

DETERMINATION OF LEVELS OF LEAD AND CADMIUM CONTAMINATION IN MEAT PRODUCTS SOLD IN NORTHERN LEBANESE MARKETS

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ABSTRACT

The levels of lead and cadmium have been determined in canned and processed meat products sold in North Lebanon. Such products are normally available all year round in the markets. Never before have these meat products been assessed for their levels of toxic metals nor have they been given the deserved attention regarding their impact on human health. Using closed-vessel microwave acid-assisted digestion and graphite furnace atomic absorption spectroscopy, the levels of lead (Pb) and cadmium (Cd) were determined in 75 brands of canned and 33 brands of processed (cold cuts) meat sold in the northern part of the country. The data provided extremely important information to whether or not Lebanese individuals are exposed to high levels of such toxic metals where specifically children were found to be more vulnerable to such exposures. In canned meat, the data showed that the levels of Pb ranged from 0.00020 to 0.8161 µg/g with a mean of 0.02696 µg/g, while 29 brands were below the detectable limit. As for Cd, the data revealed levels ranging from 0.00019 to 0.1382 µg/g with a mean of 0.01557 µg/g, while seven brands were below the detectable limit. In processed meat samples, Pb concentrations ranged from 0.00025 to 0.06135 µg/g with a mean of 0.0174 µg/g of which three brands showed non-detectable levels. Concentrations of cadmium ranged from 0.0000245 to 0.0071 µg/g where the mean concentration was found to be 0.002386 µg/g. Two major parameters, amount consumed and body weight, were found to play an important role in determining whether the provisional Tolerable Weekly Intake levels (PTWIs) of Pb and Cd were exceeded or not. Specifically for canned meats, certain samples have shown that the PTWI has been markedly exceeded in children for both metals.

Keywords: Cadmium, canned meat, GFAAS, heavy metals, lead, Lebanon, microwave digestion, processed meat.

1 INTRODUCTION

Heavy metals are naturally occurring elements in the earth's crust [1]. Such metals were used thousands of years ago for several applications, but their concentration in the environment increased since the development of both agricultural and industrial fields [2,3]. Despite their known toxicity, heavy metals are still widely used nowadays. Moreover, they are transferred into the environment through anthropogenic activities such as mining, industrial processing, waste water irrigation, agricultural activities [4], transport and fuel combustion [2], iron and steel production, coal and oil combustion, waste incineration, non-ferrous manufacturing, and cement kilns [5].

Heavy metals find their way into living organisms from dietary and non-dietary exposure, where they accumulate and persist for long time periods, thus causing various health effects [6,7]. Accumulation depends on the organ of interest and on the metal's characteristics. The uptake of heavy metals by living organisms is related to the bioavailability of such elements, represented by the characteristics of the metal, the nutritional facts, and the age of the organism [5].

Lead is a naturally occurring contaminant that can be found in rock and soil [8]. For example, Pb was used for building materials, water transportation and wine sweetening, while other metals were used widely as pigments in artists' materials as well as other applications [2].

In addition, human activities contributed a great share in increasing its amount. Measures have been taken to reduce lead emission in the environment like banning or reducing the use of leaded-gasoline, phasing-out lead from household paint and solder-sealed canned food processing industry [9]. Lead may find its way into the food chain through several pathways, which may include direct deposition from air to edible plants, meat products from livestock, which have been exposed to contaminated plants, water, and air [10].

Ingested lead is transported primarily by the blood to the soft tissues. The concentration in both blood and soft tissues is relatively lower (1% and 8%, respectively, of absorbed lead) [11] than its concentration in bones (up to 90% accumulation in bones). Possessing several similarities, lead can be mistaken for calcium. Thus, bones are considered to be a long-term storage reservoir for lead, a fact that is apparent from the half-life of lead in blood and soft tissues (approximately 30 days) in comparison to its half-life in bones, which ranges from 10 to 30 years from which Pb is released slowly overtime. Pregnancy leads to an increase in blood lead concentration for the mother caused by the release of lead from maternal bones when the calcium is mobilized to construct the fetal skeleton [9]. In addition, maternal lead is transferred to the fetus through the placenta, and later during breast feeding. The most important targets for lead toxicity in humans are the blood, nervous, and cardiovascular systems, as well as the kidneys [12]. The International Agency for Research on Cancer (IARC) has classified lead as a class 2A carcinogen [8]. For adults, lead is associated with neurological toxicity that was found to influence the central information processing mainly leading to visiospatial organization disorder and affects short-term verbal memory. It also causes psychiatric symptoms and manual agility impairment [8]. Besides neurological toxicity, lead may cause other health problems such as delayed sexual maturation, affects the reproduction by decreasing the number and quality of sperm and by increasing the frequency of abortions, developmental delays, hypertension caused by reduction of nitric oxide (NO) concentration in blood, impaired hemoglobin synthesis, tiredness, sleeplessness, anemia, irritability, osteoarthritis, headaches, constipation, weight loss, joint pain, and muscle weakness [8]. According to the United States Food and Drug Administration (FDA), provisional total tolerable daily intake levels (PTTILs) of Pb are set at 0.025 $\mu\text{g/g/day}$ for pregnant women, while for infants, it was set at 0.006 $\mu\text{g/g/day}$ [13]. Meanwhile, the Joint Food and Agricultural Organization (FAO) and the World Health Organization (WHO) Expert Committee on Food Additives (JECFA) have set the provisional tolerable weekly intake (PTWI) of lead at 0.025 $\mu\text{g/g}$ body weight (bw) for all age groups [13].

