

USE OF BIOINFORMATICS TOOLS TO STUDY PHYLOGENETIC ANALYSIS AND SEQUENCE SIMILARITY OF MALASSEZIA SP. A PATHOGEN INVOLVED IN DANDRUFF.

Abstract

One of the most prevalent species in the human skin microbiota, *Malassezia* species, has been linked to skin conditions like dandruff and seborrheic dermatitis. Although *Malassezia* plays a significant role in common skin diseases, little is understood about its molecular makeup. In this approach, bioinformatics can be quite helpful. Selected genes from *Malassezia* species can be subjected to BLAST P. It was discovered that *M. restricta* shares similarities with the distant human pathogen *Candida albicans* and the plant pathogen *Ustilago maydis* by the use of blast, a bioinformatics tool. It is possible to identify the convergent and divergent features of *Malassezia* species using the bioinformatics tool Clustal W. Researchers must create more potent bioinformatics tools to help manage the growing issue of dandruff and itchiness on the scalp in the human population.

Keywords: Bioinformatics, Microbiota, *Malassezia* Species.

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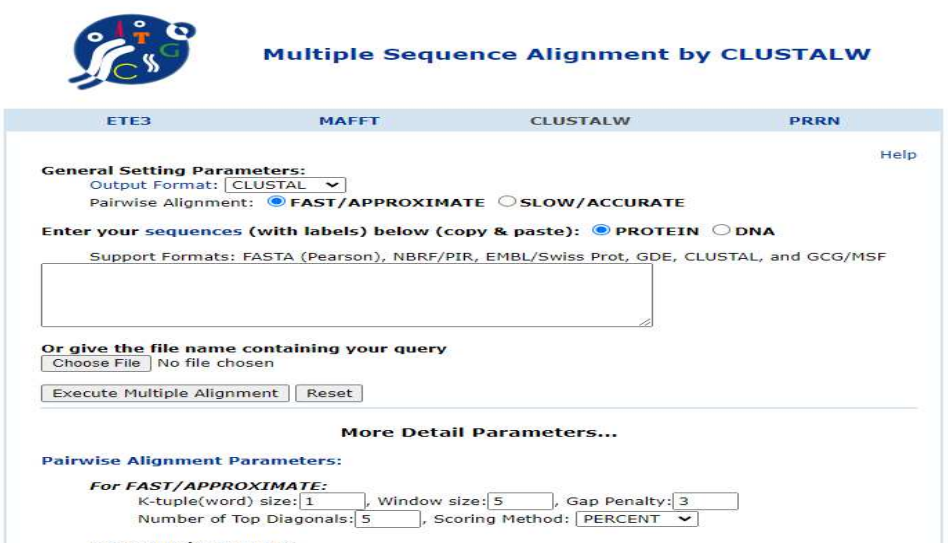
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**IndexTerms—Malassezia,Ustilago,
Candidaalbicans,CLUSTALW,BLAST,Dandruff,scalp pruritus.**

I. INTRODUCTION

One of the most important species of skin microbiota is the Malassezia species, which has been linked to a number of skin conditions, including dandruff and seborrheic dermatitis. The molecular makeup of malassezia is largely unknown. There are currently roughly 18 species of Malassezia known, including M.globosa, M.restricta are the most common species found in humans. malassezia is the eukaryotic biota of the human skin. Seborrheic dermatitis and dandruff are common skin conditions that cause itching and skin flaking. While seborrheic dermatitis is characterized by yellow flakes and irritation, dandruff is characterized by loose flakes and lack of inflammation. The proliferation of the commensal malassezia is the cause of dandruff and other diseases that affect about 50% of individuals. Three components seem to be involved in the etiology of D/SD: metabolism by microbiota, secretions from sebaceous glands, and individual vulnerability. (DeAngelis and others, 2005); Ro and Dawson, 2005). This chapter will describe the most common matches of M.globosa sequence and its phylogenetic analysis using bioinformatics tools BLAST and CLUTALW. BLAST is a tool of NCBI . It finds region of similarity between two or more sequences, the sequences can be either protein or nucleotide. BLAST stands for basic local alignment search tool. Blast is basically used to find out evolutionary and functional relationship between two individuals. it is not a single program but a family of programme like BLAST p , BLAST n , BLAST x, tBLASTn etc. it also helps to identify the member of gene families. CLUTALW is a multiple sequence alignment tool for DNA and protein sequence. it is not a tool for pairwise alignment but generally good for comparing three to four sequences. it is tool of European bioinformatics institute.

