

# Transcriptome from saffron (*Crocus sativus*) plants in Jammu and Kashmir reveals abundant soybean mosaic virus transcripts and several putative pathogen bacterial and fungal genera

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## Abstract

Saffron (*Crocus sativus*) is a spice with immense economic and medicinal relevance, due to its anticancer and chemopreventive properties. Although the genomic sequence of saffron is not publicly available, the RNA-seq based transcriptome of saffron from Jammu and Kashmir provides several, yet explored, insights into the metagenome of the plant from that region. In the current work, sequence databases were created in the YeATS suite from the NCBI and Ensembl databases to enable faster comparisons. These were used to determine the metagenome of saffron. Soybean mosaic virus, a potyvirus, was found to be abundantly expressed in all five tissues analyzed. Recent studies have highlighted that issues arising from latent potyvirus infections in saffron is severely underestimated. Bacterial and fungal identification is made complex due to symbiogenesis, especially in the absence of the endogenous genome. Symbiogenesis results in transcripts having significant homology to bacterial genomes and eukaryotic genomes. A stringent criterion based on homology comparison was used to identify bacterial and fungal transcripts, and inferences were constrained to the genus level. *Leifsonia*, *Elizabethkingia* and *Staphylococcus* were some of the identified bacteria, while *Mycosphaerella* and *Pyrenophora* were among the fungi detected. Among the bacterial genera, *L. xyli* is the causal agent for ratoon stunting disease in sugarcane, while *E. meningoseptica* and *S. haemolyticus*, having acquired multiresistance against available antimicrobial agents, are important in clinical settings. *Mycosphaerella* and *Pyrenophora* incorporate several pathogenic species. It is shown that a transcript from heat shock protein of the fungi *Cladosporium cladosporioides* has been erroneously annotated as a saffron gene. The detection of these pathogens should enable proper strategies for ensuring better yields. The functional annotation of proteins in the absence of a genome is subject to errors due to the existence of significantly homologous proteins in organisms from different branches of life.

## Keywords

Computational genomics ; RNA-seq ; transcriptome ; saffron ; *Crocus sativus* ; mosaic viruses ; metagenome ; Symbiogenesis

## Introduction

Saffron, possibly one of the most expensive spices, is an anticancer and chemopreventive agent [1,2], and is also effective for treating major depressive disorder symptoms [3,4]. The genome of saffron is not publicly available, although <http://www.crocusgenome.org/> provides an online interface to query individual sequences. The RNA-seq transcriptome [5,6] of saffron was independently sequenced by two research groups in order to gain deeper insights into the genes involved in apocarotenoid biosynthesis [7,8]. Previously, the YeATS suite identified several artifacts arising from RNA-seq assembly [9,10], and has been used to analyze the walnut transcriptome revealing the biodiversity and plant-microbe interactions in twenty different tissues from walnut in California [11].

A BLAST to the complete 'nt/nr' database, as done previously, is not time efficient [9–11]. In the current work, the NCBI and Ensembl database was used to create smaller, yet comprehensive, databases for viruses (V-DB), bacterial (B-DB), fungal (F-DB) and plants (mitochondria, chloroplast and ribosomes - CMR-DB), and were added to the YeATS suite for a time efficient query of the metagenome. In the absence of a genome, identification of saffron derived transcripts is not straightforward. Using these databases, the metagenome of saffron from Jammu and Kashmir was derived from the transcriptome of saffron. <http://nipgr.res.in/mjain.html?page=saffron> provided the ABySS [12] assembled transcriptome with 105269 contigs. Mosaic virus derived transcripts were found to be abundantly expressed in all five tissues analyzed [13]. Mosaic viruses can be responsible for significant yield losses [14]. Key organelles of eukaryotes, like the mitochondrion and chloroplast, possibly originated as a symbiosis between distinct single-celled organisms [15]. Symbiogenesis results in transcripts that have significant homology to both bacterial and eukaryotic genomes. Consequently, a stringent constraint that compared homology scores was used to choose bacterial and fungal transcripts. *Leifsonia*, *Elizabethkingia* and *Staphylococcus* were some of the identified bacteria, while *Mycosphaerella* and *Pyrenophora* were among the fungi detected. These bacterial and fungal genera incorporate several pathogenic species. A proper disease-management strategy can be devised based on this knowledge. It is also shown that protein-based annotation, based on open reading frames of transcripts, can be error prone due to the existence of significantly homologous proteins in organisms from different branches of life, like fungi, plants and viruses.