Such differences in standard levels arise first from the FDA's recommendations in being extremely cautious, while the World Health Organization argues that our bodies will metabolize and excrete lead efficiently as long as the amount of lead that we are exposed to does not exceed its PTWI [14]. For this reason, the JECFA safe levels are more widely accepted by scientists.

Cadmium (Cd), on the other hand, is a naturally occurring, relatively volatile, silvery-white soft metal with two valence states. It is classified as non-essential toxic metal having the ability to bioaccumulate in living organisms and is not easily excreted [10]. Cadmium plays a significant role in various anthropogenic applications, where it is mixed with other metals (copper, silver, tin and lead) to form metallic alloys [15]. Cd finds its application mostly in nickel/cadmium batteries, plastics stabilizers and pigments, electronic compounds, and is used as a protective coating to prevent the corrosion of other metals. Cadmium can also be found as an impurity in phosphate fertilizers, detergents and petroleum products [10]. Sources of human exposure to cadmium may be attributed to industry, air, soil, and water, which all pose an exposure risk to animals, their products as well as other food commodities [8,10,11,16,17].

Human exposure to cadmium occurs through inhalation, dermal contact, but to a greater extent through ingestion. Due to the fast uptake of cadmium by leafy vegetables and crops from contaminated soil, cadmium can be readily transferred to animals grazing on such contaminated plants, whereby it accumulates mainly in their tissues [18]. This leads to its presence in the food-chain, thus making meat and offal an important route of exposure to cadmium [19]. Absorption depends on the solubility of cadmium in the ingested compounds and also on the nutritional status of the organism where iron and calcium deficiency leads to higher cadmium absorption [20]. Possessing great similarities to zinc, a bio-essential nutrient for living organisms, Cd has the ability to displace zinc in important enzymatic and organ functions leading to the impairment or induction of such tasks. The kidney and liver are considered as target organs for cadmium accumulation although the liver is less sensitive to cadmium than the kidneys [21]. The half-life of Cd may range from 6 to 38 years depending on the organ in which they accumulate (6–38 years in human kidney and 4–19 years in human liver) [10]. Chronic exposure may lead to renal failure, presence of serum proteins in urine, osteomalacia, osteoporosis, lung dysfunction leading to lung disease, and bone damage, while acute exposure may lead to headache, vomiting, diarrhea, reduced body weight, ulcers, hemorrhages, testicular weakening, reddening of intestinal track and stomach [21], chest pain, muscle weakness, pulmonary edema, bronchitis, respiratory failure, renal failure, affects also the liver, cardiovascular system, and nervous system [20]. According to a study on the effects of cadmium on kidney cells, cadmium was found to induce changes in the cells related to the proliferation of pre-neoplastic cells, possibly leading to cadmium-induced carcinogenicity [21]. According to the JECFA, a PTWI of cadmium is calculated on the assumption that 5% of all the dietary cadmium is absorbed and is proposed to be applied to long-term consumption. This PTWI is established to be 0.007 $\mu\text{g/g}$ bw for cadmium per week [21]. The U.S. Environmental Protection Agency (EPA) established a reference dose of food intake of cadmium of 0.001 $\mu\text{g/g}$ bw/day, and the limit of intake of cadmium through drinking water of 0.0005 $\mu\text{g/g}$ bw/day. Such difference of intake of reference doses between food and water is due to the difference in the levels of absorption of cadmium [8].

As mentioned, there are many routes for human exposure to lead and cadmium. Individuals should be aware of the health effects resulting from the consumption of heavy metal contaminated food products. While the food chain constitutes a main source of accumulation of such heavy metals, meat products represent an essential part of the human diet and are consumed mostly on daily basis. This makes them a continuous source of exposure to lead and cadmium, thus leading to various diseases linked to the toxicity of such metals. Accordingly, it is important to determine their levels in meat and meat products to assess the health risks derived from their consumption. The level of heavy metals in meat and meat products depend on several factors such as the environmental conditions, grazing land, and the genetic characteristics of the animal's organism [22].

A recent study carried out in the central Bekaa plain of Lebanon [18] has revealed that soil metal contamination was mostly due to the use of contaminated water, to compensate for water shortage and to the heavy use of fertilizers. Such conditions can negatively affect animals that feed on plants grown in such lands, whereby consumption of animal meat products can also expose humans to such toxic metals. A more related study [23] was carried out to assess the impact of metal cans on food quality. However, the study does not comprehensively cover all canned meat products, nor it included any of the processed kind (cold cuts) and focuses mainly on metal contents of cans and its effects. Since there are no food safety programs in Lebanon that monitor consistently the quality of all food products, whether imported or local, and since very little research information exists on toxic metal contamination

in processed and canned meat, the objective of this work is to determine the contamination level of lead and cadmium in as many of such products available to the consumer, compare them with safe standard levels, and to also create awareness and perhaps ignite the interest of private or governmental sectors to become seriously engaged in such studies, all for the safety and development of better food quality.

