**BiodEnz: A database of biodegrading enzymes**

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**Abstract:**
Azo dyes, which are characterized by azo bonds, are a predominant class of colorants used in tattooing, cosmetics, foods, textile and consumer products. Laccases (EC 1.10.3.2), lignin peroxidases (EC 1.11.1.14), and Azo reductases (EC 1.7.1.6) of different microorganisms are mainly useful for the development of biodegradation systems as they catalyse reductive cleavage of azo groups (-N=N-) and have very broad substrate specificity with respect to the electron donor and is capable of oxidizing phenols and aromatic amines. Laccases have a very broad substrate specificity with respect to the electron donor and is capable of oxidizing phenols and aromatic amines. Azoreductase belongs to the family of oxidoreductases, acting on other nitrogenous compounds as donors with NAD+ or NADP+ as acceptor. Lignin peroxidase enzymes are highly non-specific and are well reported to decolourize various dyes. We have developed BiodEnz database by collecting information like strains that produce particular enzymes, azo dyes that are degraded, substrate specificity, molecular weight, the optimum temperature and pH, sequence data of the above enzymes, as the most effective inoculants used for bioremediation are able to degrade dyes over a broad concentration range, tolerate a range of environmental conditions of temperature, pH, and activity of the enzymes. The database can be searched by using a user friendly web interface.

**Keywords:** Azo dyes, azolinkages, bioaugmentation, biodegradation, enzyme specificity

**Availability:** [http://www.biodenzdatabase.in](http://www.biodenzdatabase.in)

**Background:**
Azo dyes which represent about one-half of all dyes in common use are widely used by the textile, leather, cosmetics, food coloring and paper production industries. They are considered recalcitrant xenobiotic compounds due to the presence of a nitrogen double bond (-N=N-) bond and other groups (i.e. sulfonic group) that are not easily biodegraded. The annual world production of azo dyes is estimated to be around one million tons [1]. During the dyeing process, approximately 10-15% of the used dye is released into wastewater. Treatment of dye-contaminated wastewater discharged from the textile and other dye-stuff industries is necessary to prevent contamination of soil and surface and ground water. Biological methods are generally considered eco friendly as they can lead to mineralization of organic pollutants effectively at very low cost [1]. Azo dyes are recalcitrant to biodegradation due to their complex structures and xenobiotic nature, and typically require an anaerobic-aerobic process to achieve complete degradation. Bioremediation of azo dyes in textile waste effluents by bacteria or fungi is a very promising area of study because of the relatively low expense involved. Bioremediation includes biodegradation and biotransformation, with a goal to...
mineralize hazardous contaminants in the environment. This is fulfilled by the azodye degrading enzymes like laccases, azoreductases, lignin peroxidases and some more which are vastly available in the variety of micro organisms like fungi, algae, bacteria. Bioaugmentation of the wastewater with highly effective strains provides a much more reliable process in which the process manager can use bacterial strains that target particular dye chemicals and metabolites to achieve complete mineralization. The most effective inoculants are able to degrade dyes over a broad concentration range, tolerate a range of environmental conditions of temperature, pH, and salinity. Laccases are usually known as benzenediol: oxygen oxidoreductase. They belong to the class of blue oxidases. Their molecular mass ranges from 60 to 85 kDa [2]. Laccases are involved in the biodegradation of lignins, which constitute the main non-carbohydrate component in wood and are among the most abundant groups of biopolymers in the biosphere. A great number of white-rot fungi have been reported to produce the lignin-degrading enzymes laccase, lignin peroxidases, and manganese peroxidases, or at least one of these enzymes [3]. Azoreductase catalyzes the reductive cleavage of azo linkages in benzidine based dyes and other compounds containing an azo bond to produce aromatic amines. Many bacterial strains possess many unspecific cytoplasmic enzymes which act as azoreductases. Azo reductases have been detected in liver cells possessing many unspecific cytoplasmic enzymes which act as azoreductases. Azo reductases have been detected in liver cells and several anaerobic bacteria. It is studied that when azodye is incubated in oxygen free buffer with NADH as a source of reduction equivalents, a slow decolorization is noted. Lignin peroxidase is a classical heme protein peroxidase containing a heme in the active site with molecular weight between 38 and 47 KDa [4]. Due to its high redox potential, LiP is able to directly oxidize non-phenolic lignin units. A characteristic of LiP, which is also shared by non-ligninolytic peroxidases, is its relative unspecificity for substrates such as phenolic compounds and dyes.

Methodology:
Data collection & curation:
A literature search was done using PubMed and the journals like ScienceDirect [9], Springerlink [10]. From that all the available information is retrieved till date. Search terms included azo dye degrading enzymes Laccase, Lignin peroxidase and Azoreductase. The data has also been collected from the database Brenda (an enzyme database). Nucleotide sequences are collected from NCBI [5] database and PDB ids are retrieved from the Protein Data Bank [6]. All the information has been curated manually.

Database structure:
The entries of our ‘BiodEnz’ database are generated from a text mining of hundreds of published articles. Our current database contains 234 entries of the micro organisms like bacteria, fungi and algae and their strains containing the azo dye degrading enzymes laccases, lignin peroxidases and azoreductases. The data base has separate sections for each enzyme. The information included the organism or strain in which the particular enzyme is available, concentration of dye the enzyme decolorizes, name of azodyes that are degraded or decolorized, percentage of decolorization, time taken by the organism to decolorize, molecular weight of the particular enzyme, optimum temperature and pH for that enzyme, enzyme specificity, electron donors, inhibitors, substrates, sequences, accession number of the organism is given from NCBI, and PDB ID from PDB.

Development & website structure:
The ‘BiodEnz’ database is developed using MySQL [7], a relational database management system that serves as the backend for storing data. APACHE 2.2 (Apache HTTP Server) is used as the web server and PHP5 (Hypertext preprocessor) [8] a widely used scripting language driven by Zend engine is used as the web interface. For the process of database creation in the PHP admin, the MySQL client version: 5.0.24a is used. The localhost of Server version: 5.0.24a, Protocol version: 10, Server: localhost via TCP/IP is used for the database creation. The local host for the process is php Myadmin of 2.9.0 as shown in the (Figure 1).

Utility:
BiodEnz database contains the detailed information about all the micro organisms (& their strains) such as bacteria, fungi and algae that contain the azo dye degrading enzymes, and also the properties of the laccases, azoreductases, and lignin peroxidases in those organisms like specificity of the enzymes, molecular weight, percentage of decolorization, enzyme activity, the optimum temperature and pH required for the enzyme to degrade the dyes. Our current database includes 89 entries of micro organisms for laccases, 57 entries of micro organisms for azoreductases and 57 entries of micro organisms for lignin peroxidases from various literatures and journals from 1990 to 2010. An activated sludge with culture of selected microbial strains is effective to degrade textile dyes in water. So our database provides a valuable information to perform bioaugmentation which is a biological cost effective method to convert the highly toxic and colourful dyes to

Figure 1: steps followed for creating database

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colourless, nontoxic substances, so that eco reserves are preserved. The sequences of the enzymes, Accession numbers, pdb ids are also included for further research about azodyes.

**Further development:**
The current database contains all the data about the three enzymes azoreductases, lignin peroxidase and laccases. In future we plan to add data about other azodye degrading enzymes and to update and improve the information from time to time from the sources.

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