

# Study of Predictors for Ventilatory Support in Patients of Organophosphorus Compound Poisoning.

Vipinsingh Thakur<sup>1</sup>, S.M Biradar<sup>2</sup>

<sup>1</sup>Junior Resident, Department of Medicine, BLDE University, Sri B M Patil Medical College Hospital and Research Centre, Vijayapur-586103, Karnataka, India.

<sup>2</sup>Associate Professor, Department of Medicine, B.L.D.E University, Sri B M Patil Medical College Hospital and Research Centre, Vijayapur-586103, Karnataka, India.

Received: November 2018

Accepted: November 2018

**Copyright:** © the author(s), publisher. Annals of International Medical and Dental Research (AIMDR) is an Official Publication of "Society for Health Care & Research Development". It is an open-access article distributed under the terms of the Creative Commons Attribution Non-Commercial License, which permits unrestricted non-commercial use, distribution, and reproduction in any medium, provided the original work is properly cited.

## ABSTRACT

**Background:** Organophosphorus compound poisoning is used most commonly for suicidal attempts which is seen frequently in farmers of younger age group of lower socio-economic status from rural areas. The aim of study is to know the clinical profile of organophosphate compound poisoning and to identify the factors which help in predicting the need for ventilatory support in this poisoning. **Methods:** A hospital based prospective cross sectional study was conducted with 100 patients to analyze the clinical profile of organophosphate compound poisoning and identify the factors which help in predicting the need for ventilatory support in organophosphorous compound poisoning. **Results:** Majority of the patients were in the age group of 21-30 years. The mean age of the patients was  $31.96 \pm 14.12$  years. The route of exposure was oral in all the patients. The most common organophosphorus compound consumed in our study was Malathion (28%), 46 (46%) patients had very poor GCS score ( $\leq 7$ ), 38 (38%) patients had low serum cholinesterase levels ( $\leq 2100$  IU/L), 9 (9%) patients had amylase levels  $>140$  U/L, 43 (43%) patients required ventilatory support while 57 (57%) patients did not require ventilator support. Among patients with fasciculations 64% were on ventilatory support. There was significant association of fasciculations and ventilatory support. 87 (87%) patients survived while 13 (13%) patients died in our study. All patients that died were on ventilatory support. **Conclusion:** Clinical and biochemical parameters such as greater the time lag from consumption of OP poison till getting specific treatment, Lower GCS scoring, Generalized Fasciculation's, Low cholinesterase levels were strong predictors for the need for Assisted Ventilation in OP poisoning.

**Keywords:** Organophosphorus, poisoning, ventilation.

## INTRODUCTION

India being an agriculture-based country, organophosphate (OP) pesticide remains the main agent for crop protection and pest control. It is therefore likely to have adverse effects on farmers who are accidentally over exposed while handling these pesticides. However, because of low cost and easy availability, it has also become an agent of choice for self poisoning.<sup>[1-3]</sup>

Organophosphates are one of the common causes of suicidality worldwide. Gunnell et al. have reported that there are 258,234 deaths each year from organophosphate poisoning, which accounts for about 30% of the suicidal cases globally.<sup>[4]</sup> OPs result in phosphorylation of serine hydroxyl residue on acetylcholine esterase enzyme, which results in the accumulation of acetylcholine.

This leads to cholinergic features, which can be classified into central and peripheral. Peripheral events include vomiting, diarrhea, miosis, muscle fasciculations, urinary incontinence, and bronchoconstriction. Central effects include respiratory depression and delirium.<sup>[5]</sup>

One of the devastating cholinergic features of organophosphate poisoning is respiratory failure. There are several explanations for respiratory failure; central, as well as peripheral mechanisms, underlie this phenomenon. However, studies have suggested that the major mechanisms regulating respiratory failure associated with OP ingestion are central in origin. The respiratory center known as the pre-Botzinger complex is situated in the ventrolateral medulla. It is composed of glutaminergic and muscarinic fibers. Excess acetylcholine can depress respiratory activity in these areas.<sup>[6-7]</sup> Injection of dichlorvos bilaterally into the pre-Botzinger complex in vagally intact rats produces a decrease in respiratory rate, a decrease in volume of inspired gas, and about 27% of the animals became apneic.<sup>[8-10]</sup>

### Name & Address of Corresponding Author

Dr. Vipin Singh Thakur,  
Room number 243, Boys Hostel  
Dr. B M Patil Road (Solapur Road),  
Vijayapur-586103, Karnataka, India.