2 MATERIALS AND METHODS

2.1 Apparatus

Prior to digestion, samples were dried overnight in a programmable oven (Venticell, W.P.Katul). Samples were digested using Thermo Ethos 1 microwave digestion oven (Milestone, Italy). For metal analysis, a Thermo-Electron M series graphite furnace atomic absorption spectrometer (GFAAS) equipped with deuterium and Zeeman background correction (Zeeman Furnace GF95Z) and an autosampler (FS95) were used (Thermo Electron Corporation, Germany). The sample atomization is carried out in specialized graphite tubes (Thermo Elemental Omega Platform Extended Lifetime Cuvettes). Coded hollow cathode lamps for lead and cadmium (Thermo-electron Corporation, Germany) were also used. 99.999% pure argon gas (Chehab Industrial and Medical Gases S.A.L., Lebanon) was used as an internal inert gas for the GFAAS with a flow rate of 300 mL/min [1,3].

2.2 Reagents

Throughout the digestion and detection procedures, double distilled de-ionized Water (ddH₂O) obtained by water treatment using Milli-Q system (Millipore) was used. Working standards for lead and cadmium were prepared using standard solutions supplied by Romil-Pure chemistry (standard solutions of 1000 ppm element reference solution). Ultrapure concentrated nitric acid (65%) was used for the sample digestion as well as for the preparation of wash solution (1%, v/v) and diluents (0.5%, v/v) for the GFAAS [1,3]. Ascorbic acid (99.99% pure, trace metal basis, Sigma-Aldrich) was used as the matrix modifier for lead detection, while magnesium nitrate (99.99% pure, trace metal basis, Sigma-Aldrich) was used as the matrix modifier for cadmium detection in the GFAAS. Certified reference material (CRM) (SRM1577C Bovine Liver NIST National Institute of Standards and Technology, USA) was used in part for quality control and validation of all analytical procedures throughout.

2.3 Optimization conditions

Both cadmium and lead are volatile elements in nature with a wide range of volatility that is dependent on the amount of heat applied. Thus, caution should be taken into consideration for choosing the optimum conditions for the heating program of the GFAAS and whether or not modifiers should be used.

2.3.1 Lead and cadmium optimization

The atomization temperature for Pb was very critical especially in the range of 1400–1600°C. Nevertheless, the type of matrix modifier also plays an important role when using complex samples. Upon careful optimization, the optimum atomization temperature was found to be best at 1400°C while the best matrix modifier was found to be 1% ascorbic acid. As for Cd,

it was shown that the ashing temperature was of importance where an increase in the ashing temperature revealed a decrease in the absorbance signal and vice versa. This pattern was observed in the presence and absence of matrix modifiers. Optimum results were obtained by using an ashing temperature of 350°C and a 2000 ppm solution of 99.99% pure and trace metal basis $\text{Mg}(\text{NO}_3)_2 \cdot 6\text{H}_2\text{O}$ as the matrix modifier.

2.4 Materials

All plastic and glass-ware were washed with soap and tap water, rinsed with ddH_2O , and then soaked overnight in 10% (v/v) nitric acid solution [1,24,25]. Prior to usage, any item was washed three times with ddH_2O . Acid-digested samples were stored in 50-mL polypropylene conical tubes (LaboTech, Lebanon), which were soaked in 10% nitric acid solution and washed with ddH_2O prior to use. Reagents and sample cups (LaboTech, Lebanon) for the autosampler of the GFAAS were also soaked and washed thoroughly with ddH_2O before filling them with the samples to be analyzed.

2.5 Samples and treatment

Samples of canned and processed meat (cold cuts) were purchased from various supermarkets in North Lebanon. Seventy-five different brands of canned meat and 33 of processed that can be usually found all year long in markets were considered for this study. All canned meat samples were classified under five categories namely chicken (Ch), beef (B), pork (P), duck (D), and mixed (M, mixed beef/pork/herbs and/or olives). Similarly, the processed which included cold cuts such as mortadella, ham, salami, pepperoni, and such, were characterized under turkey (T), pork (P), mixed chicken and turkey (C+T), mixed beef and chicken (B+C), mixed pork and beef (P+B), and mixed pork and chicken (P+C). For all samples studied, whether canned or processed, and in order to get a representative sample, a few grams of tissues were taken from separate subsamples of the same brand, pooled together, homogenized by cutting into fine pieces by using pre-soaked disposable plastic knives, placed in small pre-soaked Petri dishes, then dried at 70°C overnight until a constant sample weight is obtained [1,24,26]. Dried samples were then stored in tightly sealed acid-treated storage 50-mL conical tubes for later use. A quantity of 0.5 g of each dried sample was transferred to a high temperature high-pressure Teflon reaction vessel specific for Ethos1 microwave oven, followed by the addition of 7 mL of ultrapure concentrated nitric acid (65%) and 1 mL of 30% concentrated pure hydrogen peroxide [1,24,27]. Similarly, and for quality control purposes, 0.5 g of the CRM was also included in each batch and treated in the same manner as any other sample. All samples, as well as CRMs, were run in triplicates. As for the control blanks (two in each batch), the vessels contained 7 mL acid and 1 mL H_2O_2 , except where the 0.5 g sample weight, which was substituted with 0.5 mL ddH_2O . All prepared reaction vessels were left under a fume extraction hood for at least 15 min to allow the formed gases to escape. The vessels were then sealed and placed inside the microwave oven and digested using a three-step program. During the first step, samples were heated from room temperature to 200°C in 30 min at 1000 W. The second step consisted of maintaining the samples at 200°C for 30 min at 1000 °. In the final step, samples were brought to room temperature. The clear digested samples were transferred into 25-mL volumetric flasks and diluted with ddH_2O up to the mark prior to their transfer to 50 mL pre-soaked pre-cleaned polypropylene tubes and were finally stored in the refrigerator before analysis with GFAAS [1,3].