## Materials and methods

### Obtaining transcriptome and expression matrix from different tissues of saffron

<http://nipgr.res.in/mjain.html?page=saffron> provided the ABySS [12] assembled transcriptome with 105269 contigs. The site provides the sequences (Saffron\_transcriptome\_assembly.fa), annotation (Saffron\_transcriptome\_assembly.fa) and expression values for different tissues (Saffron\_expression\_matrix).

### Obtaining the viral sequences:

“<http://www.ncbi.nlm.nih.gov/genomes/> GenomesGroup.cgi?taxid=10239” provides a retrieve option to get all virus genomes (viral.nt.fa:n=7331 in Dataset1). For viral protein sequences, the NCBI database was queried for “Viruses, RefSeq” (viral.nr.refseq.fa, n=50021 in Dataset1)

### Obtaining the bacterial genomes:

“ftp://ftp.ncbi.nlm.nih.gov/genomes/refseq/bacteria/assembly\_summary.txt” provides details of available bacteria genomes. This was parsed, and the first occurring species of a genus was chosen randomly (getBacterialGenomes.csh:n=1355 in Dataset1).

### Obtaining plant mitochondrial, chloroplast and ribosomal sequences:

The NCBI database was queried for : Plants, RefSeq, Mitochondrion/Chloroplast/Ribosomal rRNA/Ribosomal mRNA. These were combined in a single file (DB.list\_CHLORO\_MITO\_MRIBO\_RRIBO, n=40049 in Dataset1).

### Obtaining plant mitochondrial, chloroplast and ribosomal sequences:

The fungal sequences were obtained from the Ensembl site [16]. A random species was chosen for each genus (list.ensembl.fungi.txt in Dataset1, n=222).

The YeATS suite was used extensively to query these databases using the BLAST command-line interface [17]. The BLAST bitscore was used as a comparison metric instead of the Evalue since it allows differentiation for high homologies where Evalue goes to zero.

## Results and discussion

### Viral transcripts:

Transcripts that map to virus genomes (V-DB) clearly demonstrates the presence of the several mosaic virus (MV) species (Table 1). MV are not a single taxon, although they are all ssRNA positive-strand viruses [13]. For example, tobacco MV is a Tobamovirus, while all other MVs in Table 1 are Potyviruses. Previously, saffron has been shown to be infected with bean yellow MV [18] and Narcissus MV [19]. *Iris*, which belong to the same *Iridaceae* family as saffron, was shown to be infected with *Iris* severe MV [20,21] and Narcissus MV [22]. A recent study highlights that the threat from these MVs is seriously underestimated [23].

The complete list of virus genome matches with a BLAST bitscore (BBS) cutoff=150 (~Evalue=1E-30) (see viral.anno.nucleotide.txt in Dataset1, n=8). Transcripts with the highest significance (Table 1) were BLAST'ed through the online interface to the saffron genome. Only CSTC028935 had a significant match to "gb|EX144186.1|EX144186 cr28.F19 Saffron (*Crocus sativus*) mature stigma lambda Unizap", with a 97% identity, although it is unlikely that this is part of the saffron genome. CSTC028935 encodes a 1587 long open reading frame (ORF) that is 79% identical to the soybean MV polyprotein (Accid:ADK60773.1). Expectedly, none of these viral transcripts were annotated previously in Saffron\_functional\_annotation [7].

### Bacterial transcripts:

The endosymbiont theory posits that key organelles of eukaryotes, like the mitochondrion and chloroplast, originated as a symbiosis between distinct single-celled organisms [15]. While the origin of this symbiosis remains a subject of debate, the fact that prokaryotes and eukaryotes share significant homology in their genomes is established beyond doubt [24]. This presents a certain degree of uncertainty in identifying the metagenome through nucleotide sequencing, especially when the genome of the endogenous organism is not known. This uncertainty is demonstrated in the analysis below.

First, the saffron transcripts were BLAST'ed to the locally created bacterial genome database (B-DB) (1355 bacterial genomes, see Methods). There were 430 matching transcripts (BBS=200, Evalue cutoff=~1E-50, see BACT.200.anno.sort in Dataset1). A combined database comprising chloroplast, mitochondrial and ribosomal genes (CMR-DB) using a more relaxed Evalue constraint gave 573 matching transcripts (BBS=60, Evalue=1E-10, CHLOROMITORIBO.60.anno.sort in Dataset1). Since the organism under focus here is a plant, the Evalue is more relaxed for these matches. In case a transcript matched both B-DB and CMR-DB, the percentage difference in the BBS was used as a metric to select bacterial genes. There

were about 150 transcripts where the B-DB BBS score was 60% greater than than the CMR-DB score. About 80 transcripts had no match in CMR-DB. Combined with the high cutoff required for bacterial identification, and low threshold for CMR-DB assignment, these 230 transcripts can be assigned to bacterial species (BACT.MITOCHLORO.anno.sort in Dataset1).

