Case report

Acute Mercury Poisoning: A Case Report

ABSTRACT

Public education on poisoning and, specifically, the potential hazardous of effects of mercury are of vital is an importance important aspect for of preventive community health. A woman presented to the ED-emergency department with a three-day history of abdominal pain, diarrhea, and fever. One week agopreviously, her daughter had brought mercury in the liquid form from the her school. She had put it on the and placed it in on the heating stove. One day later, her 14-month—old infant sister baby got developed fever and died before admission to the hospital. At the time of presentation, the woman's Herblood pressure was 134/87 mmHg; temperature,

40.2°C; heart rate, 105 bpm-and regular; respirationory rate, 18 bpm; and O₂ saturation, 96%.

Nothing was remarkable on The results of an examination and routine laboratory tests were unremarkable. As Because serine serum or urinary mercury levels could not be tested in the city, symptomatic chelation treatment with *N*-acetyl cysteine (NAC) was instituted with regard to on the basis of presumptive diagnosis and the woman's recent medical history. At On the 7th day of admission, she was discharged without any sequelae or complaints. At the discharge day blood was drawn and sent On the day of her discharge, her for mercury levels which turned out was determined to be 30 μg/dL (normal-reference range: 0—10 μg/dL).

Comment [A1]: The original sentence implied that you were referring to the infant's signs; hence, I have made this revision.

Comment [A2]: Are you referring to a physical examination? Please specify.

Comment [A3]: "Serine" is an amino acid. From the context, I believe you intended to say *serum* instead and have accordingly made that change.

Comment [A4]: I'm afraid this part is slightly unclear in context. Did you instead mean that there was no facility in the hospital or surrounding area that was equipped to determine serum or urinary mercury levels? Please elaborate.

Comment [A5]: Please specify if this was the blood or serum level.

BACKGROUND

Mercury is silver-colored and exists in the liquid state at room temperature. Mercury It is available in inorganic and organic forms. All its compounds of mercury are toxic but differ in their routes of absorption, and clinical findingseffects, and responses to therapytherapeutic approaches for treating toxicity caused by them also differ. Methylmercury, the a soluble form of mercury, is neurotoxic. Elemental (organic) mercury is especially hazardous for children since it is in the liquid form and can easily be found around [1]. Acute and chronic mercury exposure represents is a potential threat to community health. Mercury poisoning can occur as a result because of occupational hazards or suicide attempts.

The clinical effects of mercury poisoning depend on the form and the route of entry to the organism. Neurologic, gastrointestinal, and renal systems are may be predominantly affected depending on the route of exposure.

This article presents the case of a 36-year-old case woman admitted to the emergency department (ED) with nausea, vomiting, and diarrhea caused by accidental inhalation of and skin exposure of to metallic mercury.

CASE PRESENTATION

A woman presented to the ED with a three-day history of abdominal pain, diarrhea, and fever.

One week agopreviously, her daughter had brought mercury in the liquid form from the school without permission from her teacher. She had played with the mercury, and then put-placed it on the heating stove and watched its vaporization. Meanwhile, while her mother breast-fed her 14-month—old sisterinfant. 24 hours—One day after this event, her baby the infant got-developed fever and died before admission to the hospital, without any specific diagnosis. The autopsy

Comment [A6]: Please check if these changes convey your intended meaning. The compounds themselves cannot have *findings*, so I have used "effects" instead. Also they cannot have responses to therapy, so I have assumed you are talking about therapeutic approaches differing according to the type of compound.

Comment [A7]: The original phrase appeared contradictory. Elemental mercury is mercury in uncombined form, whereas organic mercury would refer to an organic compound of mercury. Since you've referred to the liquid form, I assume you're referring to the elemental uncombined form.

Comment [A8]: This phrase implies that mercury in its liquid state naturally occurs in a home environment, which is not true. Did you instead mean that liquid mercury is found in common household items like fluorescent light bulbs and thermometers?

report <u>disclosed a suspected suggested</u> mercury poisoning, which might have led to cardiorespiratory collapse <u>resulting in and, eventually,</u> death-of the infant.

At the time of presentation, On examination, her the woman's blood pressure was 134/87 mmHg; temperature, 40.2°C; heart rate, 105 bpm and regular; respirationory rate, 18 bpm; and O₂ saturation, 96% with as determined by pulse oximetry at room temperature. Her fever relieved resolved after administration of 1 gr paracetamol, while arterial oxygen O₂ saturation rose to 98% with supplemental oxygen.

