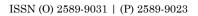
### Journal of Medical Research and Health Sciences

Received 12 May 2021 | Revised 18 June 2021 | Accepted 30 July 2021 | Published Online 13 Aug 2021

**Trace Element Contents in Thyroid of Patients with Diagnosed Nodular** 

**Goiter Investigated by Instrumental Neutron Activation Analysis** 

DOI: https://doi.org/10.52845/JMRHS/2021-4-8-5 JMRHS 4 (8), 1405–1417 (2021)



**Open Access Journal** 



JMRHS JOURNAL

#### **RESEARCH ARTICLE**

Vladimir Zaichick<sup>\*</sup>

Russia

<sup>1</sup>Radionuclide Diagnostics

Department Medical Radiological

Research Centre Korolyev St. 4.

Obninsk 249036 Kaluga Region,

Abstract

**Background :** Nodular goiter (NG) is an internationally important health problem.

**Objectives**: The aim of this exploratory study was to examine the content of ten trace elements (TE): silver (Ag), cobalt (Co), chromium (Cr), iron (Fe), mercury (Hg), rubidium (Rb), antimony (Sb), scandium (Sc), selenium (Se), and zinc (Zn) in the normal thyroid and in the thyroid tissues with diagnosed colloid NG.

**Methods :** Thyroid tissue levels of TE were prospectively evaluated in 46 patients with NG and 105 healthy inhabitants. Measurements were performed using non-destructive instrumental neutron activation analysis with high resolution spectrometry of long-lived radionuclides. Tissue samples were divided into two portions. One was used for morphological study while the other was intended for TE analysis.

**Results :** It was found that contents of Ag, Co, Cr, Fe, Hg, Sc, and Zn were significantly higher in goitrous thyroid than in normal gland.

**Conclusions :** There are considerable changes in TE contents in the goitrous transformed tissue of thyroid.

Keywords: Thyroid nodular goiter, Intact thyroid, Trace elements, Instrumental neutron activation analysis

Copyright : © 2021 The Authors. Published by Medical Editor and Educational Research Publishers Ltd. This is an open access article under the CC BY-NC-ND license (https://creativecommons.org/licenses/by-nc-nd/4.0/).

#### 1 | INTRODUCTION

o less than 10 % of the world population is affected by goiter detected during the examination and palpation and most of these thyroidal lesions are nodular goiters (NG) (1). However, using ultrasonography NG can be detected in almost 70% of the general population (2). NG is also known as endemic nodular goitre, simple goitre, nodular hyperplasia, nontoxic uninodular goitre or multinodular goiter (3). NG is benign lesions; however, during clinical examination, they can mimic malignant tu-

mors. NG can be hyperfunctioning, hypofunctioning, and normal functioning. Euthyroid NG is defined as a local enlargement of thyroid without accompanying disturbance in thyroid function (3).

For over 20th century, there was the dominant opinion that NG is the simple consequence of iodine deficiency. However, it was found that NG is a frequent disease even in those countries and regions where the population is never exposed to iodine shortage (4). Moreover, it was shown that iodine excess has severe consequences on human health and associated with the presence of thyroidal disfunctions and autoimmunity, NG and diffuse goiter, benign and malignant tumors of gland (5-8). It was also demonstrated that besides the iodine deficiency and excess many other dietary, environmental, and occupational factors are associated with the NG incidence (9-11). Among them a disturbance of evolutionary stable input of many chemical elements in human body after industrial revolution plays a significant role in etiology of thyroidal disorders (12).

Besides iodine involved in thyroid function, other trace elements (TE) have also essential physiological functions such as maintenance and regulation of cell function, gene regulation, activation or inhibition of enzymatic reactions, and regulation of membrane function (13). Essential or toxic (goitrogenic, mutagenic, carcinogenic) properties of TE depend on tissue-specific need or tolerance, respectively (13). Excessive accumulation or an imbalance of the TE may disturb the cell functions and may result in cellular degeneration, death, benign or malignant transformation (13–15).

In our previous studies the complex of in vivo and in vitro nuclear analytical and related methods was developed and used for the investigation of iodine

**Supplementary information** The online version of this article (https://doi.org/10.52845/JMRHS/2021-4-8-5) contains supplementary material, which is available to authorized users.

**Corresponding Author:** Vladimir Zaichick Radionuclide Diagnostics Department Medical Radiological Research Centre Korolyev St.- 4, Obninsk 249036 Kaluga Region, Russia Email: vzaichick@gmail.com and other TE contents in the normal and pathological thyroid (16–22). Iodine level in the normal thyroid was investigated in relation to age, gender and some non-thyroidal diseases (23, 24). After that, variations of TE content with age in the thyroid of males and females were studied and age- and gender-dependence of some TE was observed (25– 41). Furthermore, a significant difference between some TE contents in normal and cancerous thyroid was demonstrated (42–47).

