## **Genome Annotation Submission Cover Sheet**

## Pre-SM\*ART QC Phage Genome Annotation Checklist

Phage Name: Azathoth

Your Name: Melinda Harrison
Your Institution: Cabrini University
Your email: mah348@cabrini.edu

Additional emails: conboya@chc.edu

(For correspondence)

Please check each box indicating completion of each task.

1. Does the genome sequence in your final contain the same number of bases and is it the same as the posted sequence on phagesdb.org?

2. Are all the genes "valid" when you click the "validate" button?

3. Have the genes been renumbered such that they go sequentially from 1 to the highest number?

4. Have all old BLAST hits been cleared, and all gene features reBLASTed?

5. Are the locus tags the "SEA PHAGENAME"?

6. Has the Documentation been recreated to match the information in the feature table?

7. Have tRNA ends been adjusted with web-based Aragorn and/or tRNAscan SE?

8. Has the frameshift in the tail assembly chaperone been annotated (where applicable?)

9. For the items below, generate a genome profile, and review the following. For the

YourPhageName\_CompleteNotes.dnam5 file:

a. Have any duplicate genes (or any with the same stop coordinate?) been removed?

b. Does every gene have **one and only one** complete set of Notes

c. Do the functions in the Notes match the official function list?

d. Are all three lines of functional evidence described for EVERY gene?

e. Do the notes contain the initial Glimmer/GeneMark data from the autoannotation?

For the YourPhageName .dnam5 file:

a. Have any duplicate genes (or any with the same stop coordinate?) been removed?

b. Is the Notes field empty for all the features with no known function (including hidden marks?

c. Do the function names in the Notes match the official function list?

d. Is the function field EMPTY for all features?

- 9. Did you use PECAAN to annotate your phage?
  - a. If, so please describe how in the text field after question 10.
- 10. Describe any issues or specific genes that you were unable to satisfactorily resolve, and warrant further inspection in the Quality Control review.

Genes 21 and 24 were deleted, more info can be found in their notes. Another gene 21 (13504 - 13776) was added because it fills in the gap, aligns with many other phage homologs, and gives good blast results and a function. Start sites for genes #3, #9, #19 were changed. Gene 26 (15288 - 46) is a wraparound gene with large overlap. We called gene #18 a 'holin' citing a few phage homologs as well as a modest HHPred result.

