Aluminium Phophide Poisoning: Can Mode of Ingestion Predict Mortality?

A Study in North Bihar

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Abstract

Aluminium phosphide poisoning has emerged as a major problem in rural India. No specific antidote is available. This study was conducted to evaluate relation between mode of ingestion and outcome in terms of mortality.

Materials And Methods: An observational study was done on patients admitted with ALP (Aluminium phosphide) poisoning in DMCH, Bihar. Diagnosis was done clinically and biochemically. Clinical variables and outcomes were recorded and statistically analysed using appropriate test.

Results: Fatality was not related to dosage of ingestion. Patients who dissolved the tablet in water before ingestion had a highly significant better outcome in terms of mortality.

Introduction

Agrochemical poisoning is a major public health problem in developing countries particularly in setting of low education and poor regulatory frameworks. Aluminium phosphide is a solid fumigant pesticide. It is marketed in India as tablets of celphos and quickphos. Aluminium phosphide has currently aroused interest with increasing number of cases in the past three decades due to increased use in agricultural and non-agricultural purpose. In India, this poisoning was not known before 1980. The first case in India was reported in 1981 from M.G.M. Medical College, Indore. The incidence of the poisoning has been increasing steadily and is now the commonest mode of suicide in the agricultural community in northern India. Although the percentage of poisoning with AIP is low, but the mortality rate of this kind of intoxication is very high. Overall mortality in cases of aluminium phosphide poisoning varies between 37–100%. The wide variability in survival rate and inability to find out effective antidote aroused interest in search of predictors of outcome.

Background of Study

Aluminium phosphide is an efficacious and easy to use and freely available rodenticide in India in form of chalky white or brown 3 gm. tablets containing 56% of ALP and 44% of ammonium...
carbonate. The tablets are taken out of sealed container and placed on stored grains and storage container is closed for few days to combat moles and vermines in granaries. ALP has relatively high vapour pressure that allows it to penetrate porous material effectively. On coming into contact with water or moisture or OH radical of air or hydrochloric acid in stomach, 3 gm. tablet of ALP liberates 1 gm. of phosphine or phosphorus hydrogen, i.e., $\text{ALP} + 2\text{H}_2\text{O} = \text{ALP} + 3\text{H}_2\text{O} = \text{AL(OH)}_3 + \text{PH}_3, \text{ALP} + 3\text{HCL} = \text{ALCL}_3 + \text{PH}_3$

Phosphine is a colourless gas with fishy or garlic odour. Irrespective of routes of exposure, the inhalational, ingestional or ocular, the toxic effects of PH$_3$, are same. Some of ALP is directly absorbed from stomach to reach liver to liberate PH, slowly to prolong the toxic effects of poisoning. It is rapidly absorbed from stomach or lungs by simple diffusion, oxidised slowly and is excreted in urine as hypophosphite and also excreted unchanged through lungs significantly.

PH$_3$ inhibits the electron transport resulting from preferential inhibition of cytochrome oxidase leading to respiratory chain inhibition which leads to cellular hypoxia and small vessel injury which is further potentiated by cardiotoxicity due to anoxic myocardial damage and shock. Direct toxic effect of ALP leads to arrhythmias. Hypotension and shock ensue within 3-6 hours of ingestion of ALP. In survivors, the cardiotoxicity and hypoxia disappear within 5-7 days due to excretion of PH$_3$, and restoration of normal cellular metabolism. The toxic chemical myocarditis leads to varied fatal ECG changes, 6 to 24 hours after ingestion in non-survivors, in the form of VE beats, conduction disturbance, LBBB, ventricular fibrillation, aberrant conduction and idioventricular rhythm terminally leading to asystole. The non-fatal ECG changes appear within 12 to 24 hours in survivors and disappear within 56 to 80 hours. Death in first 24 hours appears to be cardiogenic as evidenced by shock and ECG abnormality. Since the survivors show complete normal ECG recovery, it denotes that the effect of poisoning is due to some reversible factor leading to disturbance in the permeability of sodium, potassium, calcium and magnesium ions leading to change is trans membrane action potential due to focal myocardial involvement and subsequent myocardial necrosis. The peripheral circulatory failure (PCF) due to wide spread small vessel injury leads to peripheral vasodilatation leading to shock. Direct toxic effects of PH$_3$, on adrenal cortex accompanied by decreased cortisol levels, leads to shock and high mortality. Injury to alveolar capillary membrane by PH$_3$, while being inhaled, leads to ARDS (Adult Respiratory Distress Syndrome) which may also occur rarely in ingestion. Wide spread capillary damage leads to bleeding diathesis, disseminated intravascular coagulation (DIC) and acute tubular necrosis (ATN). Shock and DIC lead to terminal renal failure. The clinical features are more or less the same irrespective of the mode of toxicity, except the initial symptoms pertaining to the route of entry. Ingestional Toxicity manifests as: (i) mild: Nausea, vomiting, headache, abdominal pain and discomfort. These patients usually recover. (ii) Moderate and Severe Systemic Manifestations: gastrointestinal System: Nausea, vomiting, diarrhoea, pain epigastrium, retrosternal pain and epigastric burning sensation, hepato-biliary System: Acute hepatic failure, jaundice, hepatitis and soft tender hepatomegaly. cardiovascular System (60-100%) : Increased JVP, feeble heart sounds, S3 gallop, hypotension, shock, arrhythmias, myocarditis and pericarditis, respiratory System (within 2 to 3 hours) : Cough, dyspnoea, cyanosis, bilateral basal rales and rhonchi, respiratory failure and ARDS, (24) renal System: Acute (oliguric or nonoliguric) renal failure, central Nervous System: Headache, dizziness, diplopia, paraesthesias, ataxia, altered sensorium, restlessness, intention tremors, convulsion, hypoxic encephalopathy, coma and delayed haemorrhagic stroke muscular System: Muscle pain, severe muscle weakness, myopathy with muscle wasting and proximal muscle weakness, haemopoietic System: Bleeding diathesis, DIC and jaundice endoerinal system:
hypoglycemia and hyperglycemia. Bad Prognostic Signs include intractable shock, anemia, chest infection, metabolic acidosis, severe hypoxia, electrolyte disturbances, arrhythmias, oliguria, aspiration pneumonia, haemolysis, coma and DIC.