2.6 Sample analysis by GFAAS

Before analyzing the samples in the GFAAS, the instrument was programmed according to the type of analyte to be detected. Following metal selection, the optimized parameters of the specific metal were set in the software of the instrument. Calibration standards for Pb and Cd were carefully prepared by diluting a certain amount taken from their stock solutions (1000 ppm) down to 10 ppb (mother solution) for lead, and similarly to 1 ppb (mother solution) for cadmium, by using subsequent 10-fold dilutions each time so as to diminish the analytical preparation errors as much as possible. The mother solutions were used by the GFAAS's auto-sampler to create automatically a specified calibration curve specific for each metal. The number of calibration points was automatically set by the instrument and included five points for each metal. 0.5% (v/v) nitric acid solution was prepared for the auto-sampler for use as diluent, while 1% (v/v) nitric acid solution was prepared and used by the auto-sampler to wash the injection tube between each dilution, as well as the samples to be analyzed, so as to prevent any cross-over contamination between any of them [1,3].

From each digested sample, as well as digested CRMs, 1 mL aliquots were taken and were placed in a 1-mL pre-cleaned polypropylene GFAAS sample cups, which were loaded onto the autosampler of the GFAAS. As well, standard mother solutions, diluents, and matrix modifiers were placed in 20-mL pre-cleaned polypropylene reagent cups and were also loaded onto the auto-sampler. Considering the amount of time that the GFAAS needs to analyze each sample, up to 30 samples were included in each run per each working day. All specimens were run in batches with digested blanks and CRMs. Digestion blanks were used to test for the presence of any possible contamination in the digestion procedure, while CRMs were included to calculate the recovery of the digestion process [1,3,24].

For further quality control, the instrument was programmed to periodically re-measure sample blanks and standards from the calibration curve after every 10 samples to check for any instrumental variations during the analysis, thus ensuring fidelity and consistency of the data.

3 RESULTS AND DISCUSSION

The aim of this work was to first collect and analyze all possible meat brands normally sold in Lebanese markets for their contents of Pb and Cd. Second was to compare contamination levels with standard values set by health organizations. Third, and most important, was to make the general population, aware of the health hazards that may arise from consuming such products that may potentially ignite the interest of private and governmental health organizations to engage in such studies. For this purpose, the contents of lead and cadmium in 75 brands of canned and 33 brands of processed meat found in the Lebanese markets were determined.

As in all analytical procedures, quality control (QC) is a necessary and a crucial part. The accuracy and precision of the microwave digestion oven and the GFAAS as well as the methods used have all been examined using CRM, blank samples, and replicates within every batch. The recovery percentages of the digested CRMs were all found to fall within the accepted analytical range of 80–120%, thus confirming the validity of the analytical procedure and the authenticity of the results. All digested blanks and prepared standards passed the QC checks set by the software. The calibration curves showed excellent linearity, with R^2 values ranging between 0.9954 and 0.9998, thus demonstrating the excellent performance of the instrument over the broad range of concentrations used.

In general, results from the analyzed samples have shown wide variations of Pb and Cd concentrations (Table 1). Concentrations were based on dried weight (μg of metal/g of dried sample) so as to eliminate the water content differences between all samples [1,24,26]. The numbers of brands in each category, their minimum, maximum, mean detectable concentrations, as well as the number of non-detectable samples, have all been summarized in Table 1.

Forty-six samples (61% of analyzed brands) of the canned meat were found to exhibit various levels of Pb contamination, while 68 samples (91%) showed Cd contamination. For the processed meat, 91% exhibited various Pb concentrations, while 94% revealed the presence of Cd. According to the EU directive 466/2001 [3], which regulates the amounts of Pb and Cd in meat and their products, the maximum allowed levels (MAL) of Pb and Cd in meat and meat products were set at 0.100 and 0.050 $\mu\text{g/g}$, respectively. This study has shown that certain sample brands have markedly exceeded such levels.

