

## Mini Review

# Biodegradation of Methyl Parathion and its Application in Biosensors

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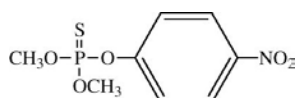
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**\*Corresponding author:** Jitendra Kumar, Nuclear Agriculture and Biotechnology Division, Mumbai, India**Received:** March 15, 2018; **Accepted:** May 09, 2018;**Published:** May 19, 2018**Abstract**

Methyl parathion is an Organo Phosphate (OP) insecticide which is being used in agriculture to protect the crops from insects. It causes many health problems in humans, related to acetyl cholinesterase inhibition. Methyl parathion is classified as Category Ia (extremely toxic) by the World Health Organization (WHO) and as Toxicity Category I (most toxic) insecticide by the United States Environmental Protection Agency (U.S. EPA). Organ Phosphorus Hydrolyses (OPH) (E.C.3.1.8.1) was discovered in soil micro-organisms and hydrolyzes methyl parathion into p-nitro phenol (PNP) and Di Methyl Thio Phosphate (DMTP). Hydrolyzed product PNP can be detected by electrochemical and optical methods. This review is a compilation of the work reported on OPH based enzymatic and microbial biosensor for detection of methyl parathion pesticide.

**Introduction**

Methyl parathion is an organophosphate insecticide, nematicide, and acaricide/miticide used to control boll weevils and many insect pests of agricultural crops [1-6]. Methyl parathion is produced by the reaction of O,O-dimethyl phosphorochloridithionate and the sodium salt of 4-nitrophenol in acetone solvent [7-8]. Bayer has developed this pesticide and has long been the 'parent' company with its well-known brand 'Folidol'. However, there have also been a number of other manufacturers globally. The main manufacturers of Methyl parathion are All India Medical Co (India), Bayer India, Bayer Mexico, Cheminova (Denmark), Rallis India, Sundat (Singapore) and Velpol Company (Mexico) [5,9]. The IUPAC chemical name of MP is O,O-dimethyl O-4-nitrophenyl phosphorothioate. Its chemical structure is shown below:



Methyl parathion kills pests by acting as a stomach poison and act as potent irreversible acetyl cholinesterase inhibitor. It was used to control a variety of insects and mites, including thrips, weevils, aphids and leafhoppers, in a very wide range of crops including cereals, fruit, nuts, vines, vegetables, ornamentals, cotton, and field crops [5,7,10]. Central Insecticide Board and Registration Committee (CIBRC) in India recommended it in two different forms either in 2% DP or 50% EC for controlling the insect pests from the cotton, paddy, wheat, pulses such as green gram and black gram and oilseeds such as ground nut and mustard crops.

Methyl parathion binds into the acyl pocket at the active site of acetyl cholinesterase enzyme. The binding of a phosphate group to the serine amino acid at the active site of acetyl cholinesterase changes the configuration of the enzyme molecule, stabilizing it and preventing it from functioning and inactivating permanently [11]. When inhaled by human being, the first adverse effects are a bloody or runny nose, coughing, chest discomfort and difficulty in breathing. Skin contact may cause localized sweating and involuntary muscle

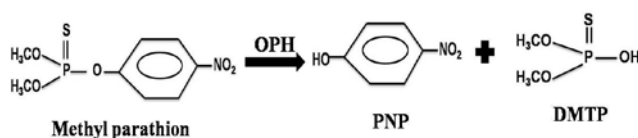


Figure 1: Structural presentation of methyl parathion hydrolysis with OPH.

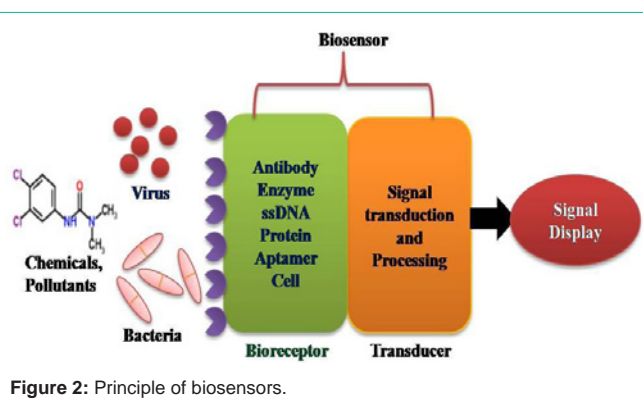


Figure 2: Principle of biosensors.

