Genome analysis

No one tool to rule them all: prokaryotic gene prediction tool annotations are highly dependent on the organism of study

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Abstract

Motivation: The biases in CoDing Sequence (CDS) prediction tools, which have been based on historic genomic annotations from model organisms, impact our understanding of novel genomes and metagenomes. This hinders the discovery of new genomic information as it results in predictions being biased towards existing knowledge. To date, users have lacked a systematic and replicable approach to identify the strengths and weaknesses of any CDS prediction tool and allow them to choose the right tool for their analysis.

Results: We present an evaluation framework (ORForise) based on a comprehensive set of 12 primary and 60 secondary metrics that facilitate the assessment of the performance of CDS prediction tools. This makes it possible to identify which performs better for specific use-cases. We use this to assess 15 ab initio- and model-based tools representing those most widely used (historically and currently) to generate the knowledge in genomic databases. We find that the performance of any tool is dependent on the genome being analysed, and no individual tool ranked as the most accurate across all genomes or metrics analysed. Even the top-ranked tools produced conflicting gene collections, which could not be resolved by aggregation. The ORForise evaluation framework provides users with a replicable, data-led approach to make informed tool choices for novel genome annotations and for refining historical annotations.

Availability and implementation: Code and datasets for reproduction and customisation are available at https://github.com/NickJD/ORForise.

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Supplementary information: Supplementary data are available at Bioinformatics online.

1 Introduction

Whole genome sequencing, assembly and annotation is now widely conducted, due predominantly to the increase in affordability, automation and throughput of new technologies (Land et al., 2015). The prediction of protein-coding genes, specifically their corresponding CoDing Sequence (CDS) in prokaryote genomes has often been seen as an established routine. This is in part due to a number of assumptions and features, such as the high density (protein-coding genes contribute ~80–90% of prokaryote DNA) and the lack of introns (Lobb et al., 2020; Salzberg, 2019). However, this process involves the complex identification of a number of specific elements, such as: promoter regions (Browning and Busby, 2004), the Shine–Dalgarno (Dalgarno and Shine, 1973) ribosomal binding site and operons (Dandekar et al., 1998), which all contribute to identifying gene position and order. Additionally, the role of horizontal gene transfer (Jain et al., 1999) and pangenomes further complicates an already difficult process and likely contributes to errors and a lack of data held in public databases (Devis and Valencia, 2001; Furnham et al., 2012). Finally, our ability to characterize the functions of regions of DNA [which has been generally reserved for model organisms]
(MOs) and core genes (Russell et al., 2017) is being outstripped by the rate of genomic and metagenomic sequence data generation from non-MOs and non-core gene DNA sequences.

Before the turn of the century, it was understood that a great deal of work was still needed to address these issues. Studies had shown that many existing CDS prediction tools systematically failed to identify or accurately report genes whose features lay outside a rigid set of rules, such as non-standard codon usage, those which overlap other genes or those below a specified length (Burge and Karlin, 1998; Guigo, 1997). Since then, a systematic overview of 1474 prokaryotic genome annotations in GenBank concluded ‘the cause of the high rates of missed genes is less clear, largely due to a lack of information about the annotation methods used’ (Wood et al., 2012). Interestingly, while the majority of missed genes reported were under 300 nt, the annotation tools, which performed the incomplete annotations were developed to report CDSs at a minimum length of 110 nt. While there has been much work to address the problem of incomplete annotation, many gene types continue to be absent or under-represented in public databases (Huvel and Stumpf, 2014; Warren et al., 2010), such as short/small-ORFs (short ORFs) (Duval and Cossart, 2017; Storz et al., 2014; Su et al., 2013). This means that CDS prediction methodologies that use information from existing sequences are in turn ill-equipped to identify genes belonging to these underrepresented/missing gene types. It is therefore of paramount importance that we understand the limits of current CDS predictors as our reliance on automated genome annotation of novel genomes continues to increase (Brenner, 1999). Measures to compare both novel and contemporary CDS prediction tools are not well established or universally employed and novel tool descriptions tend to focus on algorithmic improvements rather than carrying out a systematic assessment of where the strengths or weaknesses in their approaches lie. This prevents researchers from gaining meaningful insight into the specific features of genes, which led to them being missed or partially detected, resulting in a lost opportunity to improve our understanding of prokaryotic genome content.

Gene annotation is challenging and is not a single step process. CDS prediction, often the first step, is fast, with little user input, but may require augmentation by different methods to supplement the initial predictions. One example is a tool, such as smoRFer (Bartholomäus et al., 2021), that specializes in finding short ORFs through the use of RNA-seq, which can detect transcription events under certain environmental conditions. Further examples use sequence conservation scores and homology searches to use existing database knowledge (Badger and Olsen, 1999; Dunne and Kelly, 2017; OhéEidegarth et al., 2014). Furthermore, pipelines are constructed [such as PROKKA (Seemann, 2014) and NCBI’s PGAP (Tatusova et al., 2016)] to automate these further rounds of annotation. However, the underlying CDS prediction tools are still core components of these pipelines and are still widely used as standalone tools.

Previous studies, which have evaluated prokaryotic CDS predictors generally only compared a small number of tools, focusing on algorithm design, and did not go into depth when reporting prediction accuracy with few other informative metrics used (Al-Turaiki et al., 2011; Mathé et al., 2002). A more recent study, BEACON (Kalkatawi et al., 2016), considered a small range of metrics including genes ‘denoted as identical, similar, unique with overlap or unique without overlap’ to either a reference annotation or from the output of three pipelines (PGAP, AAMG and RAST). Unfortunately, the types of genes missed were also not investigated further, leading to a lack of understanding of not only why and how they were missed, but also the impact on our biological understanding of the genome as a whole.