To date, the pathogenesis of NG has to be considered as multifactorial. The present study was performed to clarify the role of some TE in the maintenance of thyroid growth and goitrogenesis. Having this in mind, our aim was to assess the silver (Ag), cobalt (Co), chromium (Cr), iron (Fe), mercury (Hg), rubidium (Rb), antimony (Sb), scandium (Sc), selenium (Se), and zinc (Zn) contents in NG tissue using nondestructive instrumental neutron activation analysis with high resolution spectrometry of long-lived radionuclides (INAA-LLR). A further aim was to compare the levels of these ten TE in the goitrous thyroid with those in intact (normal) gland of apparently healthy persons.

All studies were approved by the Ethical Committees of the Medical Radiological Research Centre (MRRC), Obninsk. All the procedures performed in studies involving human participants were in accordance with the ethical standards of the institutional and/or national research committee and with the 1964 Helsinki declaration and its later amendments, or with comparable ethical standards.

#### 2 | MATERIALS AND METHODS

#### Samples

All patients suffered from NG (n=46, mean age  $M\pm$ SD was  $48\pm12$  years, range 30-64) were hospitalized in the Head and Neck Department of the Medical Radiological Research Centre. Thickneedle puncture biopsy of suspicious nodules of the thyroid was performed for every patient, to permit morphological study of thyroid tissue at these sites and to estimate their TE contents. For all patients the diagnosis has been confirmed by clinical and morphological results obtained during studies of biopsy

and resected materials. Histological conclusion for all thyroidal lesions was the colloid NG.

Normal thyroids for the control group samples were removed at necropsy from 105 deceased (mean age  $44\pm21$  years, range 2-87), who had died suddenly. The majority of deaths were due to trauma. A histological examination in the control group was used to control the age norm conformity, as well as to confirm the absence of micro-nodules and latent cancer.

All tissue samples were divided into two portions using a titanium scalpel (48). One was used for morphological study while the other was intended for TE analysis. After the samples intended for TE analysis were weighed, they were freeze-dried and homogenized (49). The pounded sample weighing about 5-10 mg (for biopsy) and 50 mg (for resected materials) was used for trace element measurement by INAA-LLR. The samples for INAA-LLR were wrapped separately in a high-purity aluminum foil washed with rectified alcohol beforehand and placed in a nitric acid-washed quartz ampoule.

#### Standards and certified reference material

To determine contents of the TE by comparison with a known standard, biological synthetic standards (BSS) prepared from phenol-formaldehyde resins were used (50). In addition to BSS, aliquots of commercial, chemically pure compounds were also used as standards. Ten certified reference material IAEA H-4 (animal muscle) and IAEA HH-1 (human hair) sub-samples weighing about 50 mg were treated and analyzed in the same conditions that thyroid samples to estimate the precision and accuracy of results.

#### Instrumentation and method

A vertical channel of nuclear reactor was applied to determine the content of Ag, Co, Cr, Fe, Hg, Rb, Sb, Sc, Se, and Zn by INAA-LLR. The quartz ampoule with samples of thyroid, standards, and certified reference material was soldered, positioned in a transport aluminum container and exposed to a 24-hour neutron irradiation in a vertical channel of the WWR-c research nuclear reactor (Branch of Karpov Institute, Obninsk) with a neutron flux of  $1.3 \times 10^{13}$  n×cm<sup>-2</sup>×s<sup>-1</sup>. Ten days after irradiation samples were reweighed and repacked. The samples were measured for period from 10 to 30 days after irradiation. The duration of measurements was from 20 min to 10 hours subject to pulse counting rate. The gamma spectrometer included the 100 cm<sup>3</sup> Ge(Li) detector and on-line computerbased MCA system. The spectrometer provided a resolution of 1.9 keV on the <sup>60</sup>Co 1332 keV line. Details of used nuclear reactions, radionuclides, and gamma-energies were presented in our earlier publications concerning the INAA of TE contents in human prostate and scalp hair (51, 52).

#### **Statistical Analysis**

A dedicated computer program for INAA mode optimization was used (53). All thyroid samples were prepared in duplicate, and mean values of TE contents were used in final calculation. Using Microsoft Office Excel, a summary of the statistics, including, arithmetic mean, standard deviation, standard error of mean, minimum and maximum values, median, percentiles with 0.025 and 0.975 levels was calculated for TE contents. The difference in the results between two groups (normal and goitrous thyroid) was evaluated by the parametric Student's t-test and non-parametric Wilcoxon-Mann-Whitney U-test.

#### 3 | RESULTS

Table 1 depicts our data for Ag, Co, Cr, Fe, Hg,Rb, Sb, Sc, Se, and Zn mass fractions in ten subsamples of IAEA H-4 (animal muscle) and IAEA HH-1 (human hair) certified reference material and the certified values of this material.