contractions. Following exposure by any route, other systemic effects may begin within a few minutes, or be delayed for up to 12 hours. These may include pallor, nausea, vomiting, diarrhea, abdominal cramps, headache, dizziness, eye pain, blurred vision, constriction or dilation of the pupils, tears, salivation, sweating and confusion [7,12]. In severe cases, poisoning will affect the central nervous system, producing in-coordination, slurred speech, loss of reflexes, weakness, fatigue, and eventual paralysis of the body extremities and respiratory muscles. Death may be caused by respiratory failure or cardiac arrest [7,12,13].

Methyl parathion was initially registered in 1954 in the United States for application as insecticide [14] but its uses was restricted in 1978 as a result of detrimental effects to humans [13,14]. Environmental Protection Agency (EPA) has now classified methyl parathion as a restricted-use pesticide and has given approval

**Table 1:** Biosensors for detection of methyl parathion.

S.No.	Microbial cells/ Enzyme	Type of detector	Detection range	Ref.
<b>Based on hydrolysis of methyl parathion by immobilized microbial cells</b>				
1	<i>Escherichia coli</i> cells OPH intracellularly	Potentiometric	Not reported	[44]
2	<i>Escherichia coli</i> cells expressing OPH on cell surface	Potentiometric	0.06-0.91 mM	[45]
3	<i>Moraxella</i> sp. express INPNC-OPH on the cell surface	Amperometric	Upto 175 $\mu$ M	[46]
4	<i>Pseudomonas putida</i> JS444, express OPH on the cell surface	Amperometric	Upto 2 $\mu$ M	[47]
5	<i>Pseudomonas putida</i> JS444 expressing OPH on cell surface	Dissolve oxygen electrode	0.2-50 $\mu$ M	[48]
6	<i>Escherichia coli</i> strain surface displayed mutant OPH (S5)	Amperometric	0.08-30 $\mu$ M	[49]
7	<i>E. coli</i> was having high periplasmic expression of OPH	Cyclic voltammetric	2-80 $\mu$ M	[50]
8	<i>Flavobacterium</i> sp.	Optical	4-80 $\mu$ M	[51]
9	<i>Sphingomonas</i> sp.	Optical	4-80 $\mu$ M	[52]
10	<i>Sphingomonas</i> sp.	Optical	4-80 $\mu$ M	[53]
11	<i>Sphingomonas</i> sp.	Optical	0.1-1ppm	[54]
<b>Based on hydrolysis of methyl parathion by immobilized OPH enzyme</b>				
12	OPH Enzyme	Electrochemical impedance spectra Linear Voltammetry	5.0-200 ng/mL and 200-1000 ng/mL	[55]
13	OPH Enzyme	Potentiometric	0.1-0.43 mM	[56]
14	OPH Enzyme	Amperometric	Up to 140 $\mu$ M	[57]
15	OPH Enzyme	Amperometric	1-10 $\mu$ M	[58]
16	OPH Enzyme	Amperometric	Up to 40 $\mu$ M	[59]
17	OPH Enzyme	Amperometric Cyclic voltammetry	Up to 2 $\mu$ M	[60]
18	OPH Enzyme	Chrono amperometric	4.6-46 $\mu$ M	[61]

for outdoor use only [10]. It was classified by the World Health Organization (WHO) as a Category Ia (extremely toxic) and by the United States EPA (U.S. EPA) as a Toxicity Category I (most toxic) insecticide [10]. Although banned in developed countries, it is still being used in India as a restricted insecticide. As per CIBRC in India, formulations of MP, 50% EC and 2% DP are banned for use on fruits and vegetables (S.O.680 (E) dated 17<sup>th</sup> July, 2001), and its use is restricted to only those crops where honeybees are not acting as pollinators. (S.O.658 (E) dated 04<sup>th</sup> Sep., 1992) [15].