Many prediction methods used today are iterations of original concepts and thus are as in flux as the genomic databases themselves. Future development of CDS prediction techniques is now harnessing the recent advances in machine learning and other computational methods. While previous methods involve the construction of models either from organism-specific parameters, such as codon usage, guanine-cytosine (GC), complex motifs and average CDS length (Besemer and Borodovsky, 1999; Stanke and Morgenstern, 2005), opinions are shifting on the use and importance of MOs (Hunter, 2008; Levy and Currie, 2015; Russell et al., 2017). The volume of prokaryotic protein-coding gene sequences have enabled advanced machine-learning approaches, such as neural networks to predict CDSs that share common characteristics with a selection of previously annotated genes. One such example, Balrog (Sommer and Salzberg, 2021) predicts protein-coding genes by training from an array of non-hypothetical protein-CDSs from thousands of bacterial prokaryote genomes and aims to provide gene prediction across diverse species. Machine-learning models can be poor at making predictions for classes (e.g. genes) whose training data exhibit high levels of bias, error, are under-represented for specific groups (e.g. gene families) and groups for which they have not been trained (Schafer and Graham, 2002). In addition to this, prokaryotic gene families are chronically under-sampled (Warren et al., 2010). It is becoming clear that even with these advances in computational approaches, it is unlikely that we will ever be capable of identifying the complete picture of CDS gene diversity without exhaustive experimental work.

To address these concerns, we extensively evaluate a collection of 15 widely used CDS prediction tools that form the basis of most of the annotations deposited in public databases and therefore have largely been used to build the genomic knowledge used by the scientific community. We provide a comparison platform developed to allow researchers to compare 12 primary and a further 60 secondary metrics to systematically compare the predictions from these tools and study the effect on the resulting genome annotations for their species of interest. This allows for in-depth and reproducible analyses of aspects of gene prediction that are not often investigated and allows researchers to understand the impact of tool choice on the resulting prokaryotic gene collection.

2 Materials and methods

2.1 Current Ensembl genome annotations

Six bacterial MOs and their canonical annotations were downloaded from Ensembl Bacteria (Howe et al., 2020) (available at https://github.com/NickJD/ORForise/tree/master/Genomes), Bacillus subtilis BEST7003 strain (assembly ASM52304v1), Caulobacter crescentus CB15 strain (assembly ASM690v1), Escherichia coli K-12 ER3413 strain (assembly ASM80076v1), Mycoplasma genitalium G37 strain (assembly ASM2732v1), Pseudomonas fluorescens UK4 strain (assembly ASM73042v1) and Staphylococcus aureus 502A strain (assembly ASM59796v1) were chosen for their scientific importance, range of genome size, GC content, assumed near complete and high quality genome assembly and annotation provided by Ensembl Bacteria. They are presented in Table 1 and further information regarding these MOs can be found in Supplementary Section S1.

For each of the chosen MOs, two data files were downloaded from Ensembl Bacteria; the complete DNA sequence (*.dna.toplevel.fa) and the general feature format (GFF) file (*.gff3) containing the position of each gene. The current collection of CDS genes presented in the MO annotations from Ensembl (Current Ensembl Annotation (CEA)) was taken as the reference annotations for this study. Prokaryotic genomes exhibit high levels of gene density, often with little extraneous DNA, which is ‘commonly perceived as evidence of adaptive genome streaming’ (Sela et al., 2016). Unannotated DNA represents between ~10% and 20% of the six MO genomes selected and while an additional 0.38–2.22% is attributed to non-coding annotations, there is still a measurable portion of each genome without any annotation. This study focuses specifically on the identification of CDSs, which constitute the significant majority of annotated genomic regions in the six genomes studied (82.76–90.62%, see Table 1).

The CDSs from each of the six genomes exhibit a range of differences, which are known to impact the ability of prediction tools to identify them. These include, but are not limited to, GC content, codon usage and gene length. The GC content varies from 31.69% to 67.21% for these genomes, and even within a single genome, the CDS GC content varies widely (see Supplementary Fig. S1 for
distributions). Furthermore, the canonical ATG start codon is used between 68.58% and 90.67% of the genes for the six genomes (see Supplementary Table S1 for more detail).

Additionally, *M. genitalium* uses the codon translation table 4, meaning one of the three universal stop codons (TGA/UGA) is instead used to code for tryptophan (Dybvig and Voelker, 1996), whereas the other five MOs use the universal translation table 11 (see Supplementary Tables S1 and S3 for more detail). While a similar median CDS length is shared across the six genes, *B. subtilis* and *P. fluorescens* have a number of long genes (>8000 nt, see Supplementary Fig. S2) and *S. aureus* contains the 31,421nt ‘giant protein Ebh’ (Cheng et al., 2014), which is more than twice the length of the next largest CDS in this study. The diversity across the rest of prokaryotes is likely to be as great as, or greater than, reported here for these six.

The Sequence Ontology (Eilbeck et al., 2005) describes an ORF as ‘The in-frame interval between the stop codons of a reading frame which when read as a sequential string of amino acids’. However, it is conventional for ORFs to be reported as regions of DNA encompassed by a start and stop codon as a start codon is expected to indicate the start of DNA transcription (Brent, 2005). We acknowledge the difference in ontological definition and during this study, we refer to the region of DNA between an in-frame start and stop codon that is predicted to encode for an amino acid (protein) sequence, as a predicted CDS.