Nothing was remarkable in her The results of head-and-neck, respiratory, cardiovascular, or and abdominal examinations were unremarkable. A Nneurological examination did not reveal showed absence of any-tremors, paresthesia, ataxia, spasticity, or hearing and vision loss. No Nneuropsychiatric abnormalities were not identified detected.

The results of the Ccomplete blood count, and urinalysis were normal, and the levels of sodium, potassium, blood urea nitrogen (BUN), creatinine, aspartate aminotransferase (AST), alanine aminotransferase (ALT), and bilirubin levels were within the normal reference ranges as well. Chest X rayradiography and cranial computed tomography revealed no findings signs of disease.

As Because serime serum or urinary mercury levels could not be tested in the city, symptomatic chelation treatment with *N*-acetyl cysteine (NAC) was instituted with regard to on the basis of presumptive diagnosis and the woman's recent history of mercury exposure. At On the 7th day of admission, she was discharged without any sequelae or complaints. In On the same day, blood was drawn and sent for her mercury levels which turned out was determined to be 30 µg/dL (hospital laboratory's normal reference range: 0—10 µg/dL in accord with the hospital

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Comment [A9]: Please refer to comment A4.

Comment [A10]: Please remember to specify if this is the blood or serum level.

laboratory reference). Her symptoms guided the treatment and her laboratory results took three days to be officially reported.

A week after the discharge, the patient revisited the ED due to because of recurrent abdominal pain. The results of a Pphysical examination and laboratory tests results were unremarkable, and she was discharged after a 24-hour observation. A Ffollow-up was scheduled for one week later. In the follow-up visit, the patient was asymptomatic and without any no clinical findingsigns were noted. Therefore, NAC treatment was terminated discontinued after 14 treatment days from the time of her first discharge. The other children did not exhibit any manifestations of the disease.

CONCLUSION

Children are always often attracted to by elemental mercury with because of its bright-shiny gray appearance [2]. The compound element has a short half-life (two months) in the blood due to since it is rapid distribution distributed rapidly into among body compartments. Half life in the body is only two months. Almost all of the absorbed amount is excreted via through urination [3]. Mercury is used for the manufacturinge of industrial chemicals, paints, explosives, batteries, thermometers, sphygmomanometers, electronic instruments, etc. Different mMercury compounds are used as antiseptic and diuretic agents in medicine [1]. It is also an ingredient in the drug Tthiomersal, which is used to prevent contamination of vaccines.

Acute inhalations of mercury vapors can cause pneumonia, adult respiratory distress syndrome, progressive pulmonary fibrosis, and death. Also Further, elemental (metallic) mercury can readily pass into systemic circulation via alveoli present in mercury vapor or directly through the skin. It is also known to pass directly from nursing mothers to infants via breast milk [4]. In the

Comment [A11]: This sentence is slightly unclear. You have already mentioned earlier that she was given symptomatic treatment. Did you mean that she was asked to continue the same medication after discharge?

Further, in the previous sentence, you say that on the day of the discharge, the blood sample was taken and her mercury level determined. So the statement that her results took three days to be reported sounds contradictory.

Please clarify so that that I can suggest a suitable revision.

Comment [A12]: The time period was not very clear in the original sentence. Please check if the revision captures what you meant. If not, please return with a clarification and I will be happy to offer an alternative.

Comment [A13]: You have mentioned only one child other than the infant who died. If you are referring to this child, please replace this phrase with "other child."

If the woman had more children, please mention so early on in the case report so that this sentence does not confuse readers.

Comment [A14]: Since you are not referring to any particular disease but just toxicity, I suggest that you replace this phrase with "signs of mercury poisoning."

present case, Ppredominance of gastrointestinal symptoms and historical findings recent history of mercury exposure suggested intoxication with elemental mercury in the present case.

All kinds of Various neurological findings manifestations can be seen occur in chronic mercury exposure. Some effects of high—dose mercury inhalation are shown on-in Table 1_[4,5]. A recently published recommendation guideline stresses that "if the elemental mercury was recently heated (e.g., from stove top, oven, furnace) in an enclosed area, all people within the exposure area should be evaluated at a healthcare facility due to the high risk of toxicity (Grade C)" [1].

