

# Article Title

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## 2 ABSTRACT

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Keywords: Text Text Text Text Text Text Text Text

## 1 INTRODUCTION

Hello, how are we doing?

## 2 METHODS

### 8 2.1 Generating Predictions

Used GOMAP on condo lalala. Input files are (usually) published along results.

### 10 2.2 Clean up

All functional annotation sets were cleaned up the following way (using definitions from the Gene Ontology version 2019-07-01):

1. Any annotations where the GO accession was marked as obsolete were removed.
2. Some terms in the GO have 'alternative ids'. When naively removing duplicates, two entries will not be recognized as duplicates if they have different accessions pointing to the same GO term. Therefore, all GO accessions were changed to their respective 'main id' and the dataset was again scanned for duplicates.
3. Any annotations with modifiers (NOT, contributes.to...) were removed since no tool used in the further analysis can handle them.

Table 1 provides information on the number of annotations that were removed this way from each dataset. All further analyses were performed on the cleaned datasets since we assume the user will only be interested in still valid and non-redundant functional annotations.

## 2.3 Quantitative Evaluation

lalala lololo table xyz

## 2.4 Quality Evaluation

Quality evaluation of gene function predictions is not trivial and usually done by comparing the set of predicted functions of a gene against a *gold standard* consisting of annotations that are assumed to be correct. We used annotations that were created or in some way curated with human participation for gold standards. There are a plethora of different metrics to perform the comparison of predictions against this gold standard. When we first published GOMAP (Wimalanathan et al., 2018), we used a modified version of the hierarchical evaluation metrics originally introduced in (Verspoor et al., 2006) because they were simple, clear, and part of an earlier attempt at unifying and standardizing GO annotation comparisons (Defoin-Platel et al., 2011). In the meantime, Plyusnin et al. (2019) have published an approach for evaluating different metrics showing substantial differences within the robustness of different approaches. TODO DESCRIBE THEIR APPROACH We have applied their method on the Gold Standards available to us to determine which evaluation metric is the most appropriate in our case. The results of this analysis can be seen in TODO.

We then evaluated our predictions and the other annotation sets using the best performing metrics as well as the one we previously used (Table TODO).

## 2.5 Phylogenetic Tree Construction

To demonstrate that a more top-level and holistic use of whole-genome functional predictions can still be useful we devised some simple ways of applying phylogenetic methods to our predictions. ### Distance Based ### Character Based

## 2.6 Ensuring Reproducibility

containerization, github...

# 3 RESULTS

... a quantitative comparison of the datasets in Table.

## 3.1 Quality Evaluation

TODO If it turns out that our predictions are good with hF but bad with more appropriate metrics, explanation would be that score thresholds for the prediction tools used in the GOMAP pipeline have been chosen to maximize this hF value. It now seems reasonable to re-adjust these thresholds to maximize a different metric which will likely result in a drop in hF score but increase in other metrics. Again emphasizes the importance of choosing the right evaluation metric. Also shows how comparison between different pipelines/predictions can be difficult if chose different metric or optimized for different metric. Also: if an annotation is not present in the gold standard, there is no way of knowing whether that gene truly doesn't have that function or whether it has just never been characterized/examined. So we cannot distinguish between a biologically true negative and an actually false negative in the gold standard. This