### 2.2 Prediction tools

This study specifically investigates CDS predictors, tools which apply complex filtering after the identification of ORFs across a region of DNA. This is different to ORF finders which, without filtering ORFs (Stothard, 2000) that meet a set of pre-defined rules, can be found in Supplementary Section S3 ‘Prediction Tools’.

Table 1. An overview of genome composition for the six MOs selected to evaluate CDS prediction tools compiled from data held by Ensembl bacteria.

| Model organism [assembly] | Genome size (Mbp) | Genes [CDSs] | Genome density [CDSs] | GC content (%) |
|---------------------------|-------------------|--------------|----------------------|---------------|
| *B. subtilis* BEST7003 [ASM52304v1] | 4.04 | 4133 [4011] | 88.91% [87.60%] | 43.89 |
| *C. crescentus* CB15 [ASM690v1] | 4.02 | 3875 [3737] | 90.60% [90.23%] | 67.21 |
| *E. coli* ER3413 [ASM80076v1] | 4.56 | 4257 [4052] | 86.28% [84.35%] | 50.80 |
| *M. genitalium* G37 [ASM2732v1] | 0.58 | 559 [476] | 92.03% [90.62%] | 31.69 |
| *P. fluorescens* UK4 [ASM73042v1] | 6.06 | 5266 [5178] | 84.75% [84.20%] | 60.13 |
| *S. aureus* S02A [ASM59796v1] | 2.76 | 2556 [2478] | 83.93% [82.76%] | 32.92 |

*Note:* Data are presented for all genes and CDS genes in bold square brackets. Note the relatively broad differences in genome size, gene density (percentage covered with annotation) and GC content.

Whole genome annotation ‘pipelines’, such as PROKKA (Seemann, 2014) and NCBI’s PGAP (Tatusova et al., 2016) were not included, but the initial CDS prediction components embedded in these pipelines, such as Prodigal and GeneMarkS-2, were included in the study. Multiple separate tools from the GeneMark family (Besemer and Borodovsky, 2005) were included (some superseded) due to their extensive use and impact on genomic knowledge over the last three decades.

### 2.3 Comparison method

A systematic software platform ORForise (ORF Authorise) was built to perform a fair, comparative, and informative analysis of the different tools examining different aspects of their predictions. Version 1.0 of the platform, written in Python3 (Van Rossum and Drake, 2009), was used and is freely available at https://github.com/NickJD/ORForise. It has been designed to process the standardized GFF3 format as well as the individual output formats produced by each tool listed in this study.

In this platform, we endeavoured to choose a wide range of metrics that clearly and representatively capture the many intricacies of the predictions. A number of metrics used in previous studies, such as the number of CDSs predicted, accurate identification of start positions or the number of genes correctly detected, can give some indication of the ‘accuracy’ of each tool. However, it was found during our analysis that there were many complexities in the prediction results, which would not be represented by these high-level metrics. For example, predicted CDS regions may overlap with one or more known CEA genes but be inaccurately extended or truncated on either the 5’ or 3’ end. It is also common for smaller CEA genes to be mistakenly encompassed by larger predicted CDSs and while the nucleotide regions of these genes are technically within the predicted regions, even if in-frame, they do not represent the true protein-CDS. Furthermore, different types of inaccuracies may be more or less important, depending on the aim of any given study. Therefore, clear and specific measures of accuracy that describe the detection of the entire locus of a gene are needed. Figure 1 illustrates how we determine correct CEA gene detection, but also explains its nuances and complexities. An example of this is the definition of short ORFs, which in prokaryotes are often described as having lengths of 100–300 nt (Duval and Cossart, 2017; Storz et al., 2014; Su et al., 2013). However, due to hard-coded cutoffs in many of the tools, we chose the ‘upper-bound’ of 300 nt or 100 codons to define short
Table 2. Version number and reference for all tools used in this study

| No. | Tool name            | Version | Reference                  |
|-----|----------------------|---------|----------------------------|
| 1   | Augustus             | 3.3.3   | Keller et al. (2011)       |
| 2   | EasyGene             | 1.2     | Nielsen and Krogh (2005)   |
| 3   | GeneMark hmm         | 3.2.5   | Lukashin and Borodovsky (1998) |
| 4   | GeneMark             | 2.5     | Borodovsky and McIninch (1993) |
| 5   | FGENESB '2020'       |         | Salamov and Solovyevand (2011) |
| 6   | Prodigal             | 2.6.3   | Hyatt et al. (2010)        |
| 7   | GeneMarkS            | 4.25    | Besemer et al. (2001)      |
| 8   | GeneMarkS 2          | '2020'  | Lomsadze et al. (2018)     |
| 9   | GLIMMER 3            | 3.02    | Delcher et al. (2007)      |
| 10  | GeneMark (H.A)       | 3.25    | Besemer and Borodovsky (1999) |
| 11  | TransDecoder         | 5.5.0   | Haas et al. (2013)         |
| 12  | FragGeneScan         | 1.3.0   | Rho et al. (2010)          |
| 13  | MetaGene             | 2.24.0  | Noguchi et al. (2006)      |
| 14  | MetaGeneMark         | '2020'  | Zhu et al. (2010)          |
| 15  | MetaGene Annotator   | 2008/8/19 | Noguchi et al. (2008)    |

Note: Tools 1–5 inclusive are model-based tools. Tools 6–15 inclusive are ab initio-based tools. Where no version number is available, the year when the tool was used is listed in single quotes.

