

Explanation of *Hydractinia* RNA annotation GFF files

There are eight GFF files:

Four are for the **Hydractinia echinata** genome:

- **hech.trna.detailed.gff**: tRNA predictions, with all metadata.
- **hech.trna.minimal.gff**: tRNA predictions, without metadata.
- **hech.trna.detailed.gff**: Rfam predictions (all RNAs except tRNA), with all metadata.
- **hech.trna.minimal.gff**: Rfam predictions (all RNAs except tRNA), without metadata.

And four for the **Hydractinia symbiolongicarpus** genome:

- **hsym.trna.detailed.gff**: tRNA predictions, with all metadata.
- **hsym.trna.minimal.gff**: tRNA predictions, without metadata.
- **hsym.trna.detailed.gff**: Rfam predictions (all RNAs except tRNA), with all metadata.
- **hsym.trna.minimal.gff**: Rfam predictions (all RNAs except tRNA), without metadata.

Tab-delimited columns in detailed and minimal files:

Following from: <https://github.com/The-Sequence-Ontology/Specifications/blob/master/gff3.md>

1. **seqid**: sequence name
2. **source**: prediction method (tRNAscan-SE or cmsearch)
3. **type**: model (tRNA-**<isotype-model>**, or Rfam family accession)
4. **start**: start coordinate of prediction (always **<= end**)
5. **end**: end coordinate of prediction (always **>= start**)
6. **score**: bit score of prediction
7. **strand**: + if on positive strand, - if on negative strand
8. **phase**: uninformative in these files, always .
9. **attributes**: one or more **<key>=<value>;** strings, with possible **<key>**s listed below with explanations

Explanation of **<key>=<value>;** strings in attributes column:

Below are explanations of the **<value>** for each possible **<key>** you will find in the **attributes** column, separated into categories:

Software/database version:

- **tRNAscan-SE**: version of tRNAscan-SE used, always 2.0.5 when present, only present in **trna** files
- **Infernal**: version of Infernal used, always 1.1.2 when present, only present in **rfam** files
- **Rfam**: Rfam release used, always 14.1 when present, only present in **rfam** files

*These three keys are the only keys present in the **attributes** column in the **minimal** files, they are also present in the **detailed** files.*

Target sequence:

- **seqlen**: full length in nucleotides of target sequence

Data from cmsearch --tblout output file:

- **evaluate**: E-value of hit, in search of full genome *H. echinata* or *H. symbiolongicarpus* v1.0 assembly sequence file
- **mdlaccn**: Rfam model accession
- **mdlcoords**: <mdlstart>-<mdlend> model start and end positions of hit

Data from tRNAscan-SE -o output file:

- **type**: isotype model (tRNAscan-SE **tRNA type** column)
- **anticodon**: anticodon (tRNAscan-SE **Anticodon** column)
- **ibegin**: beginning position of intron, 0 for none (tRNAscan-SE **Intron Bounds - Begin** column)
- **iend**: ending position of intron, 0 for none (tRNAscan-SE **Intron Bounds - End** column)
- **pseudo**: **yes** if flagged as pseudogene, else **no** (**yes** if tRNAscan-SE **Note** column contains **pseudo**)

Repeat overlaps:

- **repeat_overlap**: HydSINE1(<start>-<stop> if this prediction overlaps by 10 or more nucleotides with a RepeatMasker HydSINE1 prediction from <start> to <stop> (<start> <= <stop>), else **no**

Note: only tRNA/HydSINE1 repeat overlaps are reported in this way. No overlaps between other repeat models and tRNAs are reported, nor any overlaps between repeat models and Rfam models.

Hit score data:

- **scF**: score/<max-score>, where **max-score** is the maximum score in the file for this model
- **highscoring**: **yes** if scF >= 0.9, else **no**

Hit length data:

- **lenF:** $\langle \text{length} \rangle / \langle \text{max-length} \rangle$, where $\langle \text{length} \rangle$ is $\text{stop-start}+1$, and $\langle \text{max-length} \rangle$ is the $\langle \text{length} \rangle$ for the prediction with the maximum score in the file for this model
- **fragment:** no if $\text{lenF} \geq 0.9$, else yes

Tandem array data:

- **tandem_array:** yes($\langle \text{start} \rangle - \langle \text{stop} \rangle$) if this prediction is in a tandem array that begins at position $\langle \text{start} \rangle$ and ends at position $\langle \text{stop} \rangle$ on strand **strand** ($\langle \text{start} \rangle \leq \langle \text{stop} \rangle$); ineligible if this prediction occurs on a sequence and strand with less than 10 predictions for this model and is consequently not eligible to be in a tandem array; no if this prediction is eligible to be in a tandem array and is not;
- **tandem_array_X:** number of predictions in the tandem array this prediction is in, *only exists if tandem_array value starts with yes*
- **tandem_array_N:** number of spacings between predictions in the tandem array this prediction is in that are within the range of $[\text{tandem_array_Dmin}, \text{tandem_array_Dmax}]$, *only exists if tandem_array value starts with yes*
- **tandem_array_Dmin:** minimum spacing length for the tandem array this prediction is in *only exists if tandem_array value starts with yes*
- **tandem_array_Dmax:** maximum spacing length for the tandem array this prediction is in *only exists if tandem_array value starts with yes*

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