The data also showed that the concentrations of the studied metals are generally much higher in canned meat in comparison to processed ones and that there was no correlation between the concentrations of both metals. The data also showed non-detectable levels for both metals in canned and processed meat. This suggested that toxic metal contamination was not due to a known constant factor (processing and canning) but rather due to many other factors, which may be attributed to the source of meat (i.e. type, origin, and diet of the corresponding animal). When the highest concentrations of Pb and Cd in canned and processed meat have been compared, the data showed that the canned meat contained 13.3 times more Pb than in the processed meat, meanwhile canned meats contained 19.4 times more Cd than in the processed ones. This difference may be attributed to bioaccumulation either from the environment or during the feeding period before slaughter or even from the leaching of the

Table 1: Data summary of lead and cadmium concentrations (in $\mu\text{g/g}$) analyzed in canned and processed meat products.

	Canned meat					Processed meat			
	Beef	Chicken	Duck	Mixed	Pork	Mixed	Pork	Turkey	
Lead	Number of brands	25	22	2	12	14	8	18	7
	Number of non-detectable brands	10	9	2	3	5	1	1	1
	Mean detectable value ($\mu\text{g/g}$)	0.00976	0.05080	–	0.01270	0.03980	0.01852	0.02030	0.01440
	Minimum detectable value ($\mu\text{g/g}$)	0.00030	0.00140	–	0.00020	0.00120	0.00777	0.00024	0.00347
	Maximum detectable value ($\mu\text{g/g}$)	0.06040	0.81610	–	0.03670	0.41390	0.05430	0.06300	0.02660
Cadmium	Number of brands	25	22	2	14	12	8	18	7
	Number of non-detectable brands	3	1	1	2	0	0	2	0
	Mean detectable value ($\mu\text{g/g}$)	0.01570	0.01290	0.00524	0.02140	0.01390	0.00302	0.00161	0.00340
	Minimum detectable value ($\mu\text{g/g}$)	0.00010	0.00068	–	0.00040	0.00476	0.00026	0.00002	0.00118
	Maximum detectable value ($\mu\text{g/g}$)	0.08930	0.05580	0.00524	0.13800	0.03270	0.00497	0.00630	0.00716

metals from the can itself [28], all of which are difficult to assess. Based on the above, processed meat products might seem to be a better choice for consumption; however, this one time study is not conclusive. Therefore, constant monitoring, to include every stage of the preparation process is a must and awareness campaigns to ensure quality are more than necessary. A number of recent studies have been conducted on fresh meat products and similar results were obtained suggesting that individuals may suffer health risks due to consumption of such products [29–31]. In the first study [29], fresh samples of chicken and beef showed to contain concentration of lead and cadmium which have exceeded the allowable levels. The second study [30] targeted the assessment of lead and cadmium levels in different types of fresh tissues from chicken where in some cases the levels suggested that individuals may be at risk from ingesting such products. Another report [31] presented the determination of lead and cadmium levels in Luncheon meat where levels of such metals were found to be a possible health hazard. Such observations are coherent with our results, which also suggest that individuals may be at risk from consuming such products.

The analyzed levels of lead in the canned meat samples have been presented in Fig. 1. The samples were grouped in categories according to the type of meat studied and in increasing order of concentrations detected within and between each category. Following the figure from left to right; Pb levels in duck samples have been found to be non-detectable (nd), while they varied from nd to 0.0367 $\mu\text{g/g}$ in the mixed, nd to 0.0604 $\mu\text{g/g}$ in beef, nd to 0.4139 $\mu\text{g/g}$ in pork, and finally, nd to 0.8161 $\mu\text{g/g}$ in chicken.

The results were comparable to studies elsewhere between 1983 and 2006 [32–40] and clearly indicated that certain samples (P14 and Ch21) in the pork and chicken categories exceed the Pb MAL value [40,41] and may pose a potential risk from Pb exposure when consumed.

Similarly, cadmium levels in canned meats have been analyzed and presented in Fig. 2. Once again, the results have been arranged in categories according to the type of meat studied

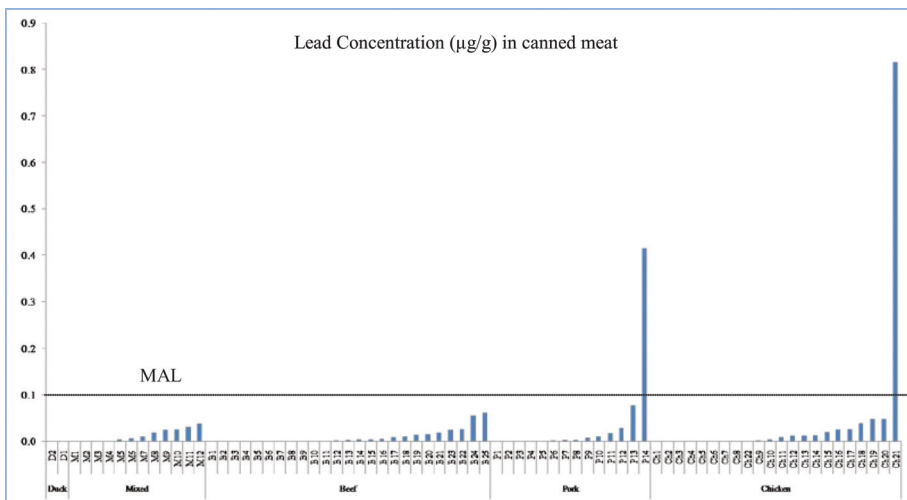


Figure 1: Lead concentration in canned meat of different categories. Concentrations of Pb are arranged from lowest to highest, within and between each category. Due to the high Pb concentration in Ch21 (0.8161 $\mu\text{g/g}$), concentrations below 0.0063 $\mu\text{g/g}$ could not be observed. The maximum allowable level (MAL) of Pb in meat (0.1 $\mu\text{g/g}$) is also shown.