ORFs. We iteratively developed 72 metrics to help provide the most accurate and informative representation of a tool’s prediction quality. Additionally, as part of the ORForise platform, we provide a number of Python3 post-analysis scripts developed to aid in the interrogation between the CEA gene annotations and the CDSs predicted by each of the tools studied. These scripts were used to extract characteristics that are useful in the investigation of why specific CEA genes are detected, missed or incorrectly reported.

2.4 Aggregated tool predictions

An extension to the ORForise comparison platform was built (Aggregate_Compare) to investigate whether an aggregation of predictions from a number of top-performing tools would perform better than individual tools. The CDS predictions from the selected tools are combined into a single data structure with duplicate CDSs filtered out, but alternative predictions for the same locus retained and ordered according to start position. The same comparison algorithm could then be employed on the set of unique CDS predictions identified by this union of the outputs of the selected tools (Prodigal, GeneMark-S-2, MetaGeneAnnotator, MetaGeneMark and GeneMark-S—chosen due to their individual performance) and as with the singular tool comparison, for every CEA gene, the CDS, which deviated the least from the correct locus was selected as the closest match.

2.5 Discovering additional ORFs

To enable the aggregation of different CDSs from contemporary and new annotations, we provide GFF_Intersector to create a single GFF representing the intersection of two existing annotations. This also provides an option to allow the retention of CDSs that have a user-defined difference (default minimum 75% coverage and in-frame). Additionally, we also provide the GFF_Adder tool, which produces a new GFF containing CDSs from an existing annotation, plus the new CDSs, filtered to remove any that overlap existing CDSs by more than 50 nt (user definable).

3 Results

3.1 Metrics for comparison of tools

A total of 72 different metrics were chosen for this exhaustive evaluation in order to give the broadest possible scope to compare and contrast the performance of the tools. The full definitions for each of these metrics can be found in Supplementary Section S5 and are intended to be used as a resource for the community when deciding which tool to apply to both novel and contemporary genome annotation work. The following are 12 of the most informative metrics selected for their ability to represent both a broad range and depth of different attributes which have been used to distinguish the prediction tools.

- M1 Percentage of Genes Detected
- M2 Percentage of Predicted CDSs that Detected a Gene
- M3 Percentage Difference of Number of Predicted CDSs
- M4 Percentage Difference of Median Predicted CDS Length
- M5 Percentage of Perfect Matches
- M6 Median Start Difference of Matched Predicted CDSs
- M7 Median Stop Difference of Matched Predicted CDSs
- M8 Percentage Difference of Matched Overlapping Predicted CDSs
- M9 Percentage Difference of Matched Short Predicted CDSs
- M10 Precision
- M11 Recall
- M12 False Discovery Rate

M1, Percentage of Genes Detected, is often used as the main indicator of tool performance in other comparisons but interpreted differently between studies. Here, it is characterized as a predicted CDS, which is in frame with a CEA gene and has captured at least 75% of its nucleotide sequence (Fig. 1A). In contrast to M1, which indicates when underprediction (or false negatives) occurs, M2 suggests when overprediction (or false positives) has occurred.

For M3, M4, M8 and M9, Percentage Difference was used to identify differences between predicted and CEA metrics: 100*(Predicted CDS metric—Ensembl Gene Metric)/Ensembl Gene Metric. The best score for a metric using the Percentage Difference calculation is 0, as 0 represents no deviation from the CEA annotations. The ‘Matched CDSs’ identifier used for M6, M7, M8 and M9 represent the CDSs, which have correctly detected an CEA gene. M6 and M7 are calculated by taking the median codon position differences recorded for mispredicted start or stop codons. Metrics, such as the Percentage of Perfect Matches (M5) can give a clearer overview of a tool’s ‘accuracy’ or performance, as it is common for a tool to misidentify either the exact start or stop locus of a detected CEA gene, while metrics, such as Median Start Difference of Matched Predicted CDSs (M6) can help establish the level of inaccuracy.
The tools were ordered by totalling the rankings for each of these 12 metrics, across the 6 MOs. Supplementary Results S1 contains the results used for the ranking. This ranking, based on a wide range of different performance measures, allows for a comparative overview of contemporary and future tools, and is presented in Figure 2. This figure also shows the Percentage of Genes Detected (M1) with an overlay of the Percentage of Perfect Matches (M5), demonstrating the inconsistency between the two metrics for each tool. Metrics, such as Percentage of Genes Detected (M1) and Percentage of Predicted CDSs that Detected a Gene (M2), are informative and can be representative of a tool’s prediction quality, however, they do not convey the complete picture when presented in isolation. This is of particular importance for those working with metageneomic or other fragmentary assemblies, as the likelihood of incomplete fragments and chimeric sequences is higher and can lead to varying mispredictions. Although the overall prediction quality of genes was high across most of the tools and genomes in this study, the additional metrics produced can be used to identify strengths and weaknesses inherent to them. For example, GeneMark.hmm (S. aureus model and genome), MetaGeneMark and MetaGeneAnnotator, GeneMarkS were all ranked highest for Percentage of Genes Detected (M1) for at least one MO, while Prodigal and GeneMarkS were ranked highest twice (GeneMarkS and GeneMark.hmm were ranked joint highest for S. aureus). However, when inspecting the 12 metrics (Supplementary Fig. S3), it was clear that there were complex differences between the prediction results of not only the highest scoring tools, but also the lower ranked tools, which were often ranked high for some metrics in some of the genomes.