The front page of CLUSTALW looks like this



The screenshot shows the web interface for CLUSTALW. At the top, there is a logo with a hand holding a DNA helix and the text "Multiple Sequence Alignment by CLUSTALW". Below the logo, there are four tabs: ETE3, MAFFT, CLUSTALW (selected), and PRRN. The main content area is titled "General Setting Parameters:" and includes a "Help" link. The "Output Format" is set to "CLUSTAL". The "Pairwise Alignment" is set to "FAST/APPROXIMATE". Below this, there is a section for "Enter your sequences (with labels) below (copy & paste):" with radio buttons for "PROTEIN" (selected) and "DNA". A text input field is provided for pasting sequences. Below the input field, there is a "Choose File" button and a "No file chosen" message. There are "Execute Multiple Alignment" and "Reset" buttons. A "More Detail Parameters..." section is partially visible, showing "Pairwise Alignment Parameters:" with fields for "K-tuple(word) size" (set to 1), "Window size" (set to 5), "Gap Penalty" (set to 3), "Number of Top Diagonals" (set to 5), and "Scoring Method" (set to PERCENT).

SOURCE: <https://www.genome.jp/tools-bin/clustalw>

Similarly the front page of blast looks like:

The screenshot shows the NCBI BLAST search interface. At the top, there is a navigation bar with the NIH logo and 'National Library of Medicine' text. Below this, the page is titled 'BLAST® » blastn suite' and 'Standard Nucleotide BLAST'. The main search area includes a text input for 'Enter Query Sequence', a field for 'Enter accession number(s), gi(s), or FASTA sequence(s)', and a 'Query subrange' section with 'From' and 'To' inputs. There are also options to 'Or, upload file', 'Job Title', and 'Align two or more sequences'. The 'Choose Search Set' section has radio buttons for 'Standard databases (nr etc.)', 'rRNA/ITS databases', 'Genomic + transcript databases', 'Betacoronavirus', and 'Experimental databases'. A prominent button says 'Try experimental taxonomic nt databases' with a 'Download' link. A 'Feedback' button is visible on the right side.

Source: https://blast.ncbi.nlm.nih.gov/Blast.cgi?PROGRAM=blastn&PAGE_TYPE=BlastSearch&LINK_LOC=blasthome

1. Material and Methods: The NCBI GENBANK BLAST server version 2.2.30 (<http://blast.ncbi.nlm.nih.gov>) was utilized for alignment and homology searches (altschul et al. 1997). The NCBI data base has complete sequence of all the 9 chromosome of *Malassezia restricta* as well as the *Malassezia restricta* mitochondrion, complete genome. A specific protein synthesized by each chromosome was selected viz lipase protein sequence from chromosome 1 zinc finger domain from chromosome 2, chitin synthetase from chromosome 3 pyruvate synthetase from chromosome 4, carboxyl methyltransferase from chromosome 5, cell division cycle protein 37 of chromosome 6, NADH dehydrogenase (ubiquinone) Fe-S protein 4 of chromosome 7, arginase protein of chromosome 8, DNA repair protein REV1 of chromosome 9 and BLASTp was carried out and homology was identified. The database used for comparison was non redundant protein databases. Multiple sequence alignment by CLUSTAL W with a K tupule word size of 1 was also carried out between secretory lipase enzymes sequences of *Malassezia restricta*, *Ustilago maydis* and *Candida albicans*. A questionnaire and random sampling technique were also used on a sample of around 243 respondents, of which 152 were female and the remaining 91 were male.

II. RESULT AND CONCLUSION

Protein BLAST of lipase protein sequence of chromosome 1 revealed similarity with *Ustilagosps* and most of the smut fungi. The zinc finger domain on chromosome 2 was found to be similar to plant pathogen *Ceratobasidium* sp and *Mycenasps*.