Vagus is the major neural pathway that interconnects the brain and lung. Mechanoreceptors provide feedback via the vagus nerve. Vagal mechanisms blunt the hypoventilation associated with OPs in spontaneously breathing animals. Vagal mechanisms also cause an increase in pulmonary secretions due to pulmonary irritants, and surgical vagotomy has been shown to decrease pulmonary secretions.<sup>[11]</sup>

OP exposure to the lung causes increased acetylcholine at the pulmonary muscarinic receptors causing pulmonary abnormalities. OP agents increase the work of breathing through an increase in pulmonary static and dynamic compliance and by causing obstruction of airways.<sup>[12-14]</sup> OPs have the tendency to cause interstitial edema, which is responsible for the decrease in pulmonary compliance and ventilation-perfusion (V/Q) mismatch.<sup>[15,16]</sup>

Hence the present study was done at our tertiary care centre to assess the clinical profile of organophosphate compound poisoning and identify the factors which help in predicting the need for ventilatory support in organophosphorous compound poisoning.

### MATERIALS AND METHODS

A hospital based prospective cross sectional study was conducted with 100 patients admitted in BLDE Hospital to analyze the clinical profile of organophosphate compound poisoning and identify the factors which help in predicting the need for ventilatory support in organophosphorous compound poisoning.

#### Statistical Analysis

Quantitative data is presented with the help of Mean and Standard deviation. Comparison among the study groups is done with the help of unpaired t test as per results of normality test. Qualitative data is presented with the help of frequency and percentage table. Association among the study groups is assessed with the help of Fisher test, student 't' test and Chi-Square test. 'p' value less than 0.05 is taken as significant.

#### Pearson's chi-squared test

$$X^2 = \sum_{i=1}^n \frac{(O_i - E_i)^2}{E_i}$$

Where X<sup>2</sup> = Pearson's cumulative test statistic.

O<sub>i</sub> = an observed frequency;

E<sub>i</sub> = an expected frequency, asserted by the null hypothesis;

n = the number of cells in the table.

Appropriate statistical software, including but not restricted to MS Excel, SPSS ver. 20 is used for statistical analysis. Graphical representation has been done in MS Excel 2010.

## RESULTS

#### Distribution of patients according to Age

Majority of the patients (41%) were in the age group of 21-30 years followed by 11-20 years (23%), 31-40 years (18%), 51-60 years (9%), 41-50 years (6%) and >60 years (3%). The mean age of the patients was 31.96 ± 14.12 years.

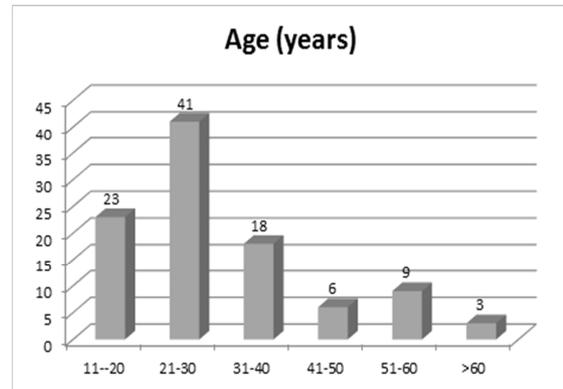


Figure 1: Distribution of patients according to Age

#### Distribution of patients according to Sex

In this study 38% patients were male and 62% were female. There was female preponderance in our study and the M:Fratio was 1:1.6. The route of exposure was oral in all the patients.