Table 1: INAA-LLR data of trace element contents in certified reference material IAEA H-4 (animal muscle) and IAEA HH-1 (human hair) compared to certified values ((mg/kg, dry mass basis)

Element	IAEA H-4	This work	IAEA HH-1	This work
	animal muscle	results	human hair	results
	95% confidence interval	M±SD	95% confidence interval	M±SD
Ag	-	0.033±0.008	0.19 <sup>b</sup>	0.18±0.05
Co	0.0027 <sup>b</sup>	0.0034±0.0008	5.97±0.42 <sup>a</sup>	5.4±1.1
Cr	$0.06^{b}$	0.071±0.010	0.27 <sup>b</sup>	=0.3
Fe	49.1±6.5 <sup>a</sup>	47.0±1.0	23.7±3.1ª	25.1±4.3
Hg	0.014 <sup>b</sup>	0.015±0.004	1.70±0.09 <sup>a</sup>	1.54±0.14
Rb	18.7±3.5 <sup>a</sup>	23.7±3.7	0.94 <sup>b</sup>	0.89±0.17
Sb	0.0056 <sup>b</sup>	0.0061±0.0021	0.031 <sup>b</sup>	0.033±0.009
Sc	0.0059 <sup>b</sup>	0.0015±0.0009	-	-
Se	$0.28{\pm}0.08^{a}$	0.281±0.014	0.35±0.02 <sup>a</sup>	0.37±0.08
Zn	86.3±11.5 <sup>a</sup>	91±2	174±9 <sup>a</sup>	173±17

1407

M – arithmetical mean, SD – standard deviation, a – certified values. b – information values.

resents certain statistical parameters (arithmetic mean, standard deviation, standard error of mean, minimal and maximal values, median, percentiles with 0.025 and 0.975 levels) of the Ag, Co, Cr, Fe, Hg,Rb, Sb, Sc, Se, and Zn mass fraction in normal and goitrous thyroid tissue.

Table 2: Some statistical parameters of Ag, Co, Cr, Fe, Hg, Rb, Sb, Sc, Se, and Zn mass fraction (mg/kg, dry mass basis) in normal thyroid and colloid nodular goiter

Tissue	Element	Mean	SD	SEM	Min	Max	Median	P 0.025	P 0.975
Normal	Ag	0.0151	0.0140	0.0016	0.0012	0.0800	0.0121	0.0017	0.0454
n=105	Co	0.0399	0.0271	0.0030	0.0046	0.140	0.0327	0.0134	0.124
	Cr	0.539	0.272	0.032	0.130	1.30	0.477	0.158	1.08
	Fe	225	100	11	51.0	512	217	67.4	456
	Hg	0.0421	0.0358	0.0041	0.0065	0.180	0.0304	0.0091	0.150
	Rb	7.37	4.10	0.44	1.11	29.4	6.49	2.60	16.7
	Sb	0.111	0.072	0.008	0.0047	0.308	0.103	0.0117	0.280
	Sc	0.0046	0.0038	0.0008	0.0002	0.0143	0.0042	0.00035	0.0131
	Se	2.32	1.29	0.14	0.439	5.80	2.01	0.775	5.65
	Zn	97.8	42.3	4.5	8.10	221	91.7	34.8	186
Goiter	Ag	0.226	0.236	0.042	0.0020	0.874	0.160	0.0020	0.849
n=46	Co	0.0627	0.0287	0.0050	0.0150	0.147	0.0623	0.0215	0.128
	Cr	0.849	0.834	0.150	0.135	3.65	0.540	0.142	2.88
	Fe	340	332	52	62.0	1350	197	68.8	1344
	Hg	0.987	0.726	0.124	0.0817	3.01	0.920	0.0968	2.36
	Rb	8.85	4.18	0.64	1.00	22.1	8.50	2.53	16.6
	Sb	0.146	0.121	0.021	0.0102	0.425	0.103	0.0128	0.419
	Sc	0.0130	0.0201	0.0040	0.0002	0.0910	0.0058	0.0002	0.0701
	Se	3.09	2.59	0.44	0.994	12.6	2.37	1.16	12.1
	Zn	121	53.4	8.2	47.0	278	109	49.1	269

M – arithmetic mean, SD – standard deviation, SEM - standard error of mean, Min - minimum value, Max - maximum value, P 0.025 - percentile with 0.025 level, P 0.975 – percentile with 0.975 level.

The comparison of our results with published data for Ag, Co, Cr, Fe, Hg, Rb, Sb, Sc, Se, and Zn mass fraction in normal and goitrous thyroid (54-74) is shown in Table 3.

The ratios of means and the difference between mean values of Ag, Co, Cr, Fe, Hg, Rb, Sb, Sc, Se, and Zn mass fractions in normal and goitrous thyroid are presented in Table 4.

#### 4 DISCUSSION

#### Precision and accuracy of results

Good agreement of the Ag, Co, Cr, Fe, Hg, Rb, Sb, Sc, Se, and Zn contents analyzed by INAA-LLR with the certified data of CRM IAEA H-4 and IAEA HH-1 (Table 1) indicates an acceptable accuracy of the results obtained in the study of TE of the thyroid presented in Tables 2-4.

The mean values and all selected statistical parameters were calculated for ten TE (Ag, Co, Cr, Fe, Hg, Rb, Sb, Sc, Se, and Zn) mass fractions (Table 2). The mass fraction of Ag, Co, Cr, Fe, Hg, Rb, Sb, Sc, Se, and Zn were measured in all, or a major portion of normal and goitrous tissue samples.