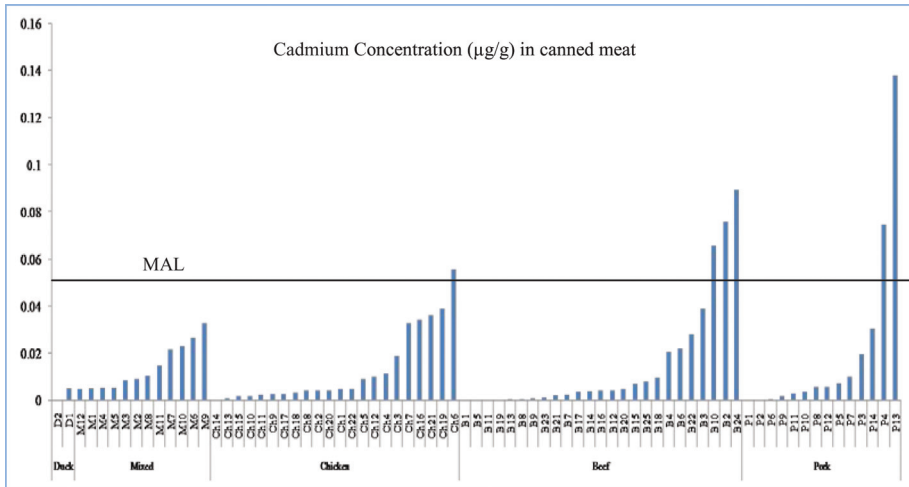


Table 2: Estimated weekly intake of Pb and Cd for canned meat.

Metal	Category	Highest concentration ($\mu\text{g/g}$)	Estimated weekly intake(g)	Weekly intake (70-kg male) ($\mu\text{g/g bw}$)	Weekly intake (60 kg-female) ($\mu\text{g/g bw}$)	Weekly intake (14 kg 2-years old) ($\mu\text{g/g bw}$)
Pb	Duck	0	400	0	0	0
			800	0	0	0
	Mixed (M13)	0.0367	400	0.000210	0.000240	0.00104
			800	0.000420	0.000490	0.00209
	Beef (B25)	0.0604	400	0.000350	0.000400	0.00173
			800	0.000690	0.000810	0.00345
Pork (P14)	0.414	400	0.002360	0.002760	0.01183	
		800	0.004730	0.005520	0.02365	
Chicken (Ch21)	0.816	400	0.004660	0.005440	0.02332	
		800	0.009330	0.010880	0.04663	
Cd	Duck (D1)	0.00524	400	0.000029	0.000035	0.00015
			800	0.000059	0.000069	0.00029
	Mixed (M9)	0.0327	400	0.000190	0.000220	0.00093
			800	0.000370	0.000440	0.00187
	Chicken (Ch6)	0.0558	400	0.000320	0.000370	0.00159
			800	0.000640	0.000740	0.00319
	Beef (B24)	0.0893	400	0.000510	0.000590	0.00255
			800	0.001020	0.001190	0.00510
	Pork (P13)	0.138	400	0.000790	0.000920	0.00394
			800	0.001570	0.001840	0.00788

intake may reach $0.04663 \mu\text{g/g}$ of bw (186.5% of the PTWI). Similarly, with a PTWI for Cd of $0.007 \mu\text{g/g bw}$ for all age groups [21], two meat samples were of concern especially for children. The consumption of 400 g of pork (P13) alone with a concentration of $0.138 \mu\text{g/g}$ has yielded an estimated weekly Cd intake of $0.00394 \mu\text{g/g bw}$ corresponding to 56% of the set PTWI. The consumption of 800 g instead has exceeded the set PTWI by 125% for children. Similarly, the consumption of 800 g beef (B24) alone under the same conditions caused 73% of the PTWI to be reached in children. Such observation suggests that awareness and regulation of consumption can play a powerful role in determining the risk of exposure to such toxic elements, especially after knowing that such toxic metals are present in our food chain.

The current situation presents a crucial stage where full awareness is required in the prevention of exposure to such health hazards especially that heavy metals first tend to bioaccumulate in the human body and second are taken from more than one source. In light of this, whether such products (canned and processed) are prepared locally or imported from elsewhere, it is crucial for the concerned authorities to place regulations for every step of the preparation procedure and before making the product available for consumers.

Figures 1 and 2 clearly show that there is no relationship between the presence of lead and cadmium within any given sample. In other words, samples that contained high levels of Pb

are not necessarily high in Cd, and vice versa. This may suggest once again that Pb and/or Cd contamination is not due to some constant factor such as the canning process alone, and that any such contamination might be due to the food, water intake, and other sources of environmental pollutions through which the animals have been exposed to during the bringing up process or during storage, processing and transportation.