Bacterial transcripts sorted based on BBS revealed the presence of several bacterial genera (Table 2). *Enterococcus* is a large genus of lactic acid bacteria, of which *E. faecalis* and *E. faecium* are common commensal organisms in the intestines of humans. *E. silesiacus* was found in two water isolates [25]. Similarly, *Staphylococcus haemolyticus* is the most frequent aetiological factors of staphylococcal infections, attributed to its ability to acquire multiresistance against available antimicrobial agents [26]. *Leifsonia xyli* is the causal agent of ratoon stunting disease in sugarcane [27], and is surprisingly genetically conserved in isolates from different countries [28]. The commonly occurring gram-negative bacillus *Elizabethkingia* is found in soil, river water and reservoirs. *Elizabethkingia meningoseptica* is resistant to most antibiotics used in the intensive care setting [29], and has been responsible for an outbreak arising from tap contamination [30]. A recent case of bacteremia has been attributed to *Elizabethkingia meningoseptica* [31].

An example of a transcript with significant homology to bacterial and plant genomes is CSTC008113. This has the best match in the 'nt' database to *Vicia americana*, a legume known commonly as American vetch and purple vetch, with BBS=957. Among the bacterial genomes, this matches to *Tenacibaculum mesophilum* with BBS=776. This has no match in the 'nt' database when the search is constrained to bacteria, demonstrating that all bacterial genomes are not included in the 'nt' database.

### Fungal transcripts:

Fungal transcripts (BBS=150, annoEnsFungi.anno.sort in Dataset1, n=795) reveal several commonly occurring genera, detected in small traces in different tissues (Table 6). The exact species can not be identified using the available data, since these genera incorporate many species. <http://www.speciesfungorum.org/Names/names.asp> provides the enumeration of fungal species. For example, *Mycosphaerella*, a genus of ascomycotaes, is one of the largest genus of plant pathogen fungi with more than 1500 species [32]. Similarly, *Pyrenophora* has ~160 species, several of them pathogenic [33]

### Caveats and advantages of ORF based annotation:

Annotation of transcripts can also be done by analyzing ORFs. Although the nucleotide sequence of CSTC042320 has no match in the complete 'nt' database, it encodes a 479 long ORF homologous (57% identity) to a cytochrome P450 protein from pineapple (*Ananas comosus*). The saffron transcriptome has been annotated accordingly (Saffron\_functional\_annotation [7]), and has significant match in the online interface to the saffron genome ([http://www.crocusgenome.org/?page\\_id=20](http://www.crocusgenome.org/?page_id=20)). However, such functional annotation of proteins in the absence of a genome is subject to errors due to the existence of significantly homologous proteins in organisms from different branches of life, like fungi, plants and viruses. CSTC018609 is one such mis-annotated transcripts, that encodes an 213 long ORF, homologous to a heat shock protein from the fungi *Cladosporium cladosporioides* with 99% identity, but is annotated as a plant gene (Saffron\_functional\_annotation [7]). The nucleotide sequence of CSTC018609 also matches to the genome of the fungi *Cladosporium cladosporioides* (Accid:HQ693482.1) with BBS=1094, 97% identity. Although the nucleotide sequence also matches to a plant (*Nicotiana tabacum*, Accid:AY372070.1) with lesser significance (87% identity), it has no significant match through the online interface provided. Interestingly, the ORF of this gene is also homologous (76% identity) to a viral HSP70 protein (*Micromonas pusilla* virus, Accid:AET43623.1).

ORF based annotation provides coverage in some cases where nucleotide homology is not detected. The viral protein RefSeq database identified additional ~20 putative viral transcripts (see viral.anno.protein.txt in Dataset1) not detected through the nucleotide comparison. One reason for such an exclusion is the smaller size of the local viral nucleotide database sequence, which does not include all strains. CSTC027157 has significant matches among the full 'nt' virus database (bean common mosaic virus complete genome, isolate

PV 0915, Accid:HG792064.1). However, some sequences (CSTC004726) have no match even in the complete 'nt' database, although they encode ORFs with significant matches to the local viral protein database. CSTC004726 encodes a polyprotein (Accid:NP\_660175.1, Evalue=9e-37) from bean common mosaic necrosis virus (Evalue=9e-37). Thus, an annotation methodology should include comparisons to both nucleotide and amino acid sequences.