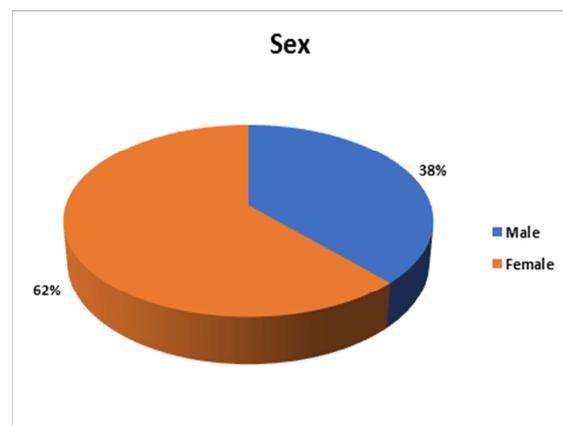


Figure 2: Distribution of patients according to Sex

#### Distribution of patients according to Type of Compound Consumed

The most common organophosphorus compound consumed in our study was Malathion (28%) followed by Dichlorovas (25%), Monocrotphos (24%), Dimethoate (6%) and Methylparathion (5%).

Table 1: Distribution of patients according to Type of Compound Consumed.

Type of Compound	N	%
Chlorpyriphos	2	2
Dichlorovas	25	25
Dimethoate	6	6
Malathion	28	28
Methylparathion	5	5

Monocrotophos	24	24
Paration	1	1
Phenylphrazole	1	1
Phorate	1	1
Phosphonic Acid	1	1
Prophenofus+Cypermethrin	2	2
Quinphos	1	1
Triazophos	2	2
Unknown Compound	1	1
Total	100	100

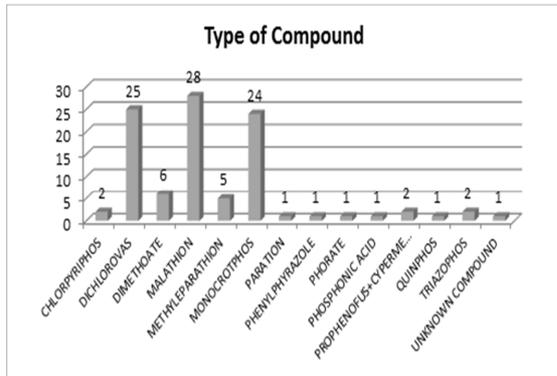


Figure 3: Distribution of patients according to Type of Compound Consumed

**Distribution of patients according to effect of OP poisoning**

The most common effect of OP poisoning was pinpoint pupils (74%) followed by respiratory failure (42%), fasciculations (11%) and intermediate syndrome (2%).

Table 2: Distribution of patients according to effect of OP poisoning

Effect of OP poisoning	N	%
Pinpoint pupils	74	74
Respiratory Failure	43	43
Fasciculations	11	11
Intermediate syndrome	2	2

**Distribution of patients according to Glasgow Coma Scale (GCS)**

46 (46%) patients had very poor GCS score ( $\leq 7$ ) while 53 (53%) patients had GCS score between 8 to 11. 1 (1%) patient had good GCS score (12-15).

Table 3: Distribution of patients according to Glasgow Coma Scale (GCS)

Glasgow Coma Scale	N	%
$\leq 7$	46	46
8-11	53	53
12-15	1	1
TOTAL	100	100

**Distribution of patients according to Serum Cholinesterase levels**

38 (38%) patients had low serum cholinesterase levels ( $\leq 2100$  IU/L) while 62 (62%) patients had serum cholinesterase levels  $> 2100$  IU/L.

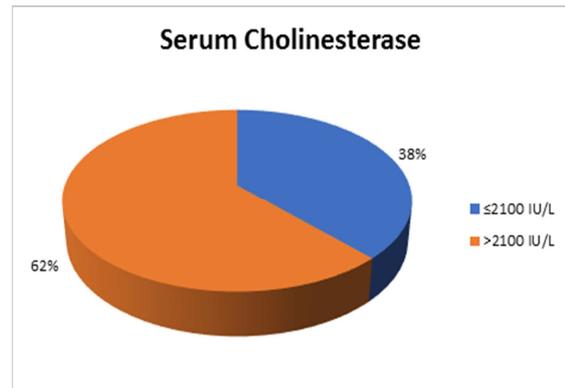


Figure 4: Distribution of patients according to Serum Cholinesterase levels

**Distribution of patients according to Amylase levels**

91 (91%) patients had amylase levels in the normal range (40-140 U/L) while 9 (9%) patients had amylase levels  $> 140$  U