The chitin synthase [*Malassezia restricta*] was found to be highly similar to *Testiculariasps.* and *Scleroderma* sps. and *Ustilagosps*, again all plant pathogen. The pyruvate synthetase gene from chromosome 4 was found to be similar to *Violaceomyces palustris* and *Ustilagosps*. Carboxyl Methyl transferase protein sequence of chromosome 5 was again found to be similar to *Ustilagosps*. Cell division cycle protein 37 protein sequence of chromosome 6 was found to be 98% to *Ustilagosps*. NADH dehydrogenase (ubiquinone) Fe-S protein 4 of chromosome 7 showed 84% similarity with *Ustilagosps*. Arginase protein of chromosome 8 was found to be 98% similar to *Ustilagomyadis*. DNA repair protein REV1 of chromosome 9 was found to be 96% to *Rhizopus* sps. and 63% to *Ustilagosps*. *M. restricta* was found to be more closely related with the plant fungal pathogen *Ustilagomyadis*. A MSA between the secretory lipase of *C. albicans*, *M. restricta* and *U. mayadis* showed *M. restricta* and *U. mayadis* to be more convergent than *C. albicans*.

A detailed summary of the result is given below in the –

Table 1: summary of BLAST P analysis of *M. restricta*

Name	RefSeq	Protein	Functional Protein Selected	SIMILARITY % WITH <i>Ustilagomyadis</i>
Chr	I	835	lipase protein	85%
Chr	II	782	zinc finger domain	Showed similarity with <i>Ceratobasidium</i> sps
Chr	III	746	chitin synthase	
Chr	IV	464	pyruvate synthetase	84%
Chr	V	532	carboxyl methyltransferase	64%
Chr	VI	410	cell division cycle protein 37	98%
Chr	VII	280	NADH dehydrogenase (ubiquinone) Fe-S protein 4	84%
Chr	VIII	233	arginase protein	98%
Chr	IX	108	DNA repair protein REV1	63%
name	RefSeq	protein	Functional Protein Selected	SIMILARITY % WITH <i>Ustilagomyadis</i>

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Chr	IX	108	DNA repair protein REV1	63%

A questionnaire and random sampling technique were also used on a sample of around 243 respondents, of which 152 were female and the remaining 91 were male. 15.9% said they flaked their scalp excessively. Dandruff was shown to be less common as people aged, with a higher incidence in the 25–34 age range. Patients with dandruff were shown to have more severe cases of scalp pruritus than those without dandruff. Among the subjects, antidandruff products of all kinds and home treatments proved to be the least effective.

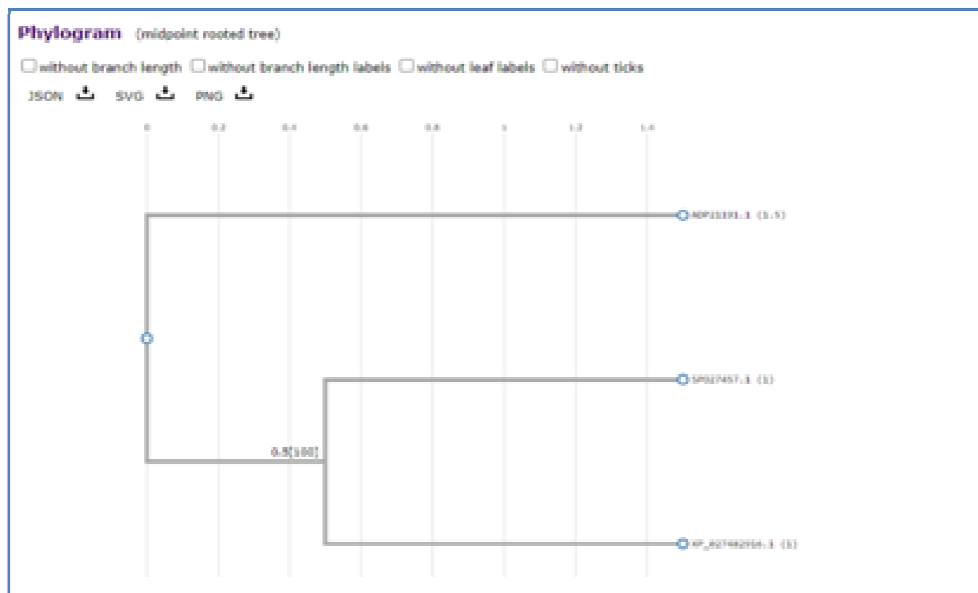


Figure 1: PhyML of Secretory Lipase

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