In the processed meat brands, the levels of lead analyzed have been presented in Fig. 3 from lowest to highest concentrations under several categories. Based on the presented figure, the lowest Pb concentrations are exhibited in turkey samples (nd to 0.0266 $\mu\text{g/g}$), followed by processed mixed (nd to 0.0543 $\mu\text{g/g}$) while the highest Pb concentration were found in pork samples (nd to 0.0613 $\mu\text{g/g}$). None of the samples have exceeded the MAL of Pb, nor have any shown to be a potential health hazard.

Cadmium was also analyzed in processed meats and the results have been presented in Fig. 4 in ascending order of concentrations detected. In the mixed category, the lowest detected level of Cd (0.000265 $\mu\text{g/g}$) was found in (C+T)1, while the highest (0.00497 $\mu\text{g/g}$) was found in (C+T)2. For pork, the concentrations ranged from non-detectable (P2, P5) to 0.0063 $\mu\text{g/g}$ (P15) while Turkey samples ranged from 0.00118 $\mu\text{g/g}$ (T3) to 0.0071 $\mu\text{g/g}$ (T7).

Recalling that the PTWI for Pb is 0.025 $\mu\text{g/g}$ of body weight, the calculations presented in Table 3 show that even with the consumption of 800 g of the highest contaminated pork sample (P18) by a 2-year-old child reveals a maximum estimated weekly intake of 0.0035 $\mu\text{g/g}$ bw, which corresponds to 14% of the set PTWI. Similarly, for T7, with a Cd concentration of 0.0071 $\mu\text{g/g}$, the consumption of 800 g by a 2-year-old child will only yield 5.7% (0.4 $\mu\text{g/g}$) of the PTWI (0.007 $\mu\text{g/g}$ bw). Accordingly, none of the samples measured in the processed meat category seemed to exhibit any high levels of lead based on the proposed estimated weekly consumption, and thus are considered to be much safer for consumption in comparison to those of the canned ones. Again, the results showed no relation between levels of Pb and Cd, which may be due to the same reasons discussed earlier and are difficult to assess.

The highest concentration from each category was considered to calculate the amount of Pb intake based on consumption of 400 or 800 g of meat type studied.

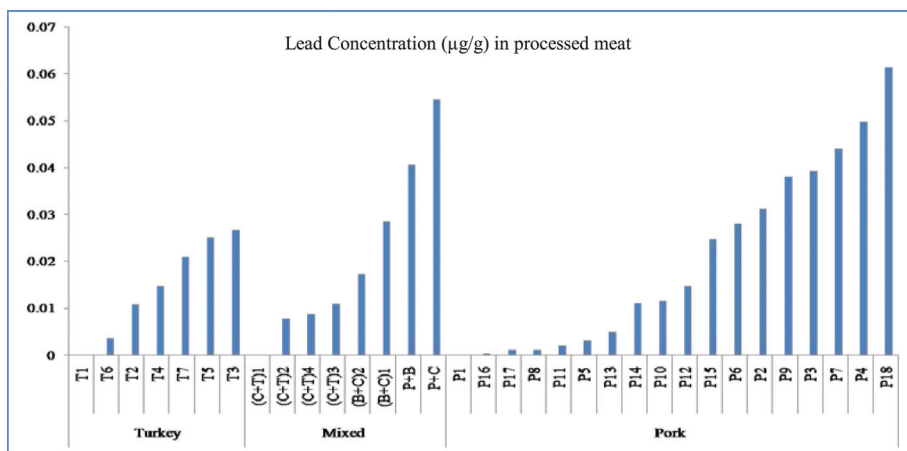


Figure 3: Lead concentration ($\mu\text{g/g}$) in processed meat (T: turkey, P: pork, C+T: mixed chicken and turkey, B+C: mixed beef and chicken, P+C: mixed pork and chicken, and P+B: mixed pork and beef).

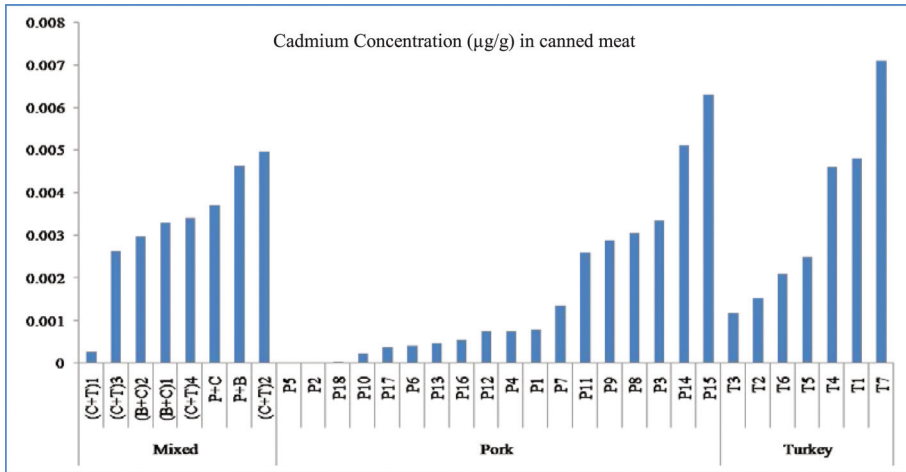


Figure 4: Cadmium concentrations (µg/g) in processed meat. The analyzed samples were arranged from lowest to highest Cd concentration within each category.