While no tool or group of tools consistently ranked highest or equally across the 12 metrics or MOs, MetaGeneAnnotator ranked best for B. subtilis and M. genitalium, GeneMarkS-2 ranked best for C. crescentus and Prodigal ranked best for E. coli, P. fluorescens and S. aureus.

The combination of multiple metrics can be used to determine which tool should be used between two candidate tools with the same or similar Percentage of Genes Detected (M1). For M. genitalium, both GeneMarkS and MetaGeneMark obtained an M1 score of 39.50%, but MetaGeneMark reported a higher Percentage of Perfect Matches (M5) (65.96% compared to 61.17%) than GeneMarkS (see Fig. 2) and is thus more accurate.

In addition, GeneMarkS is ranked first for Percentage of Genes Detected (M1) when applied to P. fluorescens with 99.29%, compared to Prodigal, which is ranked 4th with 98.49%. However, Prodigal has the highest Percentage of Perfect Matches (M5), 92.36% versus 87.03% for GeneMarkS, which means that more of the CEA genes identified by Prodigal were exact matches. In this instance, choosing either Prodigal or GeneMarkS as the overall highest performing tool is not arbitrary.

### 3.2 Model-based versus ab initio tools

It was evident that the performance of model-based tools was less consistent across the six MOs than the ab initio tools. They could perform as well as or better than a number of ab initio tools when the model selected was the same as the genome annotated. However, if genome and model were not the same, they often produced predictions of extremely low quality. For example, GeneMark with the E. coli model only predicted 71 CDSs for S. aureus’s 2478 CEA genes, of which only 18 CDSs detected a CEA gene. However, while it could be expected that mixing different models and genomes could cause poor quality predictions from model-based tools, there were instances in which both model and genome were the same and the prediction performance was also poor. In particular, in the case of EasyGene using the S. aureus model, only 49.31% of S. aureus CEA genes were detected, a contrast from the ~99% detected by the majority of ab initio tools.

Intriguingly, Augustus (a model-based tool) when employed with the E. coli model, was able to detect 96.64% of P. fluorescens genes. Both genomes are Gammaproteobacteria, and thus Augustus may be identifying common features of their genes. While this shows that model-based tools can perform well even when their model and target genomes are different, when Augustus was applied to S. aureus using the S. aureus model, it was only able to detect 20.53%, but unexpectedly detected 78.91% when using an H.sapiens model. This is indicative of the inconsistency of model-based prediction tools and the genome models they employ. In contrast, through the ranking approach, we employed, the model-based tool GeneMark.hmm with the E. coli model ranked higher (72/21) than a number of ab initio tools in both the overall ranking and for individual metrics. Furthermore, GeneMark.hmm with the S. aureus model was joint top in detecting the highest number of S. aureus CEA genes with GeneMarkS. Additionally, for each of the model-based tools, the E. coli model performed better across the six MOs than the S. aureus model.
3.3 GC content
No significant variation was observed between the CEA gene median GC content and that of the predicted CDSs from each tool, even for those with poor predictions (see Supplementary Results S2). As can be seen in Supplementary Figure S1, each of the six genomes exhibits CEA genes with a wide range of GC content profiles, irrespective of their genome’s median value. We note that the GC content of genes missed by Prodigal is lower for all six MOs, but within the 25–75th percentile range for all CEA genes (Supplementary Fig. S1 and Table S4). Notably, E.coli and P.fluorescens genes, which were missed by Prodigal are nearly 10% lower in GC content than both detected and partial genes.

3.4 Overlapping CDSs
The overall number of CDSs predicted to have an overlap with another CDS varied across each of the tools and MOs, with cases of both positive and negative percentage differences when compared to the CEA annotations (see Supplementary Results S2 ‘Full Prediction Metrics’). Proportionally, the number of overlapping CDSs reported by *ab initio* tools are closer to the number of overlapping CEA genes than those reported by the model-based group.

Most model-based tools underpredict the proportion of overlapping CDSs with the exception of GeneMark E.coli for *P.fluorescens*, which predicted 2073 overlapping CDSs compared to the 1251 reported by Ensembl (see Supplementary Tables S5 and S6 and result S1 and S2).

Correct detection of CEA overlapping genes is also a problem. By totalling and averaging the Percentage Difference of Matched Overlapping Predicted CDSs (M8), we were able to observe a clear difference between the two tool groups with respect to their ability to detect correct overlapping CEA genes (see Supplementary Tables S5 and S6). The inability of the tools to account for the unusual nature of the *M.genitalium* genome was shown again with an average M8 across all tools of −88.21%, compared to the average of −27.77% for the other five genomes.

Furthermore, when making predictions for E.coli, while model-based tools, such as Augustus and EasyGene with the *E.coli* model can closely predict the proportion of overall overlapping CDSs (Percentage Difference of −1.42% and −2.30%, respectively), due to the poorer performance of these tools for correctly detecting CEA genes, their M8 scores for matched overlapping CDSs were substantially lower than the average score of the *ab initio* tools (grouped average of −52.89% as opposed to −23.62%—see Supplementary Table S6). Prodigal exemplifies this difference between the two tool groups. It was able to predict all overlapping CEA from *P.fluorescens* and *S.aureus*, whereas even when paired with the same model and genome, model-based tools continued to perform poorly.

