## Homework 2

## Simple Count and Report

Write a program called squared\_cubed.py and prints out three columns of data, ideally, separated by tabs. A header line should be written which is labels of the columns

## N Squared Cubed

Column 1: numbers 0 -> 30 Column 2: Square (x^2) of column 1 Column 3: Cubes (x^3) of column 2

Output should look like this (but going up to at least 30 for for the N column)

- N Squared Cubed
- 0 0 0
- 1 1 1
- 2 4 8
- 3 9 27
- 4 16 64
- 5 25 125

## Genome Stats

We will compute some statistics for a tab delimited file called GFF which lists the location of genes and exons location in a genome annotation. Remember GFF is a structured format, tab delimited, which describes locations of features in a genome.

Recall eukaryotic Genes are made up of features: exons, introns, Untranslated regions (UTR). Some exons are coded as 'CDS' for CoDing Sequences - eg the ones that code for proteins.

See Wikipedia gene page and view of Gene structure in particular

Here is a GFF file for the *Penicillium chrysosporium* genome, which is the fungus which gave us one of the first antibiotics. The FungiDB database hosts genome sequences and data files for a collection of fungi.

The GFF file is available here FungiDB-54\_PchrysosporiumRP-78.gff and FastA format genome assembly is FungiDB-54\_PchrysosporiumRP-78\_Genome.fasta. These are two files related to location of genes and sequence data.

Write a script called <code>genome\_stats.py</code> to: 1. Download these file (this can be in UNIX before you run your python script or you can incorporate this into the python). I already wrote part of this for you in the template code you can start with that executes a <code>curl</code> command from within your script. But if this doesn't make sense to you, you can remove that. 2. Summarize how many exons, CDS, protein\_coding\_gene, are found in the genome annotation (GFF file) 3. Compute and print out the total length of all the protein\_coding\_gene features

(length is the END - START). 4. Compute and print out total length of all the CDS features (length is the END - START). 5. Use the FASTA file to compute the total length of genome (by adding up the length of each sequence in the file). Recall I lectured on a basic code to read in a FASTA file - you can also see that code template here 6. Print out the percentage of the genome which is coding (using the numbers calculated from the protein\_coding\_gene)