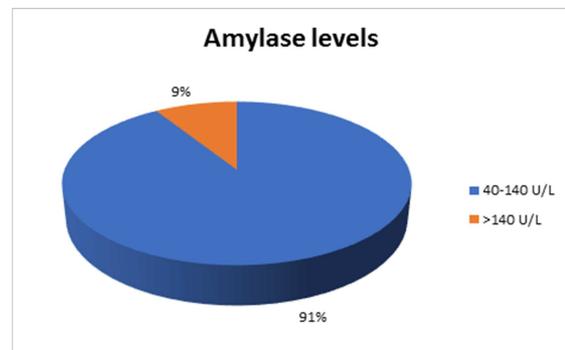


Figure 5: Distribution of patients according to Amylase levels

**Distribution of patients according to Time Interval between Consumption and Hospital Admission**

Majority of the patients (71%) were admitted in the hospital within 2-4 hours of ingesting organophosphorus poison while 18 (18%) and 9 (9%) patients were admitted within 4-8 hours and  $< 2$  hours. 2 (2%) patients were admitted after  $> 8$  hours of ingesting organophosphorus poison.

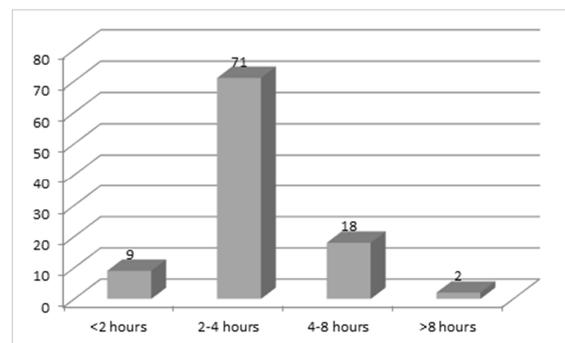
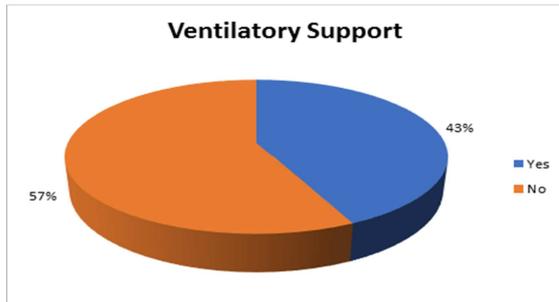


Figure 6: Distribution of patients according to Time Interval between Consumption and Hospital Admission

**Distribution of patients according to Requirement of Ventilatory Support**

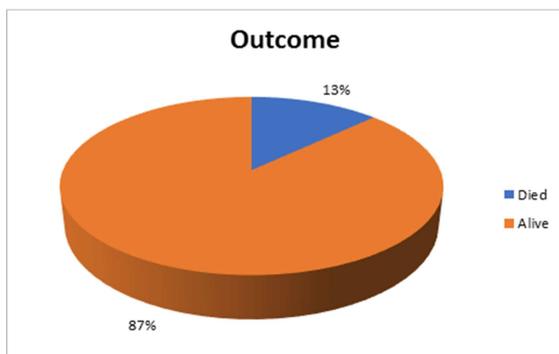
43 (43%) patients required ventilatory support while 57 (57%) patients did not require ventilator support.



**Figure 7: Distribution of patients according to Requirement of Ventilatory Support**

**Distribution of patients according to Outcome**

87 (87%) patients survived while 13 (13%) patients died in our study.



**Figure 8: Distribution of patients according to Outcome**

**Association of Pinpoint pupils and Ventilatory Support**

Among patients with pinpoint pupils 41(55.4%) were not on ventilatory support however there was no significant association of pinpoint pupils and ventilatory support as per Chi-square test (p>0.05).

**Association of Respiratory Failure and Ventilatory Support**

Among all patients 33 were on ventilatory support with respiratory failure, while 10 were not on ventilatory support with respiratory failure. There was significant association of respiratory failure and ventilatory support as per Chi-square test (p<0.05).