3.5 Short ORFs
The lengths of detected, partially matched and missed CEA genes when predicted by Prodigal are summarized in Supplementary Table S4. It shows that the CEA genes, which were missed by Prodigal for each genome were substantially shorter in length than the genes, which were detected, except for *M.genitalium*. For the other five MOs, whose combined median length of missed genes is 317, less than half the combined median length of 837.5 of those detected (Supplementary Fig. S4 and Results S2), it is an alternative start codon selection, which influences whether a predicted CEA is shortened or elongated.

The proportion of short CEA genes in the six genomes below 300 nt ranged from 4.8% to 13.6% for each of the six MOs. All tools predicted many short CDSs for *M.genitalium* because they were incorrectly truncated due to its alternative stop codon usage. On average, *ab initio* tools were shown to be more likely to correctly detect short CEA genes across the other five MOs (see Supplementary Tables S7 and S8). Interestingly, unlike overlapping genes, short ORFs were more often overpredicted, but few were actually accurate when compared to the CEA. However, *ab initio* tools were much better suited to reporting the correct proportion of short predicted CDSs for all six genomes, often reporting the same proportion (see Supplementary Table S7). While *M.genitalium* does exhibit the highest divergence in proportional reporting of short predicted CDSs, *ab initio* tools were still less divergent (see Supplementary Table S9).

3.6 Partial matches
The number of missed CEA genes was low across the tools studied, with the exception of *M.genitalium* and some outliers from the model-based tools, such as GeneMark, Augustus and EasyGene. However, we also found many genes that were incorrectly reported on the 5' or 3' end. These misannotations, which we have called ‘partial matches’ if in the correct frame and accounting for >75% of a CEA gene, constitute either an elongation or truncation of the protein product of the gene and therefore potentially have an unknown impact on the resultant sequence. A large number of genes were incorrectly reported on the 3' end for *M.genitalium* by each tool. These 3' truncated CDSs are explained by the alternative use of TGA as tryptophan in *M.genitalium* (tools incorrectly assume this encodes a stop codon). The stop codons predicted for *M.genitalium* by all the prediction tools were the same ‘TGA, TAG, TAA’ as for the CEA genes of the other five MOs. Interestingly, one CEA gene in *E.coli*, which used CTT as a stop codon, was missed by all tools except for FGENESB with its *E.coli* model. FGENESB incorrectly reported the very next codon, TGA, as the stop position. This 78 nt CEA is the only example, we found of a tool extending a CEA gene not from *M.genitalium*. Augustus with the Human model made a number of non-standard predictions due to its propensity to search for multiple CDSs for each gene but this is to be expected and is not reported in these results. Unlike 3' misprediction, a large number of genes from all six genomes were predicted with alternative start codons (see Supplementary Results S2). This was true for all tools and especially a problem for *C.crescentus* with a relatively low 68.58% ATG start codon usage for all CEA genes. The CEA genes for which Prodigal was unable to obtain a ‘Perfect Match’ (M5), was just 37.40%. Prodigal used a much higher level of ATG (80.87%) for this set of partially matches genes. This misidentification of start codon usage was a consistent problem among all the tools and genomes studied. However, for *E.coli*, the level of misidentification was lower. As an example, the number of times the correct or incorrect start codons were selected by Prodigal, across all six MOs, including the number of incorrectly chosen instances of the start codon (e.g. a different ATG further upstream of the real ATG) can be seen in Supplementary Table S2.

3.7 Aggregated tool predictions
Combined prediction approaches have previously been utilized to harness the prediction power from multiple tools to increase the number of detected CEA genes (Tatusova et al., 2016; Yok and Rosen, 2011). For each of the MOs, taking the union of the top 5 tool predictions did provide a small increase in the number of Genes Detected (M1) (and a reduction of partial matches) compared to that of the ‘best tool’ [tool with highest percentage of Genes Detected (M1)] for any particular organism. However, even with this extreme case of using the union of all predicted CDSs, the increase in M1 was negligible (average increase of 0.47%) and came at the expense of predicting a large number of additional incorrect CDSs, as can be seen in Supplementary Table S10. Even in the case of *M.genitalium*, the M1 was not improved more than 0.21% with the union prediction.

3.8 Improving historic annotations
Using the GFF_Adder tool, we investigated the potential of Prodigal to add additional CDSs to the CEA annotations. There are more than 60 additional predicted CDSs that can be found for each of our MOs, and more than 270 for *E.coli* and *P.fluorescens* (see Supplementary Table S11).
4 Discussion

4.1 Ab initio tools usually perform better than model-based

We found that ab initio tools usually perform better than model-based tools. While no one tool performed the best or worst across all metrics, the ab initio tools Prodigal, GeneMarkS-2, MetaGeneAnnotator, MetaGeneMark and GeneMarkS were ranked first–fifth, respectively, across our 12 metric ranking (Supplementary Fig. S3 and Results S1 and Supplementary Rankings).