Table 3: Estimated weekly intake of Pb for processed meat.

Metal	Meat type	Highest concentration (µg/g)	Estimated weekly intake (g)	70-kg male (µg/g bw)	60-kg female (µg/g bw)	14-kg 2-years old (µg/g bw)
Pb	Turkey (T3)	0.0266	400	0.000150	0.000170	0.000760
			800	0.000300	0.000350	0.001520
	Mixed (P+C)	0.0543	400	0.000310	0.000362	0.001550
			800	0.000620	0.000720	0.003100
	Pork (P18)	0.0613	400	0.000350	0.000410	0.001750
			800	0.000700	0.000810	0.003500
Cd	Mixed (C+T)2	0.00497	400	0.000028	0.000033	0.000142
			800	0.000056	0.000066	0.000284
	Pork (P15)	0.0063	400	0.000036	0.000042	0.000180
			800	0.000072	0.000084	0.000360
	Turkey (T7)	0.0071	400	0.000040	0.000047	0.000200
			800	0.000081	0.000094	0.000400

Lebanon is one of the countries that have recently been engaged in the process of performing total dietary studies (TDS) as a reply to the EFSA/FAO/WHO questionnaire on national TDS approach [42]. However, the conducted studies [27,43] are not enough to state whether a few or all of our samples can be considered as a potential health risk. Therefore, there is absolutely a great need for an international TDS procedure synchronization and application. Ultimately, and for this reason, one can consider that samples low or free of Pb and Cd can be considered the safest to consume, but nevertheless, this cannot be taken as a general rule since constant monitoring and screening is required prior to allowing the marketing of such products. According to the European Environment and Health Information system, most European countries had an adult Pb intake levels ranging between 10% and 30% of the lead's

PTWI and sometimes higher [44]. It is not clear which TDS to follow as a guide since the study done in Lebanon does not agree with that of the European countries and is somehow surprisingly low perhaps due to non-comprehensive nature of the TDS carried. In addition, Lebanon is a developing country in which no environmental or health regulations exist, nor does it have any food safety monitoring programs but yet its Pb intake is somehow much lower (3.2%) than that of the European countries (10–30%). In the worst case and if one takes into consideration both TDSs, two canned brands (Ch21 and P14) can be considered unsafe for consumption specifically for children. If one considers the European TDS where 30% of the Pb PTWI has been reached from the diet (i.e. 0.0075 µg/g), the remaining intake to reach 100% (17.5 µg/g) will not only be reached by consuming Ch 21 and P14, but can be markedly exceeded beyond the safe levels.

Regarding cadmium, its cadmium intake through the food chain in the Lebanese TDS is equal to 21.7% of the PTWI (0.007 µg/g bw), which is much lower than the intake percentage found in Europe that ranges between 40% and 60%. Taking into consideration both TDSs, only three canned brands (Ch6, B24, P13) may be considered unsafe for consumption especially for children. Taking into consideration the European TDS, Ch6, B24, and P13, are three brands that can compensate for the remaining amount in reaching 100% of the PTWI and perhaps higher. Therefore, such brands can also be considered to be unsafe for children.

Since the study shows that certain samples can be a real health hazard, it is therefore crucial for the concerned authorities to regulate such products on all levels by setting regulatory standards in order to manage and control such contaminants. In addition, fresh unprocessed samples should be also studied in the same region so as to compare with the current study and unleash any differences that might exist in the level of their contaminations.

4 CONCLUSION

The amount of research of toxic metal contamination in foodstuffs in Lebanon has not been given enough attention and only few reports exist within the last three decades. For this purpose, the contamination levels of Pb and Cd were assessed in meat products that are normally available to consumers all year round. The data provided extremely important information to whether or not Lebanese individuals are exposed to high levels of such toxic metals where specifically children were found to be more vulnerable to such exposures. Sixty-one percent of the analyzed brands of canned meat were found to have various levels of Pb contamination, while 91% showed Cd contamination. As for the processed, 91% exhibited various Pb concentrations, while 94% revealed the presence of Cd. Of those samples, and specifically in the canned category, two brands have exceeded the MAL for Pb while six brands have exceeded the MAL for Cd. Regarding the processed category none of the samples have exceeded the MAL of either metal.

Two major parameters, amount consumed and body weight, were found to play an important role in determining whether the PTWIs of Pb and Cd were exceeded or not. Children were found to be the most vulnerable.

Overall, the results of this study (and similar studies) have demonstrated the presence of lead and cadmium contamination in the food chain. Knowing that such toxic metals tend to bioaccumulate and may be taken from more than one source present a crucial stage in reference to health hazards. Such observation necessitates the continuous monitoring for levels of toxic metals in food products. Food products must be labeled by official organizations to indicate the levels of heavy metals present. Awareness campaigns, from official or private organizations, are a must to help citizens choose safer products so as to minimize exposure risk to such toxic elements.

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