#### Comparison with published data

In general, values obtained for Cr, Fe, Hg, Rb, Sb, Sc, Se, and Zn contents in the normal human thyroid (Table 3) agree well with median of mean values reported by other researches (54-66). The obtained means for Ag and Co were almost one order of magnitude lower median of previously reported means but inside the range of means (Table 3). A number of values for TE mass fractions were not expressed on a dry mass basis by the authors of the cited references. However, we calculated these values using published data for water (75%) (75) and ash (4.16% on dry mass basis) (76) contents in thyroid of adults.

Table 3. Median, minimum and maximum value of means Ag, Co, Cr, Fe, Hg, Rb, Sb, Sc, Se, and Zn contents in normal and goitrous thyroid according to data from the literature in comparison with our results (mg/kg, dry mass basis)

			Published data [Refer		
Tissue	El		This work		
		Median	Minimum	Maximum	
		of means	of means	ofmeans	
		(n)*	M or M±SD, (n)**	M or M±SD, (n)**	M±SD
Normal	Ag	0.21 (12)	0.000784 (16) [54]	1.20±1.24 (105) [55]	
	Co	0.306 (25)	0.016 (66) [56]	70.4±40.8 (14) [57]	0.040±0.02
	Cr	0.69 (17)	0.088 (83) [58]	24.8±2.4 (4) [59]	0.54±0.27
	Fe	252 (21)	56 (120) [60]	3360 (25) [61]	225±100
	Hg	0.08 (13)	0.0008±0.0002 (10) [62]	396±40 (4) [59]	0.042±0.03
	Rb	7.8 (9)	=0.85 (29) [62]	294±191 (14) [57]	7.37±4.10
	Sb	0.15(10)	0.040±0.003 (-) [63]	= 12.4(-) [64]	0.111±0.07
	Sc	0.009(4)	0.0018±0.0003 (17) [56]	0.014±0.005 (10) [62]	0.005±0.00
	Se	2.32 (21)	0.436 (40) [65]	756±680 (14) [57]	2.32±1.29
	Zn	110 (56)	2.1 (-) [66]	820±204 (14) [57]	97.8±42.3
Goiter	Ag	0.210 (4)	0.098±0.042 (19) [67]	2.56 (167) [68]	
	Co	0.67(12)	0.110±0.003 (64) [69]	62.8±22.4 (11) [57]	0.063±0.02
	Cr	3.66 (5)	0.72 (51) [70]	25.2 (25) [61]	0.849±0.83
	Fe	390 (5)	128±52 (13) [71]	4848±3056 (11) [57	
	Hg	-	-		0.987±0.72
	Rb	7.5 (2)	7,0 (10) [65]	864±148 (11) [57	8.85±4.18
	Sb	0.63 (1)	0.15 (19) [72]	1.10 (19) [72]	0.146±0.12
	Sc	-		. /	0.013±0.02
	Se	2.60 (8)	0.248 (41) [56]	174±116 (11) [57]	
	Zn	146 (25)	22.4 (130) [73]	1236±560 (2) [74]	1
			MEERP I		1 4 0 0
	1408				

El - element, M –arithmetic mean, SD – standard deviation,  $(n)^*$  – number of all references,  $(n)^{**}$  – number of samples.

Table 4. Differences between mean values  $(M\pm SEM)$  of Ag, Co, Cr, Fe, Hg, Rb, Sb, Sc, Se, and Zn mass fraction (mg/kg, dry mass basis) in normal thyroid and colloid nodular goiter

Element		Ratio			
	Norm	Goiter Student's t-test		U-test	Goiter
	n=105	n=46	$p \le$	р	to Norm
Ag	0.0151±0.0016	0.226±0.042	0.0000185	=0.01	15.0
Co	$0.0399 \pm 0.0030$	$0.0627 \pm 0.0050$	0.00023	=0.01	1.57
Cr	0.539±0.032	0.849±0.150	0.0507	=0.05	1.58
Fe	225±11	340±52	0.0373	=0.01	1.51
Hg	$0.0421 \pm 0.0041$	0.987±0.124	0.000000001	=0.01	23.4
Rb	7.37±0.44	8.85±0.64	0.0617	>0.05	1.20
Sb	0.111±0.008	0.146±0.021	0.119	>0.05	1.32
Sc	$0.0046 \pm 0.0008$	$0.0130 {\pm} 0.0040$	0.0377	=0.01	2.83
Se	2.32±0.14	3.09±0.44	0.105	>0.05	1.33
Zn	97.8±4.5	121±8.2	0.0155	=0.01	1.24

M – arithmetic mean, SEM – standard error of mean, Statistically significant values are in **bold**.

Data cited in Table 3 for normal thyroid also includes samples obtained from patients who died from different non-endocrine diseases. In our previous study it was shown that some non-endocrine diseases can effect on TE contents in thyroid (24). Moreover, in many studies the "normal" thyroid means a visually non-affected tissue adjacent to benign or malignant thyroidal nodules. However, there are no data on a comparison between the TE contents in such kind of samples and those in thyroid of healthy persons, which permits to confirm their identity.

Our results for goitrous tissues were comparable with published data for Ag, Fe, Rb, Se, and Zn contents (Table 3). The obtained means for Co, Cr, and Sb were approximately 10.6, 4.3, and 4.3, respectively, times lower median of previously reported means, herewith, mean for Cr was inside the range of these means, but mean for Co and Sb were outside (Table 3). No published data referring Hg and Sc contents of goitrous thyroid tissue were found.