Strains of the same species can exhibit large intraspecies variation (Van Rossum et al., 2020). Additionally, genes resulting from horizontal transfer, which is more frequent within species (Van Rossum et al., 2020), are likely to contain features from the donor strains, which the rigid model-based methods are unable to recognize. GeneMark, a model-based tool, published in 1993, even when both target genome and model were E.coli, was identified as one of the worst performing tools in this study, possibly driven by the well-known large open genome of this species (Lukjancenko et al., 2010). The same was observed for S.aureus. While model-based tools can perform well even when their model and target genomes are different, in the case of Augustus, when applied to the C.crescentus genome using the S.aureus model, it was only able to detect 3.93% of CEA genes, but unexpectedly detected 78.75% when using the H.sapiens model. Unsurprisingly, model-based predictors have therefore fallen out of development and use over the last decade and ab initio-based tools, such as Prodigal, GeneMarkS-2 and GLIMMER3 have become ubiquitous.

4.2 Codon usage has a large influence on accuracy

We found that codon usage has a large influence on accuracy due to its influence on start and stop codon choice, even in MOs.

The re-coding of a stop codon as an amino acid is rare and seems to be taxa specific (Dybvik and Voelker, 1996). While many of the tools offered the ability to change codon tables (often accounting for TGA specifically), the correct codon tables or codon preferences for each genome cannot be known in advance of annotation of a novel organism. Despite this, we would expect that they should be able to predict a significant proportion of genes, even in the absence of the knowledge of a different codon usage table. Some tools, such as Prodigal will assess a genome using both the universal and Mycoplasma translation table, however remarkably this did not increase the accuracy of the tool when analysing M.gentilium genome (see Fig. 2). Overall TGA was never predicted as tryptophan usage over the last decade and ab initio-based tools, such as Prodigal, GeneMarkS-2 and GLIMMER3 have become ubiquitous.

4.3 Metagenomic annotation approaches are suitable for whole genome sequences

Interestingly, tools made specifically for metagenomic and fragmented genome annotation performed better than most single genome tools (tools ranked third, fourth and sixth were developed for metagenome annotation), possibly indicating that even ‘complete’ genomes may themselves still harbour elements of sequencing and assembly error which these types of algorithms have been designed to account for. Most genomes submitted to databases, such as the NCBI Genome repository (Haft et al., 2018), are incomplete and can contain hundreds of fragments which can make gene prediction an even more difficult task. As S. Salzberg said in 2019 “Paradoxically, the incredibly rapid improvements in genome sequencing technology have made genome annotation less, not more, accurate” (Salzberg, 2019). This indicates that future annotation work performed on non-model and more diverse organisms may benefit from approaches implemented by metagenomic tools.

4.4 Short genes and overlapping genes are often misreported

We found that short genes and overlapping genes are often misreported and that many tools still have hard-coded limitations and weightings against these types of genes, with model-based tools performing especially poorly.

It has been well established in the literature that short genes are likely under-represented across genomic databases, and therefore, possibly even within the Ensembl data used in this study (Duval and Cossart, 2017; Storz et al., 2014; Su et al., 2013). The growing acceptance that short genes are not only common in prokaryotic genomes but also have important roles (Andrews and Rothnagel, 2014), is at odds with many tools still containing hard-coded limitations for minimum CDS length and algorithmic weights against short CDSs. As might be expected because of its re-coding of TAG, M.gentilium proved challenging for all tools to accurately identify CDSs, resulting in the early truncation of a large proportion of CEA genes and an increase in predicted short CDSs. This often led to the tools predicting additional spurious short CDSs in the missed regions (a result that can be seen in the low M10 Precision metric for this genome). However, for the other genomes, most tools also predicted too many short CDSs (9.07% and 39.10%, for ab initio- and model-based tools, respectively), but paradoxically still managed to miss a large proportion of Short CEA genes in the Ensembl annotations (missing 26.38% and 53.69% for ab initio- and model-based tools, respectively) (see Supplementary Tables S7–S9 and Results S2).

For overlapping genes, while ab initio tools performed better than model-based tools (see Supplementary Tables S5 and S6), in general they both under-predicted the number of overlapping CEA genes across the genomes (on average ~6.07% and ~30.15% for ab initio- and model-based tools, respectively) (see Supplementary Tables S5 and S6 and Results S2). No tool was able to correctly detect more than 20% of M.gentilium’s overlapping CEA genes. Overlapping and nested genes have now become an area of renewed interest for their potential impact on genomic organization and evolution (Huvet and Stumpf, 2014; Krakauer, 2000). For example, mokC in E.coli, believed to be a regulatory peptide, completely overlaps hokC and enables hokC expression (Pedersen and Gerdes, 1999) and no tool was able to detect both genes correctly.

Overall, the tools struggled to handle overlapping gene loci, and often returned either only one or neither of the overlapping coding regions in their predictions. This may be due to the manner in which many tools filter multiple candidate ORFs for a single locus leading to sub-optimal predictions. For example, Prodigal reports a CDS in C.crescentus on the positive strand at 23 760–24 074 when the CEA CDS is 23 350–24 170 on the negative strand. The unallocated space (24 074–24 170) resulted in Prodigal reporting the next downstream CDS starting at 24 091 instead of 24 133 (as in the Ensembl annotation), erroneously including 5’ UTR in the predicted CDS. There are now tools to identify putative short ORFs in both prokaryotes and eukaryotes using additional evidence, such as RNA expression data (Bartholomaius et al., 2021; Ji et al., 2020; Miravet-Verde et al., 2019).
Our results suggest that the identification of short and overlapping CDSs cannot be done independently without the potential for unforeseen consequences for annotation accuracy.