## Conclusions and future work:

Previous application of the YeATS suite to genomes [9,10] and metagenomes [11] required a BLAST to the complete 'nt' and 'nr' databases. In the current work, smaller, yet comprehensive databases have been created to accelerate the identification of the saffron metagenome from the transcriptome obtained from saffron in Jammu and Kashmir [7]. Transcripts from the soybean mosaic virus was found abundantly expressed in all five tissues analyzed. Although, these viruses have been isolated from these plants (and the related species *Iris*) previously [18–22], recent studies have shown the threat posed by these viruses to saffron cultivation is seriously underestimated. Several bacterial and fungal genera incorporating known pathogen species have also been identified.

A search through the complete set of sequences known (through a program like BLAST) is ideal, but not time efficient. The search can be accelerated by creating smaller, but comprehensive, sequence databases. Such a 'divide and conquer' strategy needs to account for the existence of homologous nucleotide sequences (and proteins) across the different trees of life. There are two methods for annotating a transcript - (a) through the nucleotide sequence or (b) through the amino acid sequence (assuming it is not a non-coding transcript). Annotation based on ORFs identifies transcripts that have no nucleotide sequence homology in the entire 'nt' database, which occurs due to the redundancy in the codon table. Finally, this work also highlights that the standard practice of excluding transcripts homologous to ribosomal RNA might exclude bona fide bacterial transcripts, in the absence of comparative analysis of the homology to different databases. While the current strategy of creating smaller, comprehensive subsets accelerates the annotation time as compared to YeATS, the existing flow still uses command-line BLAST to query the entire transcriptome with the sequence databases. A kmer based approach could accelerate this further, and will be added in a future release [34].

## Competing interests

No competing interests were disclosed.

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## References

1. Escribano J, Alonso GL, Coca-Prados M, Fernández JA (1996) Crocin, safranal and picrocrocin from saffron (*crocus sativus* l.) inhibit the growth of human cancer cells in vitro. *Cancer letters* 100: 23–30.

2. Abdullaev FI (2002) Cancer chemopreventive and tumoricidal properties of saffron (*crocus sativus* l.). *Experimental biology and medicine* 227: 20–25.
3. Lopresti AL, Drummond PD (2014) Saffron (*crocus sativus*) for depression: a systematic review of clinical studies and examination of underlying antidepressant mechanisms of action. *Human Psychopharmacology: Clinical and Experimental* 29: 517–527.
4. Hausenblas HA, Saha D, Dubyak PJ, Anton SD (2013) Saffron (*crocus sativus* l.) and major depressive disorder: a meta-analysis of randomized clinical trials. *Journal of integrative medicine* 11: 377–383.
5. Wang Z, Gerstein M, Snyder M (2009) RNA-seq: a revolutionary tool for transcriptomics. *Nature Reviews Genetics* 10: 57–63.
6. Flintoft L (2008) Transcriptomics: digging deep with RNA-seq. *Nature Reviews Genetics* 9: 568–568.
7. Jain M, Srivastava PL, Verma M, Ghangal R, Garg R (2016) De novo transcriptome assembly and comprehensive expression profiling in *crocus sativus* to gain insights into apocarotenoid biosynthesis. *Scientific reports* 6.
8. Baba SA, Mohiuddin T, Basu S, Swarnkar MK, Malik AH, et al. (2015) Comprehensive transcriptome analysis of *crocus sativus* for discovery and expression of genes involved in apocarotenoid biosynthesis. *BMC genomics* 16: 1.
9. Chakraborty S, Britton M, Wegrzyn J, Butterfield T, Martinez-Garcia PJ, et al. (2015). YeATS—a tool suite for analyzing RNA-seq derived transcriptome identifies a highly transcribed putative extensin in heartwood/sapwood transition zone in black walnut.
10. Martínez-García PJ, Crepeau MW, Puiu D, Gonzalez-Ibeas D, Whalen J, et al. (2016) The walnut (*juglans regia*) genome sequence reveals diversity in genes coding for the biosynthesis of nonstructural polyphenols. *The Plant Journal* .
11. Chakraborty S, Britton M, Martínez-García P, Dandekar AM (2016) Deep RNA-seq profile reveals biodiversity, plant–microbe interactions and a large family of NBS-LRR resistance genes in walnut (*juglans regia*) tissues. *AMB Express* 6: 1.
12. Simpson JT, Wong K, Jackman SD, Schein JE, Jones SJ, et al. (2009) Abyss: a parallel assembler for short read sequence data. *Genome research* 19: 1117–1123.
13. Koonin EV, Dolja VV, Morris TJ (1993) Evolution and taxonomy of positive-strand rna viruses: implications of comparative analysis of amino acid sequences. *Critical reviews in biochemistry and molecular biology* 28: 375–430.
14. Byamukama E, Wegulo SN, Tatineni S, Hein G, Graybosch R, et al. (2014) Quantification of yield loss caused by triticum mosaic virus and wheat streak mosaic virus in winter wheat under field conditions. *Plant Disease* 98: 127–133.
15. Margulis L (1996) Archaeal-eubacterial mergers in the origin of eukarya: phylogenetic classification of life. *Proceedings of the national academy of sciences* 93: 1071–1076.
16. Yates A, Akanni W, Amode MR, Barrell D, Billis K, et al. (2016) Ensembl 2016. *Nucleic acids research* 44: D710–D716.
17. Camacho C, Madden T, Ma N, Tao T, Agarwala R, et al. (2013) BLAST Command Line Applications User Manual .