**Table 1.** Number of removed annotations during cleanup.

Genome	Dataset	Obsolete Annotations	Duplicates	Annotations with Modifiers
<i>Arachis hypogaea</i>	GOMAP	3437	13	912
<i>Brachypodium distachyon</i>	GOMAP	2512	49	789
<i>Cannabis sativa</i>	GOMAP	1714	6	757
<i>Glycine max</i>	GOMAP	3333	10	930
<i>Gossypium raimondii</i>	GOMAP	1781	7	822
<i>Hordeum vulgare</i>	GOMAP	1877	8	815
	GoldStandard	0	4	0
<i>Medicago truncatula</i> A17	GOMAP	2673	10	798
	GoldStandard	0	7	0
	Gramene62-IEA	429	251	0
<i>Medicago truncatula</i> R108	GOMAP	4168	7	803
<i>Oryza sativa</i>	GOMAP	1642	7	869
	GoldStandard	44	581	0
	Gramene61-IEA	242	28	0
<i>Phaseolus vulgaris</i>	GOMAP	1190	6	783
<i>Pinus lambertiana</i>	GOMAP	1839	4	587
<i>Sorghum bicolor</i>	GOMAP	2384	66	783
<i>Triticum aestivum</i>	GOMAP	9624	17	1132
	GoldStandard	0	10	0
	Gramene61-IEA	706	88	0
<i>Vigna unguiculata</i>	GOMAP	1269	6	811
<i>Zea mays</i> B73.v3	GOMAP	1805	92	709
	GoldStandard	1	11	0
	Gramene49	221	6	0
	Phytozome	132	0	0
<i>Zea mays</i> B73.v4	GOMAP	2077	89	848
	GoldStandard	65	207	0
	Gramene61-IEA	600	178	0
<i>Zea mays</i> Mo17	GOMAP	2346	83	823
	GoldStandard	1	60	0
<i>Zea mays</i> PH207	GOMAP	2676	82	830
	GoldStandard	1	70	0
<i>Zea mays</i> W22	GOMAP	2681	88	840
	GoldStandard	1	52	0

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57 poses a problem when annotations are predicted that are not found in the gold standard: Is this truly a wrong  
58 prediction or is the gold standard incomplete? Especially in our case where the predictions not only contain  
59 more annotations than the gold standard, but are also more diverse. In effect this means that a quality  
60 score as calculated above may not only describe the quality of the prediction, but to some extent also the  
61 completeness of the gold standard itself. At least we can see here that gold standards with a median of 3  
62 annotations per gene resulted in higher quality scores than gold standards with less annotations per gene,  
63 even though predictions were generated the same way in all cases. TODO maybe put a figure  
64 with regression quality score/median annotations per gene or something In

**Table 2.** Quantitative metrics of the cleaned functional annotation sets. CC, BF, MP, and A refer to the aspects of the GO: Cellular Component, Biological Function, Molecular Process, and Any/All.