**Association of Fasciculations and Ventilatory Support**

Among all patients 11 were on ventilatory support with fasciculations, while there was no case without ventilatory support with fasciculations. There was significant association of fasciculations and ventilatory support as per Chi-square test (p<0.05).

**Association of Glasgow Coma Scale (GCS) and Ventilatory Support**

Majority of patients with very poor GCS score (67.4%) were on ventilatory support. There was significant association of Glasgow Coma Scale (GCS) and ventilatory support as per Chi-square test (p<0.05).

**Association of Amylase levels and Ventilatory Support**

Among patients with high amylase levels, 7 (16.3%) patients were on ventilatory support while 2 (3.5%) patients were not on ventilatory support. There was significant association of amylase levels and ventilatory support as per Chi-square test (p<0.05).

**Association of Serum Cholinesterase levels and Ventilatory Support**

Majority of the patients on ventilatory support had low serum cholinesterase levels (69.8%). There was significant association of serum cholinesterase levels and ventilatory support as per Chi-square test (p<0.05).

**Association of Time Interval between Consumption and Hospital Admission and Ventilatory Support**

There was significant association of time interval between consumption and hospital admission and ventilatory support as per Chi-square test (p<0.05).

**Association of Outcome and Ventilatory Support**

All patients that died were on ventilatory support. There was significant association of outcome and ventilatory support as per Chi-square test (p<0.05).

**Multivariate analysis for predictors of Ventilatory Support**

Logistic regression analysis was used to evaluate predictors of ventilatory support. The logistic regression analysis showed that respiratory failure, fasciculations, Glasgow Coma Scale, serum cholinesterase and time interval between consumption and hospital admission were independently associated with ventilatory support.

**Table 4: Association of Pinpoint pupils and Ventilatory Support**

Pinpoint pupils	Ventilatory Support				Total		p Value
	Yes		No				
	N	%	N	%	N	%	
NO	10	38.5	16	61.5	26	26.0	>0.05
YES	33	44.6	41	55.4	74	74.0	
Total	43	43.0	57	57.0	100	100.0	

**Table 5: Association of Respiratory Failure and Ventilatory Support**

Respiratory Failure	Ventilatory Support						p Value
	Yes		No		Total		
	N	%	N	%	N	%	
NO	10	17.5	47	82.5	57	57.0	<0.05
YES	33	76.7	10	23.3	43	43.0	
Total	43	43.0	57	57.0	100	100.0	

**Table 6: Association of Fasciculations and Ventilatory Support**

Fasciculations	Ventilatory Support						p Value
	Yes		No		Total		
	N	%	N	%	N	%	
NO	32	36.0	57	64.0	89	89.0	<0.05
YES	11	100.0	0	0.0	11	11.0	
Total	43	43.0	57	57.0	100	100.0	

**Table 7: Association of GCS and Ventilatory Support**

GCS	Ventilatory Support						p Value
	Yes		No		Total		
	N	%	N	%	N	%	
≤7	29	67.4	17	29.8	46	46.0	<0.05
8-11	14	32.6	39	68.4	53	53.0	
12-15	0	0.0	1	1.8	1	1.0	
Total	43	100.0	57	100.0	100	100.0	

**Table 8: Association of Serum Cholinesterase levels and Ventilatory Support**

Serum Cholinesterase levels	Ventilatory Support						p Value
	Yes		No		Total		
	N	%	N	%	N	%	
≤2100 IU/L	30	69.8	8	14.0	38	38.0	<0.05
>2100 IU/L	13	30.2	49	86.0	62	62.0	
Total	43	100.0	57	100.0	100	100.0	

**Table 9: Association of Amylase levels and Ventilatory Support**

Amylase levels	Ventilatory Support						p Value
	Yes		No		Total		
	N	%	N	%	N	%	
40-140 U/L	36	83.7	55	96.5	91	91.0	<0.05
>140 U/L	7	16.3	2	3.5	9	9.0	
Total	43	100.0	57	100.0	100	100.0	

**Table 10: Association of Time Interval between Consumption and Hospital Admission and Ventilatory Support**

Time Interval	Ventilatory Support						p Value
	Yes		No		Total		
	N	%	N	%	N	%	
<2 hours	2	22.2	7	77.8	9	9.0	<0.05
2-4 hours	30	42.3	41	57.7	71	71.0	
4-8 hours	9	50.0	9	50.0	18	18.0	
>8 hours	2	100.0	0	0.0	2	2.0	
Total	43	43.0	57	57.0	100	100.0	