18. Russo M, Martelli G, Cresti M, Ciampolini F (1979) Bean yellow mosaic virus in saffron/il virus del mosaico giallo del fagiolo in zafferano. *Phytopathologia Mediterranea* : 189–191.
19. Miglino R, Jodlowska A, Van Schadewijk A (2005) First report of narcissus mosaic virus infecting crocus spp. cultivars in the netherlands. *Plant Disease* 89: 342–342.
20. Yan S, Qin Z, Jin L, Chen J (2010) A new isolate of iris severe mosaic virus causing yellow mosaic in iris ensata thunb. *Journal of nanoscience and nanotechnology* 10: 726–730.
21. Nateqi M, Habibi MK, Dizadji A, Parizad S (2015) Detection and molecular characterization of the iris severe mosaic virus-ir isolate from iran. *Journal of Plant Protection Research* 55: 235–240.
22. Wylie SJ, Li H, Liu J, Jones MG (2014) First report of narcissus mosaic virus from australia and from iris. *Australasian Plant Disease Notes* 9: 1–2.
23. Caiola MG, Faoro F (2011) Latent potyvirus infections in crocus sativus artwrightianus: an underestimated problem in saffron? *Phytopathologia Mediterranea* 50: 175–182.
24. Doolittle WF (1998) A paradigm gets shifty. *Nature* 392: 15–16.
25. Švec P, Vancanneyt M, Sedláček I, Naser SM, Snauwaert C, et al. (2006) *Enterococcus silesiacus* sp. nov. and *enterococcus termitis* sp. nov. *International journal of systematic and evolutionary microbiology* 56: 577–581.
26. Czekaj T, Ciszewski M, Szewczyk EM (2015) *Staphylococcus haemolyticus*—an emerging threat in the twilight of the antibiotics age. *Microbiology* 161: 2061–2068.
27. Grisham M, Pan YB, Richard Jr E (2007) Early detection of *leifsonia xyli* subsp. *xyli* in sugarcane leaves by real-time polymerase chain reaction. *Plant Disease* 91: 430–434.
28. Young A, Petrasovits L, Croft B, Gillings M, Brumbley S (2006) Genetic uniformity of international isolates of *leifsonia xyli* subsp. *xyli*, causal agent of ratoon stunting disease of sugarcane. *Australasian Plant Pathology* 35: 503–511.
29. Jean S, Lee WS, Chen F, Ou T, Hsueh P (2014) *Elizabethkingia meningoseptica*: an important emerging pathogen causing healthcare-associated infections. *Journal of Hospital Infection* 86: 244–249.
30. Balm M, Salmon S, Jureen R, Teo C, Mahdi R, et al. (2013) Bad design, bad practices, bad bugs: frustrations in controlling an outbreak of *elizabethkingia meningoseptica* in intensive care units. *Journal of Hospital Infection* 85: 134–140.
31. Shinha T, Ahuja R (2015) Bacteremia due to *elizabethkingia meningoseptica*. *IDCases* 2: 13–15.
32. Crous P, Braun U, Groenewald J (2007) *Mycosphaerella* is polyphyletic. *Studies in Mycology* 58: 1–32.
33. Day J, Gietz RD, Rampitsch C (2015) Proteome changes induced by *pyrenophora tritici-repentis* toxa in both insensitive and sensitive wheat indicate senescence-like signaling. *Proteome science* 13: 1.
34. Compeau PE, Pevzner PA, Tesler G (2011) How to apply de bruijn graphs to genome assembly. *Nature biotechnology* 29: 987–991.