The range of means of Ag, Co, Cr, Fe, Hg, Rb, Sb, Sc, Se, and Zn level reported in the literature for normal and for goitrous thyroid vary widely (Table 3). This can be explained by a dependence of TE content on many factors, including "normality" of thyroid samples (see above), the region of the thyroid, from which the sample was taken, age, gender, ethnicity, mass of the gland, and the goiter stage. Not all these factors were strictly controlled in cited studies. However, in our opinion, the leading causes of interobserver variability can be attributed to the accuracy of the analytical techniques, sample preparation methods, and inability of taking uniform samples from the affected tissues. It was insufficient quality control of results in these studies. In many scientific reports, tissue samples were ashed or dried at high temperature for many hours. In other cases, thyroid samples were treated with solvents (distilled water, ethanol, formalin etc). There is evidence that during ashing, drying and digestion at high temperature some quantities of certain TE are lost as a result of this treatment. That concerns not only such volatile halogen as Br, but also other TE investigated in the study (77–79).

# Effect of goitrous transformation on ChE contents

From Table 4, it is observed that in goitrous tissues the mass fractions of all TE investigated Ag, Co, Cr, Fe, Hg, Rb, Sb, Sc, Se, and Zn are 15.0, 1.57, 1.58, 1.51, 23.4, 1.20, 1.32, 2.83, 1.33, and 1.24 times, respectively, higher than in normal tissues of the thyroid. However, the changes for Ag, Co, Cr, Fe, Hg, Sc, and Zn are just statistically significant. Thus, if we accept the TE contents in thyroid glands in the control group as a norm, we have to conclude that with a goitrous transformation the Ag, Co, Cr, Fe, Hg, Sc, and Zn contents in thyroid tissue significantly changed.

# Role of ChE in goitrous transformation of the thyroid

Characteristically, elevated or reduced levels of TE observed in goitrous tissues are discussed in terms of their potential role in the initiation and promotion of goiter. In other words, using the low or high levels of the TE found in goitrous tissues, researchers try to determine the goitrogenic role of the deficiency or excess of each TE in investigated organ. In our opinion, abnormal levels of many TE in NG could be and cause, and also effect of goitrous transformation. From the results of such kind studies, it is not always possible to decide whether the measured decrease or increase in TE level in pathologically altered tissue

is the reason for alterations or vice versa..

#### Silver

Ag is a TE with no recognized trace metal value in the human body (80). Ag in metal form and inorganic Ag compounds ionize in the presence of water, body fluids or tissue exudates. The silver ion Ag<sup>+</sup> is biologically active and readily interacts with proteins, amino acid residues, free anions and receptors on mammalian and eukaryotic cell membranes (81). Besides such the adverse effects of chronic exposure to Ag as a permanent bluish-gray discoloration of the skin (argyria) or eyes (argyrosis), exposure to soluble Ag compounds may produce other toxic effects, including liver and kidney damage, irritation of the eyes, skin, respiratory, and intestinal tract, and changes in blood cells (82). More detailed knowledge of the Ag toxicity can lead to a better understanding of the impact on human health, including thyroid function.

#### Cobalt

Health effects of high Co occupational, environmental, dietary and medical exposure are characterized by a complex clinical syndrome, mainly including neurological, cardiovascular and endocrine deficits, including hypothyroidism and goiter (83, 84). Co is genotoxic and carcinogenic, mainly caused by oxidative DNA damage by reactive oxygen species, perhaps combined with inhibition of DNA repair (85). In our previous studies it was found a significant agerelated increase of Co content in female thyroid (25). Therefore, a goitrogenic and, probably, carcinogenic effect of excessive Co level in the thyroid of old females was assumed. Elevated level of Co in NG tissues, observed in the present study, supports this conclusion.

#### Chromium

Cr-compounds are cytotoxic, genotoxic, and carcinogenic in nature. Some Cr forms, including hexavalent chromium ( $Cr^{6+}$ ), are toxicants known for their carcinogenic effect in humans. They have been classified as certain or probable carcinogens by the International Agency for Research on Cancer (86). The lung cancer risk is prevalent in pigment chromate handlers, ferrochromium production workers, stainless steel welders, and chrome-platers (87). Except in Cr-related industries and associated environments, Cr intoxication from environmental exposure is not common. However, it was found, that drinking water supplies in many geographic areas contain chromium in the +3 and +6 oxidation states. Exposure of animals to Cr<sup>6+</sup>in drinking water induced tumors in the mouse small intestine (88). Many other animal experiments and in vitro studies demonstrate also that Cr can induce oxidative stress and exert cytotoxic effects (89). Besides reactive oxygen species (ROS) generation, oxidative stress, and cytotoxic effects of Cr exposure, a variety of other changes like DNA damage, increased formation of DNA adducts and DNA-protein cross-links, DNA strand breaks, chromosomal aberrations and instability, disruption of mitotic cell division, chromosomal aberration, premature cell division, S or G2/M cell cycle phase arrest, and carcinogenesis also occur in humans or experimental test systems (87).