Genome	Genes	Dataset	Genes Annotated[%] <sup>a</sup>				Annotations <sup>b</sup>				Median Ann. per G. <sup>c</sup>			
			CC	BF	MP	A	CC	BF	MP	A	CC	BF	MP	A
<i>Arachis hypogaea</i>	67,124	GOMAP	85.85	84.68	100.00	<b>100.00</b>	150,525	132,144	493,145	<b>775,814</b>	2	2	6	<b>10.0</b>
<i>Brachypodium distachyon</i>	34,310	GOMAP	81.33	85.35	100.00	<b>100.00</b>	74,172	69,213	255,397	<b>398,782</b>	2	2	6	<b>10.0</b>
<i>Cannabis sativa</i>	33,677	GOMAP	94.22	95.48	100.00	<b>100.00</b>	85,755	73,614	262,741	<b>422,110</b>	2	2	6	<b>11.0</b>
<i>Glycine max</i>	52,872	GOMAP	86.95	88.92	100.00	<b>100.00</b>	126,470	113,068	416,989	<b>656,527</b>	2	2	6	<b>11.0</b>
<i>Gossypium raimondii</i>	37,505	GOMAP	93.00	92.37	100.00	<b>100.00</b>	95,419	84,910	307,470	<b>487,799</b>	2	2	6	<b>11.0</b>
<i>Hordeum vulgare</i>	39,734	GOMAP	88.57	91.76	100.00	<b>100.00</b>	86,489	79,727	272,420	<b>438,636</b>	2	2	5	<b>10.0</b>
		GoldStandard	0.02	0.05	0.05	<b>0.07</b>	7	23	45	<b>75</b>	0	1	1	<b>2.0</b>
<i>Medicago truncatula</i> A17	50,444	GOMAP	83.79	86.69	100.00	<b>100.00</b>	104,902	99,155	363,608	<b>567,665</b>	2	2	6	<b>10.0</b>
		GoldStandard	0.03	0.06	0.05	<b>0.07</b>	18	35	48	<b>101</b>	0	1	1	<b>2.5</b>
		Gramene62-IEA	34.15	50.66	38.98	<b>65.71</b>	33,350	62,800	39,230	<b>135,713</b>	1	1	1	<b>3.0</b>
<i>Medicago truncatula</i> R108	55,706	GOMAP	72.10	90.14	100.00	<b>100.00</b>	108,388	107,499	381,831	<b>597,718</b>	1	2	5	<b>9.0</b>
<i>Oryza sativa</i>	35,825	GOMAP	79.78	83.31	100.00	<b>100.00</b>	71,306	64,150	248,304	<b>383,760</b>	2	2	6	<b>9.0</b>
		GoldStandard	15.98	20.61	25.21	<b>31.79</b>	7,729	11,033	19,375	<b>38,145</b>	1	1	1	<b>3.0</b>
		Gramene61-IEA	29.76	43.21	46.63	<b>59.79</b>	14,475	32,703	39,101	<b>86,283</b>	0	1	1	<b>3.0</b>
<i>Phaseolus vulgaris</i>	27,433	GOMAP	94.48	93.06	100.00	<b>100.00</b>	70,987	64,022	229,230	<b>364,239</b>	2	2	6	<b>11.0</b>
<i>Pinus lambertiana</i>	31,007	GOMAP	92.67	95.91	100.00	<b>100.00</b>	71,247	68,315	212,248	<b>351,810</b>	2	2	5	<b>10.0</b>
<i>Sorghum bicolor</i>	34,129	GOMAP	82.44	85.98	100.00	<b>100.00</b>	75,145	69,659	259,004	<b>403,808</b>	2	2	6	<b>10.0</b>
<i>Triticum aestivum</i>	107,891	GOMAP	88.53	90.98	100.00	<b>100.00</b>	259,318	217,467	785,051	<b>1,261,836</b>	2	2	6	<b>10.0</b>
		GoldStandard	0.89	0.57	1.54	<b>1.73</b>	1,590	923	4,807	<b>7,323</b>	1	0	2	<b>3.0</b>
		Gramene61-IEA	26.47	55.03	48.72	<b>70.23</b>	38,593	109,013	109,507	<b>257,133</b>	0	1	1	<b>2.0</b>
<i>Vigna unguiculata</i>	29,773	GOMAP	91.21	91.08	100.00	<b>100.00</b>	74,791	67,734	242,847	<b>385,372</b>	2	2	6	<b>11.0</b>
<i>Zea mays</i> B73.v3	39,469	GOMAP	88.33	96.41	99.99	<b>100.00</b>	134,622	87,007	290,824	<b>512,453</b>	3	2	6	<b>11.0</b>
		GoldStandard	3.89	0.15	0.38	<b>4.10</b>	1,554	65	299	<b>1,918</b>	1	0	0	<b>1.0</b>
		Gramene49	29.98	45.58	40.03	<b>55.55</b>	20,066	30,936	30,084	<b>81,086</b>	1	1	1	<b>3.0</b>
		Phytozome	11.46	34.77	28.79	<b>40.87</b>	4,787	18,966	13,100	<b>36,853</b>	0	1	1	<b>2.0</b>
<i>Zea mays</i> B73.v4	39,324	GOMAP	93.16	94.92	100.00	<b>100.00</b>	87,648	81,665	278,305	<b>447,618</b>	2	2	6	<b>10.0</b>
		GoldStandard	21.23	25.60	30.82	<b>38.07</b>	11,505	14,986	25,732	<b>52,385</b>	1	1	1	<b>3.0</b>
		Gramene61-IEA	37.12	55.97	60.94	<b>74.11</b>	19,870	47,547	58,093	<b>126,003</b>	1	1	2	<b>3.0</b>
<i>Zea mays</i> Mo17	38,620	GOMAP	86.98	90.87	100.00	<b>100.00</b>	86,074	78,650	277,395	<b>442,119</b>	2	2	6	<b>10.0</b>
		GoldStandard	3.22	0.14	0.35	<b>3.42</b>	1,266	64	277	<b>1,607</b>	1	0	0	<b>1.0</b>
<i>Zea mays</i> PH207	40,557	GOMAP	86.55	90.61	100.00	<b>100.00</b>	88,962	84,910	288,208	<b>462,080</b>	2	2	6	<b>10.0</b>
		GoldStandard	3.15	0.14	0.33	<b>3.34</b>	1,302	63	266	<b>1,631</b>	1	0	0	<b>1.0</b>
<i>Zea mays</i> W22	40,690	GOMAP	90.77	92.58	100.00	<b>100.00</b>	93,622	84,450	289,364	<b>467,436</b>	2	2	6	<b>10.0</b>
		GoldStandard	2.92	0.13	0.29	<b>3.08</b>	1,205	59	241	<b>1,505</b>	1	0	0	<b>1.0</b>

