To remember

- * Hifiasm is not always the perfect tool
- * Need to test at least two assemblers
- * Need an optimal coverage (between 15 and 70X according to the technology)
- * Known your genome (heterozygocity, size genome...)
- * If possible, sequencing in long reads
- * Check your assembly! (reads mapping, BUSCO, D-GENIES, kmer content...)