**Table 11: Association of Outcome and Ventilatory Support**

Outcome	Ventilatory Support						p Value
	Yes		No		Total		
	N	%	N	%	N	%	
Died	13	30.2	0	0.0	13	13.0	<0.05
Alive	30	69.8	57	100.0	87	87.0	
Total	43	100.0	57	100.0	100	100.0	

**Table 12: Multivariate analysis for predictors of Ventilatory Support**

Parameters	OR	95% CI	p Value
Age	0.75	0.54–1.03	p>0.05
Sex	1.23	1.00–1.51	p>0.05
Pinpoint pupils	1.17	0.45–2.85	p>0.05
Respiratory Failure	1.70	1.15–2.50	p<0.05
Fasciculations	1.50	1.28–1.75	p<0.05
Glasgow Coma Scale	1.12	1.01–1.25	p<0.05
Serum Cholinesterase	2.70	2.33–3.13	p<0.05
Amylase	0.81	0.61–1.09	p>0.05
Time Interval between Consumption and Hospital Admission	1.42	1.26–1.60	p<0.05

## DISCUSSION

OP compounds were synthesized by von Hoffman. OP pesticide poisoning is common in developing worlds. The highest incidence is seen in India. Suicidal and non-suicidal organophosphate poisoning is a major problem in rural areas of India, with rapidly increasing incidence rate.<sup>[17]</sup>

Chowdhary AN et al,<sup>[2]</sup> prospective study reported incidence of suicidal poisoning is 98.6%, probably because it is cheap, easily available and used as a major pesticide in agricultural farming throughout India.

The leading cause of death in OP poisoning is respiratory failure 18-20 and various grading systems proposed suggest that most cases can be managed in the ICU.

Tsao TC et al,<sup>[21]</sup> prospective study evaluating various parameters that can predict outcome of patients in OP poisoning found one hundred two (76.7%) were males and 31 (23.3%) female. Most of the cases were young people 80% (< 40 years) predominantly males. There was wide variation in age ranging from a minimum of 13-68 years with mean age of 31.5 years.

Fryer AD et al<sup>[12]</sup> descriptive study assessing clinical and biochemical parameters in organophosphate poisoning, which help to predict the need for ventilator support observed maximum number of cases was 21-30 years in the age group of years, youngest patient in present study was 14 years and the oldest patient in this study was 68 years. Out of 50 cases 33 were Males (66%) and 17 were females (34%).

Rodger ML et al,<sup>[18]</sup> descriptive study assessing clinical and biochemical parameters in organophosphate poisoning, which help to predict the need for ventilator support reported Methyl parathion was the commonest poisoning encountered; out of 25 cases 15 cases required ventilator support. Among 6 cases of Dimethoate poisoning, 4 cases (66.7) required ventilator support. Shetti AN et al,<sup>[22]</sup> prospective, observational, descriptive, intention to-treat study reported around 68% of patient presented with bradycardia, 28% of patients with miosis, 28% of patients with altered sensorium, 21% of patient with tachypnea, 15% Of patient fasciculation and 10% of patients with seizures.

Chethan RAN et al,<sup>[23]</sup> prospective, observational, descriptive, intention to-treat study observed mean serum pseudocholinesterase level in mild poisoning was 5680.653 U/L, moderate poisoning was 4707 U/L, severe poisoning was 175.133 U/L.

Rajeev H et al,<sup>[24]</sup> prospective study evaluating various parameters that can predict outcome of patients in OP poisoning reported fifty three patients required ventilatory support, out of which only 11 patients survived. Patients were on ventilator support

for minimum 1 day to maximum 22 days with a mean  $6.85 \pm 4.32$  days. Mortality was higher in patients who required ventilator support >7 days [P < 0.05 statistically significant].

Shetti AN et al,<sup>[22]</sup> retrospective study correlating the serum acetylcholinesterase levels with morbidity, ventilation need, ICU stay and the final outcome of the ailment reported 10 (40%) male patients died and 2 (28.5%) female patients died.