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<sup>a</sup> How many genes in the genome have at least one GO term from the CC, BF, MP aspect annotated to them? A = How many at least one from any aspect? (A = CC ∪ BF ∪ MP)<sup>b</sup> How many annotations in the CC, BF, and MP aspect does this dataset contain? A = How many in total? A = CC + BF + MP<sup>c</sup> Take a typical gene that is present in the annotation set. How many annotations does it have in each aspect? A = How many in total? Please note that A ≠ CC + BF + MP

65 conclusion this means that truly making a statement about the quality of a prediction set would require the  
66 ideal and complete gold standard. The scores we can generate so far are by far not as meaningful.

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**Table 3.** Quality evaluation of the used GO annotation sets.

Genome	Dataset	SimGIC2			TC-AUCPCR		
		CC	BF	MP	CC	BF	MP
<i>Medicago truncatula</i> A17	GOMAP	0.260687	0.192466	0.091357	0.000332	0.000573	0.000789
	Gramene62-IEA	0.755529	0.241984	0.265019	0.000702	0.000736	0.011422
<i>Oryza sativa</i>	GOMAP	0.489140	0.460077	0.214754	0.282023	0.265899	0.143394
	Gramene61-IEA	0.399104	0.422182	0.326020	0.170803	0.255999	0.127307
<i>Triticum aestivum</i>	GOMAP	0.455167	0.421998	0.202177	0.014502	0.005619	0.009458
	Gramene61-IEA	0.367460	0.352624	0.194847	0.004466	0.006177	0.004972
<i>Zea mays</i> B73.v3	GOMAP	0.198179	0.359422	0.094640	0.034380	0.001644	0.001021
	Gramene49	0.252350	0.365922	0.163637	0.049989	0.003238	0.001972
	Phytozome	0.152410	0.355794	0.100792	0.013784	0.003064	0.000539
<i>Zea mays</i> B73.v4	GOMAP	0.486568	0.440989	0.213537	0.283686	0.247096	0.135719
	Gramene61-IEA	0.349744	0.416108	0.324970	0.157917	0.230439	0.131515
<i>Zea mays</i> Mo17	GOMAP	0.232016	0.292290	0.104776	0.040932	0.001932	0.000895
<i>Zea mays</i> PH207	GOMAP	0.234422	0.272876	0.096926	0.037857	0.001743	0.000813
<i>Zea mays</i> W22	GOMAP	0.235595	0.276437	0.103866	0.038546	0.001730	0.000868

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