Objectives
To study the effect of dissolving of aluminium phosphide tablet in water before ingestion in prediction of survival.

Methods
Information of all patient admitted with AIP poisoning to the DMCH from January 2015 to January 2017 were collected and reviewed. Only patients who were hospitalized were included in the study. Information regarding age, cause of intoxication, amount of AIP consumed, route of exposure, therapeutic intervention and laboratory tests including arterial blood gas (ABG), electrocardiogram (EKG), and outcome were obtained. The factors of positive history of ingestion, symptoms compatible with AIP ingestion and chemical test for phosphine positive in gastric aspirate in combination used for diagnosis. All patients were managed with supportive care. Data were analysed with appropriate statistical analysis.

Results: 30 patients with aluminium phosphide poisoning were studied in this study. 12 (40%) patients survived with a mortality rate of 60%. (figure 1)

![Figure 1](image)

The mean age in the study group was 20 +/- 4 years. There was no significant difference in age among survivors and non survivors. (table 1)

<table>
<thead>
<tr>
<th>Mean age (in years)</th>
<th>Survived (mean +/- SD)</th>
<th>Non survived (mean +/- SD)</th>
<th>P value</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>23.78 +/-2.6</td>
<td>24.64 +/-3.1</td>
<td>&gt;0.05</td>
</tr>
</tbody>
</table>

The mean intake of tablets were 4 +/- 0.05. There was no significant difference of ingested dose among survivors and non survivors. (table 2)

<table>
<thead>
<tr>
<th>Mean intake of tablets (in numbers)</th>
<th>Survivor(mean +/- SD)</th>
<th>Non survivor(mean +/- SD)</th>
<th>P value</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>3.41 +/-1.72</td>
<td>3.55 +/-1.24</td>
<td>0.39</td>
</tr>
</tbody>
</table>

In our study, 10 patients took dissolved tablets and 20 patients took tablets without dissolving. The survival rate was (90%) in patients who had dissolved tablets while it was (15%) in those who consumed undissolved tablets. (table 3)

<table>
<thead>
<tr>
<th></th>
<th>Survived(n=12)</th>
<th>Not survived(n=18)</th>
<th>P value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Dissolved tablet(n=10)</td>
<td>9</td>
<td>1</td>
<td>0.00013</td>
</tr>
<tr>
<td>Undissolved tablet(n=20)</td>
<td>3</td>
<td>17</td>
<td>(highly significant)</td>
</tr>
</tbody>
</table>

Table 1
Table 2
Table 3
The average mean arterial pressure on arrival was 53.73 ± 17.96 mm Hg. It was significantly high among survivors. (table 4)

**Table 4**

<table>
<thead>
<tr>
<th>Mean arterial pressure (mm of Hg)</th>
<th>survivor</th>
<th>Non survivor</th>
<th>P value</th>
</tr>
</thead>
<tbody>
<tr>
<td>73.91±/2.77</td>
<td>9</td>
<td>40.27+/8.15</td>
<td>&lt;0.0001 (highly significant)</td>
</tr>
</tbody>
</table>

Clinical manifestations at time of admission involved different systemic manifestations. They were nausea and vomiting (83.33%), dyspnoea and palpitations (40%), cyanosis (16.66%) hypotension (80%) and shock (60%). Cardiac arrhythmias were present in 46.66% cases.

**Discussion**

The aluminium phosphide is a highly lethal poison with no definite antidote till now. The LD50 dose of ALP is 10 mg/kg of body weight. The specified fatal dose is 0.15-0.5 gm. However, most of the patients present with ingestion of three or more tablets which invariably results in death. For a 70 kg man 0.5 gm. ALP is lethal. The exact mechanism of action of aluminum phosphide poisoning is still unknown. A three-gram tablet can release almost one gram of phosphine gas in contact with water. After ingestion, phosphine will be released due to contact between AlP and water/acid in the gastrointestinal (GI) tract. Some phosphide may be absorbed by the GI tract without hydrolysis and convert into phosphine. The mortality varies in different studies from 37-100%. In some studies no significant correlation was found between the dose of aluminium phosphide ingested and outcome. In our study we also found no significant difference between quantity of tablets consumed. Interestingly there was different rate of survival depending upon dissolving the tablet in water before ingestion. This is probably because a great part of the phosphine gas has escaped before its consumption.

**Conclusion**

Aluminium phosphide poisoning is a great health hazard in developing country like India due to easy and wide availability. The exact mechanism of action of aluminum phosphide and an effective antidote of this poison is still unknown. But we assume that contact with water prior ingestion may predict a better outcome. Further studies in this field is needed.
References