**Table 1: Transcripts mapping to viruses sorted based on BLAST bitscore (BBS):** There are several transcripts which match to mosaic viruses (MV). Only CSTC028935 has a significant match (gb|EX144186.1) to the unpublished saffron genome, queried through the online interface ([http://www.crocusgenome.org/?page\\_id=20](http://www.crocusgenome.org/?page_id=20)). It is very unlikely that the transcript is a part of the saffron genome since it encodes an open reading frame (1587 amino acid long) with a very significant match (79% identity) to the soybean MV polyprotein (Accid:ADK60773.1).

TranscriptID	Accession ID	Description	BBS
CSTC028935	NC_003397.1.	Bean common MV	1328
CSTC098263	NC_001367.1.	Tobacco MV	952
CSTC029938	NC_001367.1.	Tobacco MV	876
CSTC042852	NC_007216.1.	Wisteria vein MV	651
CSTC006658	NC_007216.1.	Wisteria vein MV	608
CSTC103119	NC_001367.1.	Tobacco MV	553
CSTC003939	NC_007216.1.	Wisteria vein MV	459
CSTC045789	NC_009742.1.	Telosma MV	176

**Table 2: Bacterial transcripts sorted based on BLAST bitscore (BBS):** These were identified by comparing the saffron transcriptome to the local bacterial database (B-DB). The homology of these bacterial transcripts to the CMR-DB (Chloroplast-Mitochondrion-Ribosome-DB) provides support for the endosymbiont theory. The transcripts matching the bacterial database were subsequently BLAST'ed to the complete 'nt' database and the 'plant nt' database. Although, these bacterial transcripts also have significant matches to plant genomes, they are significantly lower than the matches to bacterial genomes. Interestingly, apart from the only mitochondrial transcript CSTC048724 which had a different genus (*Enhydrobacter* vs *Moraxella*), all other were chloroplast transcripts, and had the same genus when compared to the selected bacterial database versus the complete 'nt' database. As expected, none of these transcripts were annotated previously [7].

Transcript ID	DB	Accession ID	Description	BBS
CSTC006383	B-DB	NC.004668.1	<i>Enterococcus faecalis</i> V583 chromosome	3659
	nt.Bact	gb CP013614.1	<i>Enterococcus silesiacus</i>	3947
	nt.Plants	L42852.1	<i>Pseudoscourfieldia marina</i> large subunit ribosomal rna	706
	CMR-DB	NC.000927.1	<i>Nephroselmis olivacea</i> chloroplast	686
CSTC017835	B-DB	NC.007622.1	<i>Staphylococcus aureus</i> RF122	3175
	nt.Bact	gb CP011116.1	<i>Staphylococcus haemolyticus</i>	3312
	nt.Plants	KX013545.1	<i>Cymbomonas tetramitiformis</i> chloroplast	468
	CMR-DB	NC.030169.1	<i>Cymbomonas tetramitiformis</i> chloroplast	468
CSTC036051	B-DB	NC.006087.1	<i>Leifsonia xyli</i> subsp. <i>xyli</i> str.	3003
	nt.Bact	ref NR_076082.1	<i>Leifsonia xyli</i> subsp. <i>xyli</i> 23S ribosomal RNA	3003
	nt.Plants	FN563100.1	<i>Cymbomonas tetramitiformis</i> plastid 16s rna gene	486
	CMR-DB	NC.030169.1	<i>Cymbomonas tetramitiformis</i> chloroplast	481
CSTC075959	B-DB	NZ_BARD01000031.1	<i>Elizabethkingia meningoseptica</i>	2963
	nt.Bact	gb JX219387.1	<i>Elizabethkingia</i> sp. 16S ribosomal RNA gene	3016
	nt.Plants	DQ455171.1	<i>Medicago truncatula</i>	405
	CMR-DB	NC.023835.1	<i>Chlorella sorokiniana</i> chloroplast	318
CSTC038433	B-DB	NC.002516.2	<i>Pseudomonas aeruginosa</i> chromosome	2937
	nt.Bact	gb CP002881.1	<i>Pseudomonas stutzeri</i>	3386
	nt.Plants	DQ115326.1	<i>Gossypium hirsutum</i>	944
	CMR-DB	NC.025547.1	<i>Pseudochloris wilhelmii</i> chloroplast	294
CSTC012158	B-DB	NC.016818.1	<i>Rahnella aquatilis</i> CIP 78.65 = ATCC 33071	2850
	nt.Bact	gb CP003403.1	<i>Rahnella aquatilis</i> HX2	2850
	nt.Plants	DQ115326.1	<i>Gossypium hirsutum</i>	826
	CMR-DB	NC.025547.1	<i>Pseudochloris wilhelmii</i> chloroplast	351
CSTC023991	B-DB	NC.010611.1	<i>Acinetobacter baumannii</i>	2828
	nt.Bact	gb CP012608.1	<i>Acinetobacter</i> sp.	2952
	nt.Plants	DQ115326.1	<i>Gossypium hirsutum</i>	1417
	CMR-DB	NC.025547.1	<i>Pseudochloris wilhelmii</i> chloroplast	303
CSTC048724	B-DB	NZ_ACYI01000006.1	<i>Enhydrobacter aerosaccus</i>	2819
	nt.Bact	gb KT865797.1	<i>Moraxella</i> sp. 23S ribosomal RNA gene	2824
	nt.Plants	DQ115326.1	<i>Gossypium hirsutum</i>	1421
	CMR-DB	NC.022863.1	<i>Roya obtusa</i> mitochondrion	296
CSTC038044	B-DB	NZ_ACQT01000330.1	<i>Acidovorax delafieldii</i>	2763
	nt.Bact	gb KT865866.1	<i>Acidovorax</i> sp. 23S ribosomal RNA gene	2789
	nt.Plants	DQ115326.1	<i>Gossypium hirsutum</i>	473
	CMR-DB	NC.025547.1	<i>Pseudochloris wilhelmii</i> chloroplast	292
CSTC028569	B-DB	NC.011071.1	<i>Stenotrophomonas maltophilia</i> R551-3	2737
	nt.Bact	gb CP016294.1	<i>Stenotrophomonas rhizophila</i>	2896
	nt.Plants	DQ115326.1	<i>Gossypium hirsutum</i>	621
	CMR-DB	NC.025547.1	<i>Pseudochloris wilhelmii</i> chloroplast	246

Table 3: **Counts of transcripts mapping to viruses (V) and bacteria (B) sorted based on BLAST bitscore:** Viral transcripts from mosaic viruses are abundantly present in all five tissues analyzed. In comparison, bacterial transcripts are fewer in numbers.

	TranscriptID	TrLen	CORM	TEPAL	LEAF	STIGMA	STAMEN
V	CSTC028935	4762	167821.9	242246.9	163341.6	1067049.2	29630.3
	CSTC098263	781	1	49.9	0	0	0
	CSTC029938	889	0	85.4	0	0	0
	CSTC042852	7175	78307.1	87094.2	62997.3	289732.9	8391.9
	CSTC006658	3519	40341.8	86292.2	48059.2	237711.2	4193.8
	CSTC103119	759	0	60.4	0	0	0
	CSTC003939	2758	11938.7	34691.7	8590.5	51597	1335.8
	CSTC045789	558	25667.9	8440	10437.1	69069.3	2162.8
B	CSTC006383	2924	430.7	785.7	25.7	280.2	263.7
	CSTC017835	1898	247.1	221.6	38.0	139.4	8.6
	CSTC036051	2315	177.3	307.0	19.5	14.3	12.9
	CSTC075959	1885	263.8	148.7	129.3	175.8	14.3
	CSTC038433	1864	896.8	196.7	70.8	220.2	30.1
	CSTC012158	1646	472.4	490.2	16.4	362.4	322.5
	CSTC023991	1780	810.2	497.9	119.1	203.7	10.0
	CSTC048724	1648	199.2	251.3	32.9	52.2	21.5
	CSTC038044	1761	248.2	270.5	40.0	215.9	18.6
	CSTC028569	2054	115.8	159.2	17.4	117.2	18.6

Table 4: **Transcripts with the highest cumulative counts :** Apart from one viral transcript (CSTC028935), and one cytochrome P450 encoding ORF (CSTC042320), all other transcripts are ribosomal RNA or from the chloroplast/mitochondria. These eight transcripts also have significant matches to the bacterial database, although their homology to the plant genomes is significantly higher. The cytochrome P450 transcript is annotated by analyzing the open reading frame, since the nucleotide sequence has no match in the complete 'nt' database.