In the present case, Findings in the patient's recent history of mercury exposure played a critical role in the diagnosis in the present case. Inquiry for additional acid, alkali, arsenic, phosphorus or iron ingestion did not yield any suspicious finding. The Hhistory of exposure to mercury exposure, gastrointestinal symptoms, and suspicious death of the breast-fed baby infant led us to the presumptive diagnosis of acute mercury poisoning. It can be postulated that in the present case, neurotoxicity in the woman was prevented by the NAC treatment which that was instituted empirically based on clinical symptoms and history although blood and urine mercury levels were not determined at the time of admission.

Death of the a previously healthy baby in 24 hours prompts consideration of necrotizing bronchitis, pneumonia, or respiratory distress syndrome [7]. Inhalation of mercury by the baby infant in this case can be was thought to be the main reason cause of death.

Initial treatment The immediate precautionary measure is to keeping the isolate the patient away from the contaminated environment and the toxic agents. NAC is used for chelation of mercury, due to lack in the absence of other treatment options. Basically itNAC binds mercury by through its cysteing groups [1]. The Other mercury-chelating drugs commonly with used worldwide

Comment [A15]: "Historical findings" is not a very standard phrase. Hence, I have made this

Comment [A16]: I have not made any changes to this portion since it is a direct quote from a referenced paper.

Comment [A17]: I'm not altogether sure what you mean by "inquiry for..." Did you question the patient if she had ingested any of the items mentioned? Or were tests performed to determine these levels? Please elaborate.

Comment [A18]: This is a repetition of what you've already said at a couple of previous instances Please review if you need to omit one of the previous mentions.

Comment [A19]: You have not cited reference [6] anywhere in the text. Please correct this discrepancy.

application are dimercaprol or British anti-Lewisite (BAL), dimercaprosuccinic acid (DMSA), and 2,3-Ddimercapropropane-1-sulphfonate (DMPS) British Anti-Lewisite (BAL) (2.5 mg/kg) is also commonly used in the treatment [1,8].

This case report emphasizes the importance of public education on poisoning and specifically, the potential hazardous effects of mercury for preventive community health.

REFERENCES

1. Caravati E.M.,EM, Erdman A.R., Christinson AR, Christinson G., Nelson L.S.,LS, Woolf A.D.,AD, Booze L.L.,LL, Cobaugh D.J., Chy ka P.A.,DJ, Chyka PA, Scharman E.J.,EJ, Manoguerra A.S.,AS, Troutman W.G.;WG: American Association of Poison Control Centers. Elemental mercury exposure: an evidence-based consensus guideline for out-of-hospital management.

2. Nakyama H, Shono M, Hada S: **Mercury exanthem.** *J Am Acad Dermatol* 1984, **13:**848-52.

Clin Toxicol (Phila) 2007, 46:11-21.

3. Fischbach FT: A manual of laboratory & diagnostic testing, 4th edition. Philadelphia: J.B. Lippincott Company; 1992:214-6.

 Ford MD: Metals and Metalloids: Mercury. In Emergency Medicine: A Comprehensive Study Guide. 5th edition. Edited by Tintinalli JE, Kelen GD, Stapczynski JS. Newyork, McGraw Hill; 1999:1191-119393.

RisherJFRisher JF, Amler SN: Mercury Exposure: Evaluation and Intervention. In:
 The Inappropriate Use of Chelating Agents in Diagnosis and Treatment of Putative

 Mercury Poisoning. Weurotoxicology.26: 691-9.

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6. L.R. Goldman, M.W. LR. Shannon. 2001 MW: American Academy of Pediatrics: Committee on Environmental Health. Technical Report: Mercury in the Environment: Implications for Pediatrics. Pediatrics 2001, 108:197-205.

7. Tchounwou PB, Ayensu WK, Ninashvili N, Sutton D: Environmental exposure to mercury and its toxicopathologic implications for public health.

Environmental Toxicology Environ Toxicol 2003, 148-75.

8. Blanusa M, Varnai VM, Piasek M, Kostial K, Chelators as antidotes of metal toxicity: therapeutic and experimental aspects.

Current Medical Chemistry Curr Med Chem 2005, 12:-2771-94.

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