Rajeev H et al,<sup>[24]</sup> descriptive study assessing clinical and biochemical parameters in organophosphate poisoning, which help to predict the need for ventilator support reported mortality of 16% (4 out of 24 Ventilated patients)

Chethan RAN et al,<sup>[23]</sup> prospective, observational, descriptive, intention to-treat study reported most of the patients (73%) recovered completely and discharged without ventilator support, their serum pseudocholinesterase level was above 5000 U/L, POPScore was below 5 and Mean duration of hospital was 5 days. The longest hospital stay was of a male patient who had consumed parathion and was in hospital for 55 days (POP scale 9). He presented with respiratory failure within 6 hours of consumption and was intubated for 13 days and tracheostomy was done and had complete recovery and discharged after 55 days. 27% of patients required ventilator support. Death has occurred in 10% of patients. POP Score in death patients was above 8, mean pseudo-cholinesterase level was 571 U/L.

Patil SL et al,<sup>[25]</sup> descriptive study assessing clinical and biochemical parameters in organophosphate poisoning, which help to predict the need for ventilator support reported survival rate after ventilator support was better with methyl parathion poisoning, 25 out of 50 (50%) while Fenithrothion (Tik -20) poisoning was second commonest; 12 out of 50 (24%) cases. The survival rate after ventilator support was better with methyl parathion poisoning 22 out of 25 patients survived while in Dimethoate poisoning where 4 out of 6 patients survived

## CONCLUSION

OP insecticide poisoning is a life threatening condition that needs rapid diagnosis and treatment. Since most of the patients present with respiratory failure, early initiation of mechanical ventilation plays a vital role in the treatment of such cases. Emphasis must also be given to good supportive care and monitoring for the prevention and management of acute and delayed complications that occur during the course of stay in ICU.

Clinical and biochemical parameters such as greater the time lag from consumption of OP poison till getting specific treatment, Lower GCS scoring, Generalized Fasciculation's, Low Pseudo

cholinesterase levels were strong predictors for the need for Assisted Ventilation in OP poisoning. Grading of the severity of the OP compound poisoning taking the above parameters into consideration can help to identify high risk patients who may go in for Respiratory failure and require ICU admission and Ventilator support.

## REFERENCES

1. South-East Asia: World Health Organisation; 2009. WHO. Health implications from monocrotophos use: A review of the evidence in India; pp. 1–60.
2. Chowdhary AN, Banerjee S, Brahma A et al. Pesticide poisoning in nonfatal, deliberate selfharm: A public health issue. *Indian J Psychiatr.* 2007;49:117-20.
3. Srivastava A, Peshin SS, KaleekalTet al. An epidemiological study of poisoning cases reported to the National Poisons Information Centre, All India Institute of Medical Sciences, New Delhi. *Hum ExpToxicol.* 2005;24:279-85.
4. The global distribution of fatal pesticide self-poisoning: systematic review. Gunnell D, Eddleston M, Phillips MR et al. <https://bmcpublihealth.biomedcentral.com/articles/10.1186/1471-2458-7-357>. *BMC Public Health.* 2007;7:357–371.
5. Central respiratory failure during acute organophosphate poisoning. Carey JL, Dunn C, Gaspari RJ. <http://www.sciencedirect.com/science/article/pii/S1569904813002668?via%3Dihub>. *RespirPhysiolNeurobiol.* 2013;189:403–410.
6. Dichlorvos induced central apnea: effects of selective brainstem exposure in rat. Gaspari RJ, Paydarfar D. <https://www.ncbi.nlm.nih.gov/pubmed/21241738>. *Neurotoxicology.* 2011;32:206–214.
7. Organophosphate neurotoxicity: chronic effects of sarin on the electroencephalogram of monkey and man. Burchfiel JL, Duffy FH. <http://psycnet.apa.org/record/1983-32999-001>. *NeurobehavToxicolTeratol.* 1982;4:767–778.
8. Effect of cholinesterase inhibitor when injected into the medulla of rabbit. Stewart WC, Anderson EA. <http://jpet.aspetjournals.org/content/162/2/309.short>. *J PharmacolExpTher.* 1968;162:309–318.
9. Pontine cholinergic respiratory depression in neonatal and young rats. Fung ML, St John WM. <http://www.sciencedirect.com/science/article/pii/S0024320598002033>. *Life Sci.* 1998;62:2249–2256.
10. Pedunclopontine stimulation alters respiration and increases ACh release in the pontine reticular formation. Lydic R, Baghdoyan HA. <http://ajpregu.physiology.org/content/264/3/R544>. *Am J Physiol.* 1993;264:544–554.
11. Respiratory failure induced by acute organophosphate poisoning in rats: effects of vagotomy. Gaspari RJ, Paydarfar D. *Neurotoxicology.* 2009;30:298–304.
12. Fryer AD, LeinPJ, HowardAS et al. Mechanisms of organophosphate insecticide induced airway hyperreactivity. <http://ajplung.physiology.org/content/286/5/L963.article-info>. *Am J Physiol Lung.* 2004;286:963–969.
13. Abbrecht PH, Kyle RR, Bryant HJ. Pulmonary mechanical responses to cholinesterase inhibitor. <http://www.sciencedirect.com/science/article/pii/027205908902972>. *FundamApplToxicol.* 1989;13:593–604.
14. Thompson JW, Stocks RM. *Arch Otolaryngol Head Neck Surg.* Vol. 123. Surg: 1997. Brief bilateral vocal cord paralysis after insecticide poisoning: a new variant of toxicity syndrome; pp. 93–96.
15. Segura P, Chanez J, Montano LMet al. Identification of mechanisms involved in acute airway toxicity induced by parathion. *NaunynSchmiedebergs Arch Pharmacol.* 1999;360:699–710.
16. Delaunois A, Gustin P, Ansay M. Altered capillary filtration coefficient in parathion- and paraoxon-induced edema in isolated and perfused rabbit lungs. *ToxicolApplPharmacol.* 1992;116:161–169.
17. Yanagisawa N, Morita H, Nakajima T. Sarin experiences in Japan: Acute toxicity and long-term effects. *J Neurol Sci.* 2006;249:76–85.
18. Rodgers ML. OP poisoning. *Am J Emerg Med.* 2006;22:335–44.
19. Gupta SK, Peshin SS, Srivastava A et al. An epidemiological pattern of poisoning in India. *Pharmacoepidemiol Drug Saf.* 2002;11:73–4.
20. Bardin PG, Van Eaden SF, Moolman JA. Organophosphate Poisoning and Carbamate poisoning *Arch Intern Med.* 1994, 154, 1433-1441.
21. Tsao TC, Juang V, Lan R et al. Respiratory failure of acute OP Poisoning", *Chest* 1990, 98, 6.31-636.
22. Shetti AN, Bhumika R, Singla B et al. Correlation of serum acetylcholinesterase with the ventilation need, ICU stay and outcome in organophosphorus poisoning – a retrospective study. *Anaesth Pain & Intensive Care.* 2017;21(2):199-203
23. Chethan Kumar RAN, Sahna E. Correlation of Serum Pseudocholinesterase Level and Peradeniya Organophosphorus Poisoning Scale with the Severity and Inhospital Outcome of Acute Organophosphorus Poisoning. *International Journal of Contemporary Medical Research.* 2017, Volume 4, Issue 8, ICV: 77.83.
24. Rajeev H, Arvind MN. "Study of clinical and biochemical parameters in predicting the need for ventilator support in organophosphorus compound poisoning". *Journal of Evolution of Medical and Dental Sciences* 2013; Vol2, Issue 49, December 09; Page: 9555-9570.
25. Patil SL, Vasepalli P. Prognostic value of clinical and lab parameters in assessing the severity of organophosphorous compound poisoning *Indian Journal of Basic and Applied Medical Research* 2014;4:77- 91.

**How to cite this article:** Thakur V, Biradar SM. Study of Predictors for Ventilatory Support in Patients of Organophosphorus Compound Poisoning. *Ann. Int. Med. Den. Res.* 2019; 5(1):ME23-ME29.

**Source of Support:** Nil, **Conflict of Interest:** None declared