miRNA annotation

- A uniform system for microRNA annotation
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Expression criteria

- A. Detection of a distinct ~ 22-nt RNA transcript by hybridization to a sizefractionated RNA sample (ordinarily by the Northern blotting method).
- B. Identification of the \sim 22-nt sequence in a library of cDNAs made from size-fractionated RNA. Such sequences must precisely match the genomic sequence of the organism from which they were cloned

Biogenesis criteria

- C. Prediction of a potential fold-back precursor structure that contains the \sim 22-nt miRNA sequence within one arm of the hairpin.
 - the hairpin must be the folding alternative with the lowest free energy,
 - The hairpin must include at least 16 bp involving the first 22 nt of the miRNA and the other arm of the hairpin.

- The hairpin should not contain large internal loops or bulges,
 - Animal: these fold-back precursors 60-80 nt
 - Plant: more variable, upto a few hundred nt

- D. Phylogenetic conservation of the \sim 22-nt miRNA sequence and its predicted fold-back precursor secondary structure.
 - Minimal pairing requirement
- E. Detection of increased precursor accumulation in organisms with reduced Dicer function.

- A+D+E but A+D is sufficient in the absence of processing data
- A>B
- D>C

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miRNA: single molecule fold-back structure

 siRNA: RNA that are processed by Dicer from a hybrid between two antiparallel transcripts

- miR prefix + unique identifying number
 - miR-1, miR-2
- The genes that encode the miRNA:
 - Three letter prefix
 - Some cases, hypen
 - Capitalization
 - Italic
 - MIR156, mir-1

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- The identifying numbers: sequentially
 - with identical miRNAs having the same number, regardless of organism.
 - Nearly identical ortholog?
- Identical or very similar miRNA sequences within a species can also be given the same number, with their genes distinguished by letter and/or numeral suffixes
 - mir-13a mir-13b