## **Documentation**

Table 1: General information about the suite of software to finish genome assemblies

Available at:	http://github.com/dcbmariano/scripts		
General requirements:	<ul> <li>Operational system: Linux 64bit</li> <li>Perl</li> <li>Python</li> <li>Biopython library         <ul> <li>http://biopython.org</li> </ul> </li> <li>NCBI-BLAST+         <ul> <li>http://blast.ncbi.nlm.nih.gov/Blast.cgi?PAGE_TYPE=BlastDocs&amp;DOC_TYPE=Download</li> </ul> </li> <li>Mira 4.0 Assembler         <ul> <li>http://mira-assembler.sourceforge.net</li> </ul> </li> </ul>		
Scripts:	<ul> <li>CONTIGuatorD.py</li> <li>contiginfo.py</li> <li>cut_letf.pl</li> <li>mapRepeat.py</li> <li>mcontig.py</li> <li>moveDNAA.py</li> </ul>		
Advanced install:	Paste the scripts on the folder ~/bin/ Open the folder on terminal and give permission to running: sudo chmod +x * Now you can run the scripts of two ways:  1. Typing "python ~/bin/script_name.py parameter_1 parameter_n"  2. Typing just "script_name parameter_1 parameter_n"		

**Table 2: Description of the Scripts** 

Script	Language	Function	Syntax
CONTIGuatorD.py	Python	Scaffolding contigs using a reference genome	python CONTIGuatorD.py -c [contigs_fasta_file] -r [reference_fasta_file] -g [reference_genbank_file]
contiginfo.py	Python	Analyze fasta files. It generates information about: number of contigs, min contig, max contig, length of genome and N50 value	python contiginfo.py [fasta_file]
cut_letf.pl	Perl	Allow performing cuts on the left region of fasta or fastq files	perl cut_left.pl [len_cut] [fasta_or_fastq_file]
mapRepeat.py	Python	Allow the scaffolding of the contigs obtained by <i>de novo</i> assembly, including repetitive	python mapRepeat.py [contigs aligned file] [reference file] [fastq xml folder] [contig left name]

		regions based on the extraction of the consensus sequence from the reads mapped into the reference genome	[contig right name]
mcontig.py	Python	Create fasta sequences separating N's regions.	python mcontig.py [fasta_file]
moveDNAA.py	Python	Correct the beginning of circular genomes finding the gene dnaA by a reference fasta file	python movednaa.py [seq] [reference fasta file]