#### Iron

It is well known that Fe as TE is involved in many very important functions and biochemical reactions of human body. Fe metabolism is therefore very carefully regulated at both a systemic and cellular level (90, 91). Under the impact of age and multiple environmental factors the Fe metabolism may become dysregulated with attendant accumulation of this metal excess in tissues and organs, including thyroid (25, 26, 29–35). Most experimental and epidemiological data support the hypothesis that Fe overload is a risk factor for benign and malignant tumors (92). This goitrogenic and oncogenic effect could be explained by an overproduction of ROS and free radicals (93).

#### Mercury

Hg is one of the most dangerous environmental pollutants (94). The growing use of this metal in diverse areas of industry has resulted in a significant increase of environment contamination and episodes of human intoxication. Hg damages the central nervous system and has irreparable effects on the kidneys (95). Hg may also harm a developing fetus and decrease fertility in men and women (96). Besides these effects, Hg has been classified as certain or probable carcinogen by the International Agency for Research on Cancer (86). For example, in Hg

polluted area thyroid cancer incidence was almost 2 times higher than in adjacent control areas (97).

Negative effects of Hg are due to the interference of this metal in cellular signaling pathways and protein synthesis during the period of development. Since it bonds chemically with the sulfur hydride groups of proteins, it causes damage to the cell membrane and decreases the amount of RNA (98). Moreover, it was shown that Hg may be involved in four main processes that lead to genotoxicity: generation of free radicals and oxidative stress, action on microtubules, influence on DNA repair mechanisms and direct interaction with DNA molecules (99).

#### Scandium

Sc is a rare earth element. Information about its physiological role is very limited. However, toxic effects concerning Sc propensity to displace calcium in many biochemical events and its carcinogenic potential have been reported (100, 101).

#### Zinc

Zn as a trace metal plays an important role in normal and pathophysiology. This TE is a constituent of more than 3000 proteins and is a cofactor for over 300 enzymes (102). Zn is an essential mediator of cell proliferation and differentiation through the regulation of DNA synthesis and mitosis. Zn also affects DNA repair pathways by regulating multiple intracellular signaling pathways and altering proteins involved in DNA maintenance (103). This metal also maintenance the balance of a cellular redox (104). Thus, Zn is important cofactors in diverse cellular processes, but its high concentrations are toxic to the cells. The elevated level of Zn mass fractions in thyroid tissue may contribute to harmful effects on the gland. There are good reasons for such speculations since. experimental and epidemiological data support the hypothesis that Zn overload is a risk factor for benign and malignant tumors (103, 105-107).

Our findings show that mass fraction of Ag, Co, Cr, Fe, Hg, Sc, and Zn are significantly higher in NG as compared to normal thyroid tissues (Tables 4). Thus, it is plausible to assume that levels of these TE in thyroid tissue can be used as NG markers. However, this subjects needs in additional studies.

#### 5 | LIMITATIONS

This study has several limitations. Firstly, analytical techniques employed in this study measure only ten TE (Ag, Co, Cr, Fe, Hg, Rb, Sb, Sc, Se, and Zn) mass fractions. Future studies should be directed toward using other analytical methods which will extend the list of chemical elements investigated in normal and goitrous thyroid. Secondly, the sample size of NG group was relatively small. It was not allow us to carry out the investigations of TE contents in NG group using differentials like gender, histological types of colloid NG, stage of disease, and dietary habits of healthy persons and patients with NG. Lastly, generalization of our results may be limited to Russian population. Despite these limitations, this study provides evidence on goiter-specific tissue Ag, Co, Cr, Fe, Hg, Sc, and Zn level alteration and shows the necessity to TE research of NG.

#### 6 | CONCLUSION

In this work, TE analysis was carried out in the tissue samples of normal thyroid and NG of thyroid using INAA-LLR. It was shown that INAA-LLR is an adequate analytical tool for the non-destructive determination of Ag, Co, Cr, Fe, Hg, Rb, Sb, Sc, Se, and Zn content in the tissue samples of human thyroid, including needle-biopsy cores. It was observed that in goitrous thyroid content of Ag, Co, Cr, Fe, Hg, Sc, and Zn were significantly higher than in normal tissues. In our opinion, the increase in levels of Ag, Co, Cr, Fe, Hg, Sc, and Zn in goitrous tissue might demonstrate an involvement of these TE in etiology and pathogenesis of thyroid goiter. It was supposed that elevated levels of Ag, Co, Cr, Fe, Hg, Sc, and Zn in thyroid tissue can be used as NG markers.

#### **Declaration of Conflicting Interests**

The author has not declared any conflict of interests.

#### 7 | FUNDING

The authors received no financial support for this study and for publication of this article.

#### ACKNOWLEDGEMENTS

The authors are extremely grateful to Profs. Vtyurin BM and Medvedev VS, Medical Radiological Research Center, Obninsk, as well as to Dr. Choporov Yu, Head of the Forensic Medicine Department of City Hospital, Obninsk, for supplying thyroid samples.

