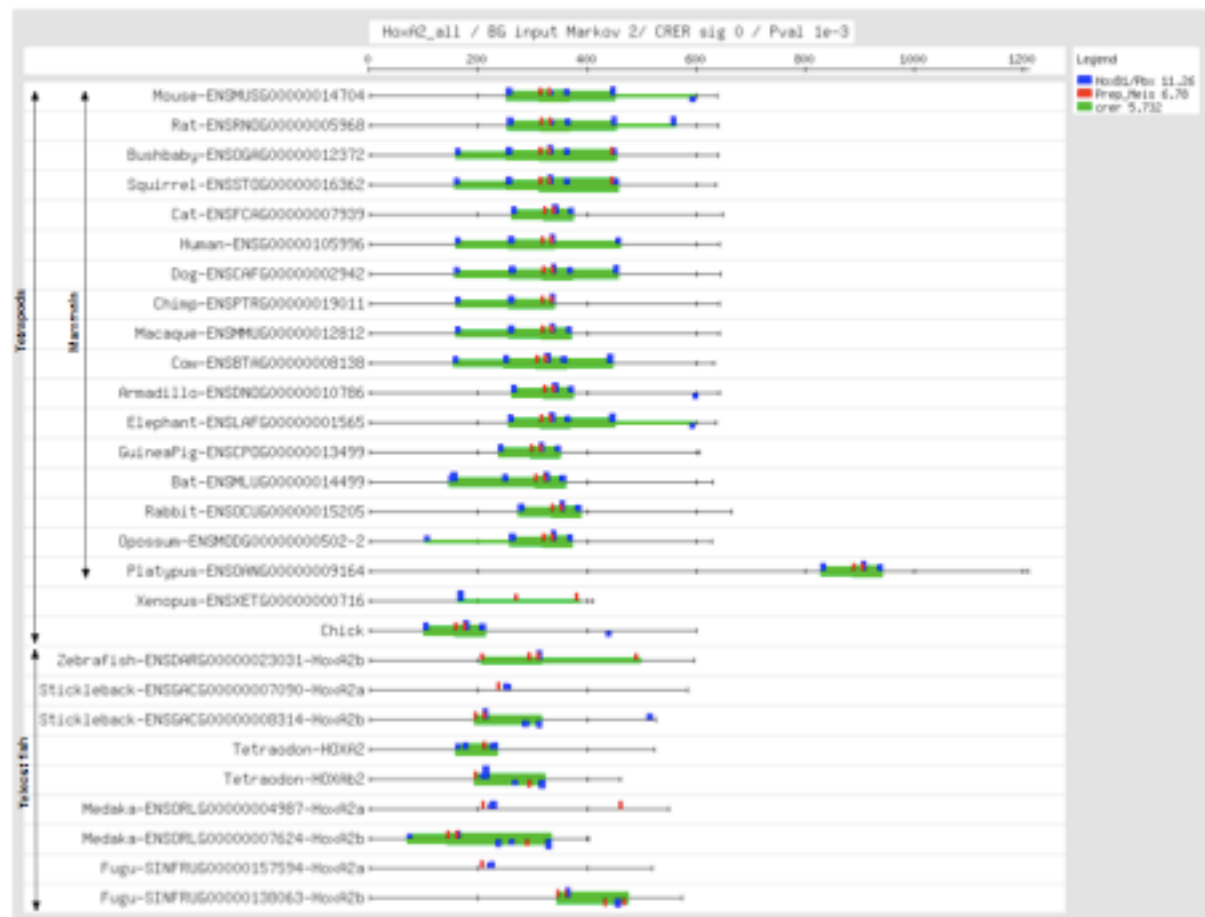
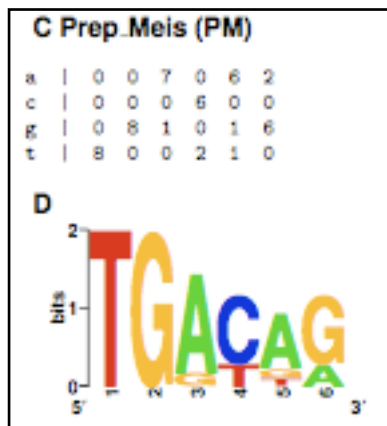
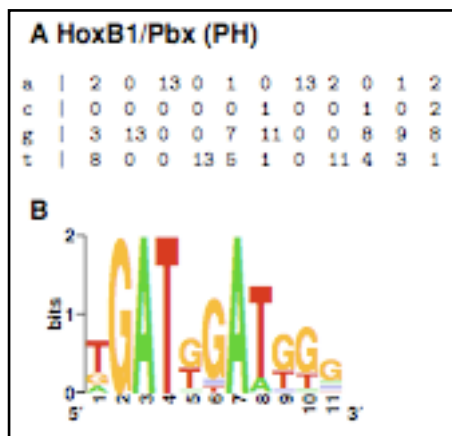


Figure 6.3: Cross-species predictions with matrix-scan in the HoxA2 intron. Predictions of TFBSs and CRERs in the HoxA2 intron of various vertebrate species. The height of each site is proportional to its weight score. CRER heights are proportional to their significance score. The numbers in the legend correspond to the highest weights for PH and PM matrices, and to the highest significance for the CRERs.