4.5 Historic bias affects gene prediction today

Overall, we have observed an increase in accuracy in tools over time as can be seen with the different versions of GeneMark compared here: the overall rankings of model-based GeneMark (1993) (with E.coli/S.aureus models), ab initio GeneMarkS (2001) and ab initio GeneMarkS-2 (2018) are 20/17, 5 and 2, respectively. However, GeneMarkS (2001) performed better than its successor GeneMarkS-2 (2018) for 5 out of the 12 metrics in Supplementary Figure S3 including Percentage of Genes Detected (M1) in P.fluorescens, M.genitalium and B.subtilis (see Supplementary Results S1 and Supplementary Rankings). The performance of GeneMarkS (2001) in M1 may reflect its use for an extended period of time in the NCBI Prokaryote Annotation Pipeline. Possibly as a result of this, many of the CEA genes GeneMarkS (2001) detected were originally identified by the tool itself. Similarly, all model-based tools performed at their best across the 12 metrics and 6 MOs when using their E.coli model, hinting at the impact of historical research in this organism. Advances in the realms of machine learning and statistical modelling have the greatest potential to address these issues but are also likely to be the most prone to historical biases in the databases. Many of the rules, such as standard CDS length and codon usage, are inferred from previously identified CDSs. The existence of annotation errors and omissions in various sequence databases is well established and unlikely to be resolved in the near future without significant coordination between repositories (Klimke et al., 2011). Additionally, much of the sequence information derived from MOs will become less relevant as greater numbers of novel organisms are sequenced (Hunter, 2008).

These issues have been raised previously: In 2009, the ‘Best Practices in Genome Annotation’ meeting report listed a number of areas of concern put forward by attendees (Madapu et al., 2010) including tool choice, strategy to update and correct previous annotations, tracking of changes in databases, prioritization of certain genes for experimental evaluation, documenting processes and keeping up with technological advances. The work presented here addresses the issue of tool choice, but many of the recommendations are yet to be realized. The lack of any previous detailed systematic overview of method performance may also have played a part in these biases not being addressed to date. Our study has shown that tool selection needs to be fully informed by its intended purpose and the tool’s weaknesses.

4.6 Current and future techniques are needed to continue annotation improvements

It is clear from both this and previous studies that combinatorial approaches are fundamental in bridging the gap to the next stage of genome annotation. This has clearly already begun with pipelines, such as PROKKA and PGAP, which utilize a collection of techniques, most notably, advanced homology searching to complete annotations where traditional CDS predictions fail or produce competing predictions. However, this can also lead to conflicting annotations. As noted, homology searches are only as good as the database being used. The presence or absence of homology does not indicate whether an ORF is a true CDS gene, especially in the nuanced field of alternative ORFs (Orr et al., 2020). Further complications involving alternative ORFs, many of which are overlapping, can be found with new evidence in E.coli, where ‘Ribosome profiling revealed out-of-frame internal minimal ORFs in 13 E. coli genes. Mutation of the start codon... in one gene, yecJ, resulted in an increase in translation of the main-ORF, suggesting that these minimal ORFs can also modulate translation of the main-ORF’ (Meydan et al., 2019).

As users of computational genomic techniques, we must realize when we have reached the limit of what is possible with the contemporary data available. This, together with other studies, proposes that the linchpin required for the next step in genome annotation, is not new and more techniques reliant on current genomic knowledge, but instead more experimental work and species agnostic approaches. However, the near unlimited scope of growth conditions, environmental pressures et cetera, has made the prospect of experimentally validating all potential CDS regions unfeasible. Finally, while great strides have been made in experimentally validating difficult to characterize gene types, one such study ‘...suggest[s] substantial numbers of small proteins remain undiscovered in E. coli, and existing bioinformatics techniques must continue to improve to facilitate identification’ (VanOrdel et al., 2018).

5 Conclusion

We have presented a comprehensive set of metrics, which distinguish CDS prediction tools from each other and make it possible to identify which performs better for specific use-cases. The ORForise evaluation framework enables users to evaluate new and existing annotations and generate consensus and aggregate gene predictions. We have demonstrated that certain types of genes, such as short genes, overlapping genes and those with alternative codon usage, are still elusive, even to the most advanced ab initio techniques. Worryingly, the performance of any tool seems to depend on the genome that is being analysed. For instance, Prodigal, which ranked best overall, was ranked first for E.coli, S.aureus and P.fluorescens, MetaGeneAnnotator was ranked first for B.subtilis and M.genitalium and GeneMarkS-2 was ranked first for C.crescentus (see Supplementary Rankings). Additionally, no individual tool ranked as the most accurate across all genomes for the Percentage of Genes Detected (M1) (the single metric historically used to assess tool performance) or any other individual metric. This is likely to have a measurable impact on downstream genomic and pangenomic studies. However, overall, we found Prodigal to be one of the most well-rounded tools, not only detecting the highest number of CEA genes for two very diverse MOs (E.coli and M.genitalium), but also performing overall best when ranked across the 12 metric rankings and 6 MOs. It was also overall best for Perfect Matched genes (M5). However, as outlined earlier, it was not always ranked first for all genomes, further suggesting that users should choose tools carefully, based on the organism and question they are studying. Finally, we advise against generating aggregated ab initio annotations from multiple tools where no existing annotation is available for the genome, as this results in poor overall performance. However, additional cycles of annotation with tools designed to identify putative CDSs in the unannotated regions, show promise for improving current prokaryotic genomic knowledge.

Author contributions

All authors discussed the conceptualization of the comparison platform and its impact. N.J.D. wrote the code. All authors contributed to the manuscript.

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