### Biodegradation of methyl parathion

Organ Phosphorus Hydrolase (OPH) (E.C.3.1.8.1) was first discovered in soil micro-organisms *Pseudomonas diminuta* MG and *Flavobacterium* sp. and hydrolyzes methyl parathion into P-Nitro Phenol (PNP) and Di Methyl Thio Phosphate (DMTP) [16-21]. This hydrolytic reaction is the first steps of degradation of methyl parathion by soil microorganism which was extensively studied. OPH was first found in *Pseudomonas diminuta*, and then in *Flavobacterium* sp., both are soil microbes [22]. OPH also has been found in a variety of organisms such as squid, protozoa, mammals, yeast, fungi and soil bacteria [22,23]. OPH is also known as phosphotriesterase, parathion hydrolase, paraoxonase, DFPase, somanase, sarinase, phosphorothiolase and parathion aryl esterase [24-27]. The gene encoding OPH, *opd* (organophosphate degradation), has been expressed in various systems including *Escherichia coli*, *Drosophila melanogaster*, *Streptomyces lividans* and insect cells [23,24,28]. OPH has been studied extensively over the years [24,29-31] and several genetically engineered variants have been produced in an effort to improve its catalytic ability [31-33].

Below is the structural presentation of methyl parathion hydrolysis with OPH into P-Nitro Phenol (PNP) and Di Methyl Thio Phosphate (DMPT) (Figure 1).

PNP is an optically detectable product which can be detected by electrochemical and colorimetric methods. Thus, this hydrolytic step has been extensively exploited to develop the biosensor for detection of methyl parathion.

### Biosensors for detection of methyl parathion using biodegradation step

A biosensor is a self-contained integrated device which is capable of providing specific quantitative or semi-quantitative analytical information using a biological recognition element which is in direct contact with a transducer element. Biosensors consists of three basic components: (i) a biological component to interact with analyses which generate some physico-chemical signal, (ii) a transducer to convert the generated signal in to output signal (mostly electrical) and (iii) a signal processing system to process the output signal into the appropriate form that can be displayed on device (Figure 2) [5-6,34-36].

Biological materials play a very significant role in biosensor field as one of the main components which provide selectivity and specificity of the system for interest of analyze. The principle of detection is the specific binding of the analyze of interest to the complementary biological material immobilized on a suitable support matrix. The specific interaction results in a change in one or more physico-chemical properties like change in pH, electron transfer, mass changes, heat transfer, uptake or release of gases or specific

ions, which can be detected and produce an electronic signal, which is proportional to the concentration of a specific analyte or group of analyses. Now days, biosensor plays an important role as alternative detection method replacing the traditional analytical methods such as spectrophotometer, gas-liquid chromatography, thin-layer chromatography, high-performance liquid chromatography, capillary electrophoresis and mass spectrometry etc [37-41]. Biosensor facilitate onsite detection of large number of sample with no or low preparation, less time requirement and no requirement of expensive apparatus and trained personnel which are generally limitation in traditional analytical methods.

On the basis of bio component, if enzyme is used as bio component, it is known as enzymatic biosensor and if microbial cells are used, it is called microbial biosensor. Both bio components (enzyme/microbial cells) have certain limitations and advantages. Purified enzymes have very high specificity for their substrates or inhibitors, their application in biosensor construction may be limited by the tedious, time-consuming and costly enzyme purification steps and requirement of cofactor/coenzyme to generate the measurable product. Microbes provide an ideal alternative to these bottle-necks [35,42,43]. The enzymes and co-factors that co-exist in the microbes give the ability to consume and hence detect large number of analyses. Microbial cells can be easily manipulated and adapted to consume and degrade new substrates under certain cultivating condition. Additionally, the progress in molecular biology/recombinant DNA technology has opened endless possibilities of tailoring the microorganisms to improve the activity of an existing enzyme or express foreign enzyme/protein in host cell. All of the above makes microbial cells an excellent biosensing element for developing biosensor. The use of microbial cells has been demonstrated as an alternative biological catalyst without compromising on cost of purifying enzymes [35,42,43].

OPH enzyme and the microbial cells having OPH have been extensively utilized for developing biosensor to detect methyl parathion pesticide (Table 1). Enlist the publications where different microbial cells expressing OPH and purified OPH enzyme have been immobilized on a suitable matrix and further used as biosensor.

## Conclusion

This review focuses on biodegradation of methyl parathion using OPH enzyme which is present in soil microorganism. It lists many reports where microbe carrying OPH and purified OPH were used for hydrolysis of methyl parathion which further applied in biosensors for detection of methyl parathion pesticide.

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