#### REFERENCES

- Carlé A, Krejbjerg A, Laurberg P. Epidemiology of nodular goitre. Influence of iodine intake. Best Pract Res Clin Endocrinol Metab. 2014;28(4):465–79.
- 2. Kant R, Davis A, Verma V. Thyroid nodules: Advances in evaluation and management. Am Fam Physician. 2020;102(5):298–304.
- 3. Hoang VT, Trinh CT. A Review of the Pathology, Diagnosis and Management of Colloid Goitre. Eur Endocrinol. 2020;16(2):131–135.
- Derwahl M, Studer H. Multinodular goitre: 'much more to it than simply iodine deficiency. Baillieres Best Pract Res Clin Endocrinol Metab. 2000;14(4):577–600.
- 5. Zaichick V. Iodine excess and thyroid cancer. J Trace Elem Exp Med. 1998;11(4):508–509.
- Zaichick V, Iljina T. Dietary iodine supplementation effect on the rat thyroid 1311 blastomogenic action. Die Bedentung der Mengenund Spurenelemente 18 Arbeitstangung Jena: Friedrich-Schiller-Universitat. 1998;p. 294–306.
- Kim S, Kwon YS, Kim JY, Hong KH, Park YK. Association between Iodine Nutrition Status and Thyroid Disease-Related Hormone in Korean Adults: Korean National Health and Nutrition Examination Survey VI. Nutrients. 2013;11(11):2757–2757.
- 8. Vargas-Uricoechea □, Pinzón-Fernández MV, Bastidas-Sánchez BE, Jojoa-Tobar E,

Ramírez-Bejarano LE, Murillo-Palacios J. Iodine Status in the Colombian Population and the Impact of Universal Salt Iodization: A Double-Edged Sword? J Nutr Metab. 2019;p. 6239243–6239243.

- Stojsavljević A, Rovčanin B, Krstić D, Jagodić J, Borković-Mitić S, Paunović I, et al. Cadmium as main endocrine disruptor in papillary thyroid carcinoma and the significance of Cd/Se ratio for thyroid tissue pathophysiology. J Trace Elem Med Biol. 2019;55:190–195.
- Fahim YA, Sharaf NE, Hasani IW, Ragab EA, Abdelhakim HK. Assessment of Thyroid Function and Oxidative Stress State in Foundry Workers Exposed to Lead. J Health Pollut. 2020;10(27).
- Liu M, Song J, Jiang Y, Lin Y, Peng J, Liang H, et al. A case-control study on the association of mineral elements exposure and thyroid tumor and goiter. Ecotoxicol Environ Saf. 2021;208:111615–111615.
- Zaichick V. Medical elementology as a new scientific discipline. J Radioanal Nucl Chem. 2006;269:303–309.
- 13. Moncayo R, Moncayo H. A post-publication analysis of the idealized upper reference value of 2.5 mIU/L for TSH: Time to support the thyroid axis with magnesium and iron especially in the setting of reproduction medicine. BBA Clin. 2017;7:115–119.
- Beyersmann D, Hartwig A. Carcinogenic metal compounds: recent insight into molecular and cellular mechanisms. Arch Toxicol. 2008;82(8):493–512.
- 15. Martinez-Zamudio R, Ha HC. Environmental epigenetics in metal exposure. Epigenetics. 2011;6(7):820–827.
- Zaĭchik VE, Yus R, Melnik AD, Cherkashin VI. Neutron-activation analysis in the study of the behavior of iodine in the organism. Med Radiol (Mosk). 1970;15(1):33–36.

- Zaĭchik VE, Matveenko EG, Vtiurin BM, Medvedev VS. Intrathyroid iodine in the diagnosis of thyroid cancer. Vopr Onkol. 1982;28(3):18–24.
- Zaichick V, Tsyb AF, Vtyurin BM. Trace elements and thyroid cancer. Analyst. 1995;120(3):817–821.
- Vye Z, Yuya C. Determination of the natural level of human intra-thyroid iodine by instrumental neutron activation analysis. J Radioanal Nucl Chem. 1996;207(1):153–161.
- 20. Zaichick V. In vivo and in vitro application of energy-dispersive XRF in clinical investigations: experience and the future. J Trace Elem Exp Med. 1998;11(4):509–510.
- Zaichick V, Zaichick S. Energy-dispersive Xray fluorescence of iodine in thyroid puncture biopsy specimens. J Trace Microprobe Tech. 1999;17(2):219–232.
- 22. Zaichick V. Relevance of, and potentiality for in vivo intrathyroidal iodine determination. Ann N Y Acad Sci. 2000;904:630–632.
- 23. Zaichick V, Zaichick S. Normal human intrathyroidal iodine. Sci Total Environ. 1997;206(1):39–56.
- 24. Zaichick V. Human intrathyroidal iodine in health and non-thyroidal disease. New aspects of trace element research. 1999;p. 114–119.
- 25. Zaichick V, Zaichick S. Age-related changes of some trace element contents in intact thyroid of females investigated by energy dispersive X-ray fluorescent analysis. Trends Geriatr Healthc;2017(1):31–38.
- Zaichick V, Zaichick S. Age-related changes of some trace element contents in intact thyroid of males investigated by energy dispersive Xray fluorescent analysis. MOJ Gerontol Ger. 2017;1(5):28–28.
- Zaichick V, Zaichick S. Age-related changes of Br, Ca, Cl, I, K, Mg, Mn, and Na contents in intact thyroid of females investigated by neutron activation analysis. Curr Updates Aging. 2017;1:5–6.

- Zaichick V, Zaichick S. Age-related changes of Br, Ca, Cl, I, K, Mg, Mn, and Na contents in intact thyroid of males investigated by neutron activation analysis. J Aging Age Relat Dis. 2017;1(1):1002–1002.
- 29. Zaichick V, Zaichick S, Age, Ag, Co, Cr, et al. Se, and Zn contents in intact thyroid of females investigated by neutron activation analysis. J Gerontol Geriatr Med. 2017;3:15–15.
- Zaichick V, Zaichick S, Age, Ag, Co, Cr, et al. Se, and Zn contents in intact thyroid of males investigated by neutron activation analysis. Curr Trends Biomedical Eng Biosci. 2017;4(4):555644–555644.
- 31. Zaichick V, Zaichick S; 2018.
- Zaichick V, Zaichick S. Neutron activation and X-ray fluorescent analysis in study of association between age and chemical element contents in thyroid of males. Op Acc J Bio Eng Bio Sci. 2018;2(4):202–212.
- Zaichick V, Zaichick S. Variation with age of chemical element contents in females' thyroids investigated by neutron activation analysis and inductively coupled plasma atomic emission spectrometry. J Biochem Analyt Stud. 2018;3(1):1–10.
- Zaichick V, Zaichick S. Association between Age and Twenty Chemical Element Contents in Intact Thyroid of Males. SM Gerontol Geriatr Res. 2018;2(1):1014–1014.
- Zaichick V, Zaichick S. Associations between age and 50 trace element contents and relationships in intact thyroid of males. Aging Clin Exp Res. 2018;30(9):1059–1070.
- Zaichick V, Zaichick S. Possible role of inadequate quantities of intra-thyroidal bromine, rubidium and zinc in the etiology of female subclinical hypothyroidism. EC Gynaecology. 2018;7(3):107–115.

- Zaichick V, Zaichick S. Possible role of inadequate quantities of intra-thyroidal bromine, calcium and magnesium in the etiology of female subclinical hypothyroidism. Int Gyn and Women's. Health. 2018;1(3).
- Zaichick V, Zaichick S. Possible role of inadequate quantities of intra-thyroidal cobalt, rubidium and zinc in the etiology of female subclinical hypothyroidism. Womens Health Sci J. 2018;2(1):108–108.
- 39. Zaichick V, Zaichick S. Association between female subclinical hypothyroidism and inadequate quantities of some intra-thyroidal chemical elements investigated by X-ray fluorescence and neutron activation analysis. Gynaecology and Perinatology. 2018;2(4):340–355.
- 40. Zaichick V, Zaichick S. Investigation of association between the high risk of female subclinical hypothyroidism and inadequate quantities of twenty intra-thyroidal chemical elements. Clin Res: Gynecol Obstet. 2018;1(1):1–18.
- 41. Zaichick V, Zaichick S. Investigation of association between the high risk of female subclinical hypothyroidism and inadequate quantities of intra-thyroidal trace elements using neutron activation and inductively coupled plasma mass spectrometry. Acta Scientific Medical Sciences. 2018;2(9):23–37.
- 42. Zaichick V, Zaichick S. Trace element contents in thyroid cancer investigated by energy dispersive X-ray fluorescent analysis. American Journal of Cancer Research and Reviews. 2018;2:5–5.
- Zaichick V, Zaichick S. Trace element contents in thyroid cancer investigated by instrumental neutron activation analysis. J Oncol Res. 2018;2(1):1–13.
- 44. Zaichick V, Zaichick S. Variation in selected chemical element contents associated with malignant tumors of human thyroid gland. Cancer Studies. 2018;2(1):2–2.
- 45. Zaichick V, Zaichick S. Twenty chemical element contents in normal and cancerous thyroid. Int J Hematol Blo Dis. 2018;3(2):1–13.

- Zaichick V, Zaichick S. Levels of chemical element contents in thyroid as potential biomarkers for cancer diagnosis (a preliminary study). J Cancer Metastasis Treat. 2018;4:60–60.
- 47. Zaichick V, Zaichick S. Fifty trace element contents in normal and cancerous thyroid. Acta Scientific Cancer Biology. 2018;2(8):21–38.
- Zaichick V, Zaichick S. Instrumental effect on the contamination of biomedical samples in the course of sampling. The Journal of Analytical Chemistry. 1996;51(12):1200–1205.
- Zaichick V, Yuv T. A simple device for biosample lyophilic drying. Lab Delo. 1978;2:109–110.
- 50. Zaichick V. Applications of synthetic reference materials in the Medical Radiological Research Centre. Fresenius J Anal Chem. 1995;352:219– 223.
- Zaichick S, Zaichick V, Ag, Co, Cr, Fe, et al. Se, and Zn contents in intact human prostate investigated by neutron activation analysis. J Appl Radiat Isot. 2011;69:827–833.
- 52. