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Role of Total Antioxidant Capacity and Troponin I in Predicting Outcome of Acute Zinc Phosphide Poisoning: A Prospective Study

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Omv	cisity, Zagaz	ig, Lgypt.					
Corr	esponding	author*:	ABSTRACT:				
Aya	Ahmed	Elsayed	Background: Zinc phosphide (Zn3P2) is cheap, easily available, highly potent				
Moha	imed	•	rodenticide. Zinc phosphide toxicity is considered one of the most common				
Emai	1:		poisonings, particularly in developing countries, with multiorgan toxicity that				
avaah	med203065@	ngmail com	deeply affects the poisoned patient outcome.				
ayaar		ginan.com	Aim: The study aimed to assess the effectiveness of total antioxidant capacity				
Submit	Date 10-05-2024	5	(TAC) and Troponin I for early prediction of acute (Zn3P2) toxicity outcome.				
Revise	Date23-05-2025	5	Methods: This prospective cohort study was conducted on 53 cases of acute				
Accept	Date 26-05-2025		(Zn3P2) toxicity presented at Zagazig University Hospitals (emergency room				
			and intensive care units) over the period from July 2023 to January 2024.				
			Patients were evaluated by a thorough history, clinical assessment, routine				
			laboratory investigations, focusing attention to the role of TAC, Troponin I in				
			predicting severity.				
			Results: This study included patients with an age range from 17 to 57 years.				
			Females were more than males representing 67.9% of patients. Regarding the				
			clinical picture, 73.6% of patients were symptomatic, abdominal pain was the				
			most, representing 61.5% of cases followed by vomiting 59% respectively. The				
			Receiver operator characteristic curve (ROC) curve for Troponin I and TAC				
			showed that Troponin I had better validity as a severity predictor of Zn3P2				
			poisoning followed by TAC according to their AUC.				
			Conclusion: TAC and Troponin I are simple, easy and reliable markers that				
			save time. They could be good predictors of poor outcomes in zinc phosphide				
			poisoning.				
			Recommendations: It is recommended to use TAC and Troponin I as				
			predictors for severity and need for ICU admission in acute (Zn3P2) poisoning.				
			Keywords: Zinc phosphide; Acute Toxicity; Total Antioxidant Capacity;				
			Troponin I; Outcome.				
	INT	RODUCT	ION suicidal, is regarded as a health risk in Egypt				
inc phosphide is one of			the most common, because it is an inexpensive, readily access				

Inc phosphide is one of the most common, highly potent rodenticides, which has been used since 1940 in agricultural, urban and industrial environments [1]. Because of its potent garlic-like scent, which attracts mice but repels other animals, it is used to control rodents like rats, ground squirrels, prairie dogs, nutria, and muskrats [2]. These days, zinc phosphide poisoning, whether accidental or suicidal, is regarded as a health risk in Egypt because it is an inexpensive, readily accessible toxin with a high fatality rate of 37–100% and no laws regulating its purchase [3]. Zinc phosphide reacts with the acid of the stomach or its water content leading to the release of phosphine gas (PH3) [4]. Because it inhibits cytochrome C oxidase, phosphine gas disrupts oxidative phosphorylation and is quickly absorbed by the stomach and lungs, resulting in

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acute poisoning [1]. This causes lipid peroxidation, protein denaturation, and the production of extremely reactive oxygen molecules, which harm cells and cause damage to multiple organs [5].Congestive heart failure, severe metabolic acidosis, cardiac arrhythmia, liver necrosis, nausea, vomiting, and abdominal discomfort are the typical clinical signs of zinc phosphide overdose [1]. Cardiovascular collapse is the primary cause of mortality.

Treatment for zinc phosphide poisoning is supportive and symptomatic since there is no specific antidote for the condition [6]. Since cardiac troponins are very sensitive and specific biomarkers that can forecast the prognosis of cardiotoxicity, they are the most suitable biomarkers for the diagnosis of heart damage [7]. Furthermore, the entire antioxidant status in biological samples is referred to as total antioxidant capacity (TAC). It is among the most significant bio-analytical indicators of the body's oxidative stress [8]. Initial TAC values in patients with acute metal phosphide poisoning may help predict the onset of cardiotoxicity and oxidative damage, which could lower mortality and improve patient outcomes [9]. The purpose of the study was to evaluate Troponin I and total antioxidant capacity (TAC) as predicting tools for acute zinc phosphide toxicity outcome.

METHODS

This is a prospective cohort study conducted on 53 patients with acute zinc phosphide toxicity presented at Zagazig University Hospitals (emergency room and intensive care units) during seven months' period from July 2023 to January 2024.Inclusion criteria include clear history of intake of zinc phosphide, clinical picture of acute zinc phosphide poisoning (severe vomiting, hypotension, abdominal pain, cardio-pulmonary disturbance), patient age above 16 y. Exclusion criteria include Patients with unclear history, comorbidities (such as cardiovascular disease, renal or hepatic failure), w All enrolled patients were subjected to the following:

A-History taking from which the poisoning history (including the mode of poisoning, the amount, and the interval between consumption and hospital presentation) and sociodemographic information (particularly age and sex) were gathered.

B-Clinical assessment of the Glasgow coma scale (GCS) to determine the conscious state and vital indicators (blood pressure, heart rate, respiration rate, and temperature) [10] was performed.

C-Investigatory workup:

1) Biochemical studies:

Blood samples were withdrawn as follows: About 2 cm3 of arterial blood were withdrawn in a heparinized syringe to perform arterial blood gases (ABG) with sodium (Na+) and potassium (K+) electrolytes by an automated analyzer (Cabs B 221) which is present in Zagazig University Hospitals according to the method described by [11]. Approximately 10 cm3 of blood was collected on admission from the intake, 3 cm3 were separated into an EDTA tube to perform a complete blood count (CBC) test, and 7 cm3 were moved to a plain tube. The tube was left upright to allow the blood to clot, and after 10 minutes of centrifugation at 5000 rpm, samples of the serum supernatant were taken. A portion was used for routine investigations, and the remaining portion was preserved at -80°C to be used later for specific tests. All patients were evaluated by the following tests:

A-Nonspecific (routine investigations done for all zinc phosphide poisoned cases in Zagazig University Hospitals): Complete blood picture (CBC) by using Siemens Advia 120, hematology analyzer [12], liver function tests (LFT), namely ALT, AST, Total bilirubin, Direct bilirubin were assessed according to the method described by [13], [14], [15], [16] respectively, and kidney function tests (KFT) by spectrophotometry, namely serum creatinine as described [17] and blood urea nitrogen (BUN) by the colorimetric method [18].

B-Specific investigations:

- Plasma total antioxidant capacity (TAC): by the described colorimetric method according to [19]by spectrophotometry at wavelength 500-510 nm.
- Serum cardiac Troponin I level by Colloidal Gold Assay using the Fast Test kit (Getein Biotech. Inc.; USA) with Getein 1100 immunofluorescence quantitative analyzer [20].
- Serum creatine kinase-MB fraction level by a sandwich electrochemiluminescence immunoassay method, using Full automated cs200florometric analyzer [21].

2) Standard twelve-lead Electrocardiography (ECG) that was routinely recorded for all acute zinc phosphide poisoned cases on arrival at the university hospital ER by using CardiMax FCP-7101 ECG machine (Fukuda Denshi Co., Ltd., Tokyo, Japan) [22].

Case Management:

All cases were treated according to the management protocol adopted in Poisoning **RESULTS**

This was a prospective cohort study of acute zinc phosphide (Zn3P2) poisoning cases that were brought to Zagazig University Hospitals' poison treatment unit and intensive care units through the period from the beginning of July 2023 to the end of January 2024 and fulfilled the inclusion criteria. Fifty-three patients, ages ranging from 17 to 57 years, with a median age of 23 years, were included in this study. females were more than males representing 67.9% of patients. About 71.7% of patients came from rural areas. About 67.9 % of patients were students and 90.6% of patients were intentionally poisoned. All patients exposed to (Zn3P2) via ingestion. Most patients 71.7% were presented early <5 hours. As regard the

Treatment Unit (ZUH) including initial patient stabilization (ABC), decontamination was done through gastric lavage using coconut oil or castor oil along with using antioxidants such as N-acetyl cysteine (NAC) because it has been demonstrated to reduce organ toxicity, especially hepatotoxicity, by acting as a precursor or restorer of glutathione (GSH), and it has also been proven to be helpful in treating Zn3P2 poisoning. Additional supportive therapies included intravenous fluids, vasopressors, infusions of sodium (NaHCO3), bicarbonate antiemetic, antispasmodics, proton pump inhibitors, and H2 blockers. omen who were pregnant and those with co ingestion.

Statistical Methods

The Statistical Package for Social Science (SPSS) program version 28.0 (IBM, 2021) IBM crop was used to do the statistical analysis. Released in 2021: Version 28.0 of IBM SPSS statistics for Windows. NY: IBM crop in Armonk. The chi-square test, Mann Whitney (MW) test, Fisher's Exact Test, Pearson's and spearman's correlation coefficients, the receiver operating characteristic (ROC) curve analysis, the independent-samples T-test, and other tests were used.

amount of (Zn3P2), it ranged from 0.25 –4 sachets with mean 2(1-3) (Table 1). Regarding the clinical picture, 73.6% of patients were symptomatic, abdominal pain was the most, representing 61.5% of cases followed by vomiting 59%, dyspnea, hypotension and drowsiness representing 35.9%, 33.3% and 25.6 % of cases respectively (Table 1).

Regarding the vital data, mean systolic, diastolic and man arterial blood pressure were105.57, 67.64 and 79.94mmHg respectively. Mean heart rate and respiratory rate were 79.08 beat/minute and 13.45 breath /minute respectively. Mean temperature was 37.1°c and mean peripheral oxygen saturation was 98.79%. The mean arterial PH was 7.34, with 69.8% of cases had normal acid base

balance,17% had metabolic acidosis, 7.5% had respiratory alkalosis and 5.7% had respiratory acidosis (Table 2). Normal sinus rhythm was recorded in 77.4% of cases, the most frequent abnormal ECG finding among the studied cases was sinus tachycardia in 13.2% of cases, sinus bradycardia represented 5.7% and Wide QRS was 3.8% (Table 2).