TranscriptID	CORM	TEPAL	LEAF	STIGMA	STAMEN	Description
CSTC043388	1311047.7	4516814.8	423456.4	134222.0	1403873.5	26S rRNA
CSTC048227	642066.9	2044589.8	255631.2	77501.7	509226.4	18S rRNA
CSTC031703	572899.7	1513910.7	150241.9	40571.2	998203.4	26S rRNA
CSTC009284	360533.7	1334369.3	83657.0	23579.3	628958.8	26S rRNA
CSTC028935	167821.9	242246.9	163341.6	1067049.2	29630.3	Bean Mosaic Virus
CSTC018503	212853.0	482137.8	39453.3	13557.8	277618.9	26S rRNA
CSTC042320	57.4	143.9	86.2	805965.3	9905.4	Cytochrome P450 (Protein)
CSTC044767	33477.2	42562.8	696846.6	20970.8	14874.6	Chloroplast
CSTC050677	50947.8	477696.2	23367.5	12243.9	99507.3	Mitochondria
CSTC019287	17553.0	29246.6	467484.7	14051.0	7976.2	Chloroplast

**Table 5: Transcripts with highest counts - comparing the significance of homology to plant and bacterial genomes:** These transcripts are from Table 4, barring a virus derived transcript and the cytochrome P450 gene. They have been BLAST'ed to the complete 'nt' database. The endosymbiont theory is amply supported by the significant matches of these plant mitochondrial and chloroplast transcripts to bacterial genomes.

CSTC043388	AY095471.1	<i>Bulbine succulenta</i> 26S ribosomal RNA gene, partial sequence	2492
CSTC043388	NZ_LDOD01000051.1	<i>Tenacibaculum mesophilum</i> strain HMG1 scaffold51, whole genome	990
CSTC048227	HM640735.1	<i>Hesperocallis undulata</i> voucher Cranfill Schmid s.n.(JEPS) 18S	1664
CSTC048227	NZ_LDOD01000055.1	<i>Tenacibaculum mesophilum</i> strain HMG1 scaffold55, whole genome	178
CSTC031703	KM117261.1	<i>Wisteria floribunda</i> 18S ribosomal RNA gene, internal transcribed	1068
CSTC031703	NZ_ATXL01000109.1	<i>Bavariicoccus seileri</i> DSM 19936 H541DRAFT_scaffold00109.109_C,	549
CSTC009284	AF389262.1	<i>Dicentra eximia</i> 26S ribosomal RNA gene, partial sequence	1033
CSTC009284	NZ_ATXL01000109.1	<i>Bavariicoccus seileri</i> DSM 19936 H541DRAFT_scaffold00109.109_C,	651
CSTC018503	HG970335.2	<i>Fusarium graminearum</i> chromosome 4, complete genome	1993
CSTC018503	NZ_ATXL01000109.1	<i>Bavariicoccus seileri</i> DSM 19936 H541DRAFT_scaffold00109.109_C,	1509
CSTC044767	KP899622.1	<i>Dioscorea zingiberensis</i> voucher DZING20150203V3 chloroplast,	5288
CSTC044767	NC_006576.1	<i>Synechococcus elongatus</i> PCC 6301 DNA, complete genome	1999
CSTC050677	JN375330.1	<i>Phoenix dactylifera</i> mitochondrion, complete genome	3192
CSTC050677	NZ_BBVC01000023.1	<i>Caedibacter varicaedens</i> DNA, contig: Cva_contig000023, whole	638
CSTC019287	KM014691.1	<i>Iris gatesii</i> voucher T06-52 plastid, complete genome	6335
CSTC019287	NZ_GL890965.1	<i>Moorea producens</i> 3L scf52112, whole genome shotgun sequence	544

**Table 6: Fungal transcripts:** These are sorted based on BLAST bitscore (BBS). Although, the best match specifies a genus and the species, this identification needs validation. The counts in the different tissues are lesser than those of viral and bacterial transcripts. STI: STIGMA, STA: STAMEN.

TRID	Accid	Description	BBS	CORM	TEPAL	LEAF	STI	STA
CSTC021516	XM_002619271.1	<i>Clavispora lusitaniae</i>	1897	2	581	0	31	11
CSTC086633	XM_013574385.1	<i>Aureobasidium namibiae</i>	1306	4	345	5	398	25
CSTC072893	XM_001551736.1	<i>Botryotinia fuckeliana</i>	1290	5	360	2	57	1
CSTC101026	DQ677898.1	<i>Cladosporium cladosporioides</i>	1210	1	142	1	64	8
CSTC023760	XM_001930596.1	<i>Pyrenophora tritici-repentis</i>	1051	4	221	0	92	2
CSTC064623	XM_003856981.1	<i>Mycosphaerella graminicola</i>	1083	0	203	1	105	4
CSTC026015	X81860.1	<i>Cladosporium herbarum</i>	1066	4	175	1	103	8
CSTC016711	XM_016737515.1	<i>Penicillium expansum</i>	885	3	99	0	48	0
CSTC041979	XM_003854273.1	<i>Mycosphaerella graminicola</i>	824	2	126	0	101	0