Regarding routine investigations, WBCs, RBCs and platelet count were 8.4 (103/mm3), 4.67 (106/mm3) and 4.9 (103/mm3) respectively. Mean hematocrit was 39.16%. The mean serum sodium was 143.06 mmol/L and the mean serum Potassium was 3.7 mmol/L. This study revealed that PT, PTT was 12.38 sec and 33.27 sec respectively and the mean INR was 1.21. Regarding liver and kidney functions, the median ALT and AST was 12 (U/L) and 17.1(U/L) respectively. The median creatinine and urea was 0.59 (mg/dl) and 8.3(mg/dl) respectively. Mean GCS was 14.67 (Table 2).

Median total antioxidant capacity (TAC) was 0.35 mm/L. Median troponin I and CK -MB were 0.25 ng/ml and 2.23 ng/ml. There was statistically high significant relation(p<0.001) between ICU admission and all of TAC, Troponin I and CK-MB. TAC, Troponin I and CK-MB were in high significant relation with the outcome of the patients. TAC (AUC 0.829,80% sensitivity and 86% specificity), Troponin I (AUC 0.907, 90% sensitivity and 88.4% specificity) and CK-MB (AUC 0.814,70% sensitivity and 79.1 % specificity) respectively. (Tables 3,4 Figures 1,2,3). In terms of fatality, only 18.9% of patients were admitted to the intensive care unit, and no deaths were reported in the current study.

Table (1): Statistical analysis of demographic data and poisoning history in acute zinc phosphidepoisoned patients admitted to Zagazig University Hospitals (ZUH) from the beginning of July 2023 to the end of January 2024

•	No ICU admission	ICU admission	χ2	р	
	N=43(%)	N=10 (%)			
Sex:					
Female	29 (67.4%)	7 (70%)	Fisher	>0.999	
Male	14 (32.6%)	3 (30%)			
Residence					
Rural	32 (74.4%)	6 (60%)	Fisher	0.442	
Urban	11 (25.6%)	4 (40%)			
Education					
Illiterate	2 (4.7%)	10 (100%) Fisher		>0.999	
Educated	41 (95.3%)	0 (0%)			
Occupation					
Not worker	3 (7%)	0 (0%)			
Student	29 (67.4%)	7 (70%)			
Worker	1 (2.3%)	1 (10%)	0.025§	0.873	
Farmer	1 (2.3%)	0 (0%)			
Employee	9 (20.9%)	2 (20%)			
Mode of exposure					
Suicidal	35 (81.4%)	8 (80%)	Fisher	>0.999	
Accidental	8 (18.6%)	2 (20%)			
Clinical picture					
Asymptomatic	13 (30.2%)	1 (10%)	Fisher	0.258	
Symptomatic	ptomatic 30 (69.8%)				
	Median(IQR)	Median(IQR)	Z	р	
Age (year)	19 (18 – 37.2)	18 (17.75 – 30.5)	-0.368	0.713	
Came within (hour)					
Early (<5 hours)	38 (62.8%)	0 (0%)	31.225	<0.001**	
Late (>5 hours)	5 (37.2%)	10 (100%)			
Amount (sachet)					
0.25 - 0.5	12 (28%)	0 (0%)			

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	No ICU admission	ICU admission	χ2	р
1 – 1.5	12 (28%)	2 (20%)		
2 - 2.5	10 (23.3%)	4 (40%)	4.861§	0.027*
>2.5	8 (16.3%)	3 (30%)		
Unknown	3 (9.3%)	1 (10%)		

Non-significant (p>0.05), *: Significant (p<0.05), **: Highly significant (p<0.001)

Mann Whitney test(Z) Chi square test(χ 2) interquartile range (IQR) Monte Carlo test (MC) Chi square for trend test (§) Intensive Care Unit (ICU), Zagazig University Hospitals (ZUH).

Table (2): Statistical analysis of vital data, Arterial Blood Gases (ABG), Electrocardiogram (ECG) and laboratory data in acute zinc phosphide- poisoned patients admitted to Zagazig University Hospitals from the beginning of July **2023** to the end of January **2024**

	No ICU admission	No ICU admission ICU admission		р	
	Mean \pm SD	Mean ± SD		^	
Systolic blood pressure	110.12±9.03	86.00±2.108	8.334	< 0.001**	
(mmHg)					
Diastolic blood pressure	71.05 ± 7.83	53.00± 3.49	7.081	<0.001**	
(mmHg)					
Mean arterial blood	83.72 ± 8.02	63.67 ± 3.04	7.730	<0.001**	
pressure (mmHg)					
Heart rate (beat/minute)	77.0 ± 6.66	88.0 ± 15.13	-2.249	0.049*	
Respiratory rate (/min)	13.4 ± 1.87	13.7 ± 2.11	-0.454	0.652	
Temperature	37.1 ± 0.18	37.12 ± 0.27	-0.275	0.788	
PaO2	88.23 ± 9.3	81.2 ± 8.7	2.177	0.034*	
Arterial PH	7.37 ± 0.04	7.21 ± 0.08	6.056	<0.001**	
Peripheral oxygen	98.93 ± 1.06	98.16 ± 1.09	2.066	0.044*	
saturation					
ABG			χ2		
Normal acid balance	37 (86%)	0 (0%)			
Metabolic acidosis	3 (7%)	6 (60%)			
Respiratory alkalosis	1 (2.3%)	3 (30%)	16.641	<0.001**	
Respiratory acidosis	2 (4.7%)	1 (10%)			
ECG			χ2		
Normal sinus rhythm	38 (88.4%)	3 (30%)			
Sinus bradycardia	3 (7%)	0 (0%)		0.001.00	
Sinus tachycardia	2(4.7%)	5 (50%)	25.504	<0.001**	
Wide QRS	0 (0%)	2 (20%)			
WBCs (103/mm3)	4.83 ± 0.67	5.19 ± 1.02	-1.154	0.254	
RBCs (106/mm3)	4.67 ± 0.44	4.69 ± 0.42	-0.142	0.887	
Hemoglobin (g/dl)	13.4 ± 1.87	13.7 ± 2.11	-1.371	0.176	
Platelet count (103/mm3)	243.98 ± 39.76	237.4 ± 23.37	0.501	0.619	
НСТ	38.95 ± 3.53	40.03 ± 3.89	-0.853	0.398	
PT (sec)	12.3 ± 2.04	12.75 ± 1.47	-0.664	0.51	
PTT (sec)	33.59 ± 7.67	31.92 ± 4.34	0.66	0.512	
INR	1.22 ± 0.3	1.19 ± 0.23	0.327	0.745	
Sodium	143.09 ± 5.59	142.9 ± 6.32	0.096	0.924	
Potassium	3.73 ± 0.43	3.53 ± 0.76	1.153	0.254	
GCS	14.98 ± 0.15	13.40 ± 0.69	13.832	<0.001**	
	Median (IQR)	Median (IQR)	Z	р	
ALT (U/L)	12(10 - 16)	305(230.88 - 477.75)	-4.876	<0.001**	
AST (U/L)	16.8(15 - 27)	202.7(117.38-263.75)	-4.753	<0.001**	
Direct bilirubin (mg/dl)	0.17(0.12 - 0.24)	0.15(0.12 - 0.33)	-0.205	0.837	
Total bilirubin (mg/dl)	0.42(0.32 - 0.64)	0.55(0.25 - 0.78)	-0.148	0.882	
Creatinine (mg/dl)	0.57(0.49 - 0.73)	0.88(0.52 - 1.32)	-2.108	0.035*	
Urea (mg/dl)	8.1(6.5 - 10.2)	10.3(7.68 - 19.95)	-1.968	0.049*	

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Table (3): Statistical analysis of Total Antioxidant Capacity (TAC), Troponin I and Creatine Kinase-MB(CK-MB) data in acute zinc phosphide- poisoned patients admitted to Zagazig University Hospitals from the beginning of July **2023** to the end of January **2024**

	No ICU admission	ICU admission ICU admission		р	
	Mean ± SD	Mean ± SD			
TAC (mm / L)	0.38 ± 0.11	0.24 ± 0.09	3.77	<0.001**	
Troponin I (ng/ml)	0.23(0.2 - 0.29)	0.9(0.33 - 22.59)	-3.988	<0.001**	
CK-MB (ng/ml)	1.9(1.18 - 2.78)	4.13(2.64 - 46.37)	-3.524	<0.001**	

Mann Whitney test (Z) interquartile range(IQR) independent sample t test(t) $*p\leq0.001$ is statistically highly significant Zagazig University Hospitals(ZUH), Total Antioxidant Capacity (TAC), Creatine kinase –MB(CK-MB), Intensive Care Unit (ICU).

Table (4): Diagnostic Validity of Total Antioxidant Capacity (TAC), Troponin I and creatine kinase-MB in prediction of need for Intensive Care Unit (ICU) admission in acute zinc phosphide- poisoned patients admitted to Zagazig University Hospital from the beginning of July **2023** to the end of January **2024** by Receiver Operating Characteristic (ROC) curve analysis

	Cutoff	AUC	Sensitivity	Specificity	PPV	NPV	Accuracy	р
TAC	≤0.265	0.829	80%	86%	57.1%	94.9%	84.9%	0.001**
Troponi n I	≥0.305	0.907	90%	88.4%	64.3%	97.4%	88.7%	<0.001* *
CK-MB	≥6.2150	0.814	70.0%	79.1%	43.8%	91.9%	77.35%	<0.001* *

positive predictive value (PPV) negative predictive value (NPV) area under curve (AUC) $**p \le 0.001$ is statistically highly significant,, Intensive Care Unit (ICU), Total Antioxidant Capacity (TAC), Creatine kinase – MB(CK-MB)



Figure (1): Receiver operating characteristic (ROC) curve showing validity of Total Antioxidant Capacity (TAC) in diagnosis of need for Intensive Care Unit (ICU) admission in zinc phosphide (Zn3P2) poisoned patients admitted to Zagazig University Hospitals (ZUH) from the beginning of July 2023to the end of January 2024.

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Figure (2): Receiver operating characteristic (ROC) curve showing validity of troponin I in diagnosis of need for Intensive Care Unit (ICU) admission in zinc phosphide (Zn3P2) poisoned patients admitted to Zagazig University Hospitals (ZUH) from the beginning of July 2023 to the end of January 2024

Figure (3): Receiver operating characteristic (ROC) curve showing validity of Creatine Kinase Myocardial Band (CK-MB) in diagnosis of need for intensive care unit (ICU) admission in zinc phosphide (Zn3P2) poisoned patients admitted to Zagazig University Hospitals (ZUH) from the beginning of July 2023 to the end of January 2024.

DISCUSSION

In Egypt, zinc phosphide-containing rodenticides are frequently employed as grain preservatives. Because they are inexpensive and readily available, they are also frequently used as a potent suicidal tool, contributing to one-third of suicides globally. Zinc Phosphide is a highly toxic rodenticide with a high mortality rate between 37-100% [6, 23].

This work aimed to evaluate the most common prognostic parameters in acutely (Zn3P2) poisoned cases, over the period from the beginning of July 2023 to the end of January 2024 which can help the physicians evaluate the cases, predict the prognosis and decide the best plan of care, and so improving outcome of patients. The tested parameters included are total antioxidant capacity (TAC), troponin I, creatine kinase myocardial band (CK-MB), ABG, electrolytes, CBC, PT, PTT, INR, liver functions, kidney functions and ECG. Fiftythree cases were included in this prospective cohort study with inclusion and exclusion criteria.

In terms of baseline patient characteristics, the included cases' ages ranged from 17 to 57 years. A similar prevalent age group was reported by previous studies [9, 24, 25, 26] that explained this age may be more susceptible to social, occupational and economic stresses.

The majority of poisoned cases in the current study were females (67.9%) which is consistent with similar studies done by [9, 26, 27, 28] they explained this by higher liability of female to stress. On the contrary, Etemadi-Aleagha et al. [29] and Trakulsrichai et al. [30] revealed that compared to females, males were more exposed to acute phosphide poisoning.

According to the current study's findings, the majority of patients (71.7%) came from rural areas, which is consistent with research by Tassew et al. [26] and El Gendy et al. [31] who reported that most of the patients under study were from rural areas, which were agricultural communities with easy access to harmful pesticides. However, Chaudhary et al. [32] and El Masry and Tawfik [33] discovered that the majority of their patients came from cities and explained this by globalization and migration of people to these areas leading to overcrowding and exposure to more social stress.

Most included cases gave a history of suicidal exposure (90.6 %). Similarly, Abdel Wahab et al. [9] and Trakulsrichai et al. [30] reported that suicide was the most common manner of poisoning with (Zn3P2).

The estimated amount ingested ranged from 0.25 sachet to 4 sachets, with a mean value of 2(1-3), This finding was like those recognized by Abdel Wahab et al. [9] who stated that most individuals with (Zn3P2) poisoning had only been exposed to one sachet. Additionally, a statistically significant correlation has been found between the quantity of zinc phosphide and the patients' outcomes, as reported by Mashali et al. [7] who reported that the patient with acute (Zn3P2) poisoning is considered at risk if the ingested amount > 2.5 sachets.

The current study illustrated that 67.9% of our cases were students. This agrees with various studies done by Akhtar et al. [34], Sagah et al. [35] and Abdel Wahab et al. [9] who discovered that most patients with acute phosphides poisoning were students, and they attributed this to teacher scolding and low exam scores.

The time lag between ingestion and hospital presentation, less than 5h, was also reported by Wahdan and Elmadah [36] and Shahin et al. [37] who clarified that most people who attempt suicide do not truly mean to die; rather, they seek medical assistance quickly in an attempt to attract compassion and support from their loved ones. Furthermore, there was a statistically significant correlation between the patients' outcomes and the delay time. These findings align with those published by Mashali et al. [7] who reported that zinc phosphide poisoned patient is considered at risk if time till hospitalization > 5 h.

In this study, there was no statistical relationship between ICU admission and either respiratory rate and temperature, However, there was a statistically significant correlation between patients' outcomes and heart rate. Additionally, there was a statistically significant correlation between the patients' outcomes and their systolic, diastolic, and mean arterial blood pressure. Prasad et al. [38] and Yogendranathan et al. [39] found that vital signs may be used to predict the fate of Zn3P2 poisoning at presentation. However, Hassanian-Moghaddam et al. [28] claimed that the prognosis for Zn3P2 poisoning has little to do with vital signs.

Moreover, there was a statistically highly significant relation between PH and the outcome of patients as PH was significantly lower among patients who needed ICU admission, Wahdan and Elmadah [36] found that metabolic acidosis was present in 32% of cases included in their study.

In the current study, most patients were fully conscious with GCS ranged between 13 and 15 and a mean value of 14.67 ± 0.70 and there was a statistical high significant relation between GCS and outcome of patients. These findings were in accordance with Khater and Sarhan [27] who found that all (Zn3P2) studied patients were oriented and fully conscious.

The current study's findings demonstrated a statistically significant increase in ALT and AST levels among ICU patients, as well as a statistically significant relationship between the two parameters and the outcome since they are much higher among patients who required ICU care. Elevated ALT, a sign of hepatotoxicity, was discovered in numerous research by [7,27 ,40].

The result of this study revealed that serum creatinine level ranged from (0.44-1.32) mg/dl with a median value of 0.61 and serum urea between (4-78.6) mg/dl with a median value of 8.3. There was a significant relation between serum urea and creatinine level and the outcome, as it was significantly higher among patients needed ICU admission. This was in harmony with Yogendranathan et al. [39] who stated that repeated hemodialysis helped the patients recover from severe acute renal damage induced by zinc phosphide poisoning, with blood creatinine levels exceeding 1000 µmol/L.

The ECG findings among studied patients in this study revealed normal sinus rhythm (77.4%), sinus tachycardia (13.2%), sinus bradycardia (5.7%) and wide QRS complex (3.8%). There was statistically high significant relation between ECG findings of patients and the outcome as 50% of patients admitted to ICU had sinus tachycardia and 20% had wide QRS complex. Khater and Sarhan [27] matched with our research as they discovered that 50% of their patients with (Zn3P2) intoxication showed dysrhythmia and roughly half had normal ECGs. Regarding total antioxidant capacity (TAC), in this study the median TAC was 0.35 (0.26-0.43) mm / L, which decreased in most patients. This result is in accordance with Abdel Wahab et al. [9] who discovered that acute phosphide poisoning was associated with a drop in TAC. The current study's optimal TAC cutoff point was <0.265 mM/L, with an area under the curve (AUC) of 0.829, 80% sensitivity, and 86% specificity. Since TAC significantly dropped among ICU patients, there was a strong correlation between it and patient outcomes. Abdel Wahab et al. [9] said that their study's optimal TAC cutoff point was ≤0.07 mM/L, and that it was 100% sensitive and 63.54% specific for mortality. However, Halvaei et al. [41] found no difference in the value of TAC between control and treatment group with antioxidant after ALP exposure.

Troponin I is one of the excellent markers for myocardial injury. In the present study, there was a statistically significant increase in troponin I found among the ICU cases and a highly significant correlation between the cardiac troponin I and the outcome. This was consistent with study done by Abdel Wahab et al. [9] who found that individuals with severe zinc phosphide poisoning had significantly higher serum troponin I levels than patients with mild or moderate intoxication.

In the present study, there was a statistical increase in CK-MB found among the poisoned cases and a highly significant correlation between the CK-MB and the outcome. This is consistent with studies done by Mashali et al.

[7], Khater and Sarhan [27] and El-Nahhal [42] who reported that cardiac enzymes were elevated and significant for prediction of the outcome in acute (Zn3P2) poisoning. Mostafalou et al. [43] and Hakimoğlu et al. [44] are against the current work because they found that while elevated levels of several cardiac enzymes, such as CK-MB or troponin T (Tn T), might confirm cardiotoxicity and fatality, their absence did not rule out cardiotoxicity following acute exposure to zinc phosphides.

Thankfully, the current study found no fatalities and only 18.9% of patients were admitted to the intensive care unit. The appropriate supportive therapy that was started upon arrival of (Zn3P2) intoxicated patients may have contributed to this positive outcome. This was consistent with the findings of Abdel Wahab et al. [9], who reported that every (Zn3P2) intoxicated patient recovered fully and was discharged.

CONCLUSIONS

In conclusion, Troponin I, TAC and CK-MB respectively are good predictors for severity and need for ICU admission in acute zinc phosphide (Zn3P2) poisoning. They are quick, easy, reliable, and useful markers that save time so they might be useful tools for determining the prognosis of zinc phosphide.

Considerations: Ethical The study was performed after approval from the ethical committee of scientific research (Institutional Research Board "IRB" number **ZU-IRB** #10545/7-3-2023. Written informed consent was obtained from the patient, if they were conscious, or from their family members if unconscious. Throughout the whole trial, patient records were kept private

Conflict of interest: The authors declare that they have no competing interest.

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REFERENCES

 Juárez-Martínez A, Madrigal-Anaya J. D. C, Rodríguez-Torres, Y. P, Dorado-García R, Montes-Ventura D. M. and Jiménez-Ruiz A.

Mohamed, et al

Zinc phosphide poisoning: from A to Z. Toxics, 2023, 11(7): 555.

- 2-Gupta, R. C. Non-anticoagulant rodenticides. In Veterinary toxicology, 2018: (pp. 613-626). Academic Press.
- **3.** 3-Hamade H, Sahin A, Sukhn C, El Tawil C, Rizk J, Kazzi Z. and El ZahranT. Human zinc phosphide exposure in Lebanon: a case report and review of the literature. Clinical Practice and Cases in Emergency Medicine. 2021, 5(1): 50.
- 4-Trukhan V. M, Izotov A. D, Shoukavaya T. V. "Compounds and solid solutions of the Zn-Cd-P-As system in semiconductor electronics". Inorganic Materials. 2014, 50 (9): 868–73.
- 5-Hinds L. A, Henry S, VAN DE Weyer N, Robinson F, Ruscoe, W. A. and Brown, P. R. Acute oral toxicity of zinc phosphide: an assessment for wild house mice (Mus musculus). Integrative Zoology. 2023, 18(1): 63–75.
- 6-Bilics G, Héger J, Pozsgai É, Bajzik G, Nagy C, Somoskövi C. and Varga C. Successful management of zinc phosphide poisoning—a hungarian case. International Journal of Emergency Medicine. 2020, 13:48.
- 7- Mashali A. A, Salama N. H, Elsobky H. A. and Sobh Z. K. Prediction of zinc phosphideinduced hepatotoxicity and cardiotoxicity from clinical, laboratory, and radiological indicators. Environmental Science and Pollution Research. 2020, 27: 39547-59.
- 8-Alshatteri A. H, Ali G. K. and Omer K. M. Enhanced peroxidase-mimic catalytic activity via cerium doping of strontium-based metal– organic frameworks with design of a smartphone-based sensor for on-site salivary total antioxidant capacity detection in lung cancer patients. ACS Applied Materials and Interfaces. 2023, 15(17): 21239-51.
- 9-Abdel Wahab M, Shalaby S, El Awady E, Hussien R. and Salah Eldin W. Assessment of the role of total antioxidant capacity and troponin I as possible predictors for phosphidesinduced cardiotoxicity. Ain Shams Journal of Forensic Medicine and Clinical Toxicology. 2020 34(1): 82-94.
- 10. 10-Nair S. S, Surendran A, Prabhakar R. B. and Chisthi M. M. Comparison between four score and GCS in assessing patients with traumatic head injury: a tertiary centre study. International Surgery Journal. 2017, 4(2): 656-62.
- 11. 11-Dominiczak M. H. and Szczepańska-Konkel M. Chapter regulation of hydrogen ion concentration balance). Medical Biochemistry E-Book. 2014: 332-6.

- 12-Meintker L, Ringwald J, Rauh M. and Krause S. W. Comparison of automated differential blood cell counts from Abbott Sapphire, Siemens Advia 120, Beckman Coulter DxH 800, and Sysmex XE-2100 in normal and pathologic samples. American journal of clinical pathology. 2013, 139(5): 641-50.
- 13. 13-Sonntag O. and Scholer A. Drug interference in clinical chemistry: recommendation of drugs and their concentrations to be used in drug interference studies. Annals of clinical biochemistry. 2001, 38(4): 376 385.
- 14. 14-HuangX. J, Choi Y. K, Im H. S, Yarimaga O, Yoon E. and Kim H. S. Aspartate aminotransferase (AST/GOT) and alanine aminotransferase (ALT/GPT) detection techniques. Sensors. 2006, 6(7): 756-82
- 15. 15-Löhr B, El-Samalouti V, Junge W, Maatouk H, Halabi A, Fahle A. and Domke I. Reference range study for various parameters on Roche clinical chemistry analyzers. Clinical laboratory.2009, 55(11-12): 465-71.
- 16. 16-McPherson RA and Pincus MR. Henry's Clinical Diagnosis and Management by Laboratory Methods.21st ed. Saunders Elsevier.2007:1405.
- 17. 17-Salazar J. H. Overview of urea and creatinine. Laboratory medicine. 2014, 45(1): e19-e20.
- 18. 18-Laterza O. F, Price C. P. and Scott M. G. Cystatin C: an improved estimator of glomerular filtration rate. Clinical chemistry.2002, 48(5): 699-707.
- 19. Koracevic D, Koracevic G, Djordjevic V, Andrejevic S. and Cosic V. Method for the measurement of antioxidant activity in human fluids. Journal of clinical pathology .2001, 54(5): 356-61.
- 20-Pantaghini M. International federation of clinical chemistry and laboratory medicine (IFCC) scientific division committee on standardization of marker of cardiac damage. Clinical Chemistry and Laboratory Medicine.1998, 36:887-93.
- 21. 21-Elmisbah T. E. and Aiderous M. Levels of troponin and creatine kinase MB in myocardial infarction patients. Int J Med Lab Res, 2018, 3(3): 18-22.
- 22. 22-Elgazzar F. M, Shama M. A, Shoeib O. and Hafez A. S. The role of echocardiographic findings in estimating survival probability of intensive care unit admitted aluminum phosphide poisoned patients. Journal of medical toxicology. 2022, 18(2): 128-138.
- 23. 23-Kaur R, Choudhary D, Bali S, Bandral S. S, Singh V,Ahmad M. A. and Chandrasekaran B. Pesticides: An alarming detrimental to health

and environment. Science of The Total Environment. 2024, 915: 170113.

- 24. 24-Nekoukar Z, Moghimi M, Rasouli K, Hoseini A, Zakariaei Z, Tabaripour R. and BanimostafaviE. S. Suicide attempt using zinc phosphide rodenticide: A case report and literature review. Clinical case reports.2021: 9(10): e04932.
- 25. 25-GhasempouriS. K, Zakariaei Z, Hoseininejad S. M, Chinian F, Soleymanii M, Pashaei S. M. and Sadeghi M. Clinical manifestations and treatment management of hospitalized patients with zinc phosphide poisoning, Mazandaran Province, Northern Iran. BMC Emergency Medicine. 2022, 22(1): 104.
- 26. 26-Tassew S. F, Haile B. A. and Amera Birlie T. Outcome of rodenticide poisoning and its associated factors among adult patients admitted with rodenticide poisoning at the emergency unit of Debre Tabor Comprehensive Specialized Hospital, Debre Tabor, North Central Ethiopia. Open access emergency medicine.2023,15: 189-97.
- 27. 27- Khater A. and Sarhan N. Cardiotoxicity in acute zinc phosphide intoxicated patients (a prospective study). Ain Shams Journal of Forensic Medicine and Clinical Toxicology.2015, 24(1): 44-52.
- 28.
- 28- Hassanian-Moghaddam H. and Zamani N. Therapeutic role of hyperinsulinemia/ euglycemia in aluminum phosphide poisoning. Medicine.2016, 95(31): 1–7.
- 30.
- 29-Etemadi-Aleagha A, Akhgari M. and Iravani F. S. Aluminum phosphide poisoning-related deaths in Tehran, Iran, 2006 to 2013. Medicine. 2015,94(38): e1637.
- 32. 30-Trakulsrichai S, Kosanyawat N, Atiksawedparit P, Sriapha C, Tongpoo A, Udomsubpayakul U, Rittilert P. and Wananukul W. Clinical characteristics of zinc phosphide poisoning in Thailand. Therapeutics and clinical risk management. 2017, 13:335–40.
- 33. 31- El Gendy M. A, Alfadaly, N. and Mohamed, I. N. Retrospective and statistical study of pattern of acute poisoning among cases presented to emergency department of Kafr Elsheikh Governorate hospitals. The Egyptian Journal of Hospital Medicine 2018, 73(3): 6272-6282.
- 34. 32-Chaudhary S, Momin S.G, Vora D. H, Modi P, Chauhan V, Chotaliya D. An epidemiological study of fatal aluminum phosphide poisoning at Rajkot. IOSR Journal of pharmacy. 2013, 3(1): 17-23.

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- 35. 33-El Masry M.K. and Tawfik H.M. 2011 Annual report of the poison control Centre of Ain Shams University Hospital, Cairo, Egypt. Ain Shams J Forensic Med Clinical Toxicology.2013, 20(1): 10-17.
- 36. 34- Akhtar S, Rehman A, Bano, S. and Haque, A. Accidental phosphine gas poisoning with fatal myocardial dysfunction in two families. Journal of the College of Physicians and Surgeons Pakistan 2015, 25 (5): 378-379.
- 37. 35-Sagah GA, Oreby MM, El-Gharbawy RM, Ahmed Fathy AS. Evaluation of potential oxidative stress in Egyptian patients with acute zinc phosphide poisoning and the role of vitamin C. Int J Health Sci (Qassim). 2015 Oct;9(4):375-85.
- 38. 36-Wahdan A. and Elmadah E. Methemoglobinemia and intravascular hemolysis; unusual presentations of metal phosphides poisoning. Ain Shams Journal of Forensic Medicine and Clinical Toxicology. 2016, 26(1): 129-39.
- 39. 37-Shahin M, Abuelfadl A. and Zaki A. The potential role of S 100β protein in evaluation of CNS affection and prediction of mortality in acute phosphides intoxication. Ain Shams Journal of Forensic Medicine and Clinical Toxicology 2016, 26(1): 7-15.
- 40. 38-Prasad S, Karunanidhi K, Manohar V, Mohan N. AST level in zinc phosphide poisoning. Eurasian J Emerg Med .2016, 15:39– 43.

- 39-Yogendranathan N, Herath H. M. M. T. B, Sivasundaram T, Constantine R. and Kulatunga A. A case report of zinc phosphide poisoning: complicated by acute renal failure and tubulo interstitial nephritis. BMC pharmacology & toxicology.2017,18(1): 37.
- 42. 40- Marashi, S. M. What is the real cause of hepatic dysfunction after zinc phosphide containing rodenticide poisoning? Indian Journal of Gastroenterology. 2016, 35(2): 147-148.
- 43. 41-Halvaei Z, Tehrani H, Soltaninejad K, Abdollahi M. and Shadnia S. Vitamin E as a novel therapy in the treatment of acute aluminum phosphide poisoning. Turkish journal of medical sciences.2017, 47(3): 795-800.
- 44. 42-El-Nahhal Y. Accidental zinc phosphide poisoning among population: A case report. Occupational Diseases and Environmental Medicine. 2018, 6(2): 37-49.
- 45. 43-Mostafalou S, Karami-Mohajeri S. and Abdollahi M. Environmental and population studies concerning exposure to pesticides in Iran: a comprehensive review. Iranian Red Crescent Medical Journal. 2013: 15(12): e13896.
- 46. 44-Hakimoğlu S, Dikey İ, Sarı A, Kekeç L, Tuzcu K. and Karcıoğlu M. Successful management of aluminum phosphide poisoning resulting in cardiac arrest. Turkish journal of anesthesiology and reanimation.2015, 43(4): 288.

Citation

Elsayed, A., Ibrahim, M., Hussieny Mahmoud, A., Ahmed, M. G. Role of Total Antioxidant Capacity and Troponin I in Predicting Outcome of Acute Zinc Phosphide Poisoning: A Prospective Study. *Zagazig University Medical Journal*, 2025; (2635-2646): -. doi: 10.21608/zumj.2025.383600.3944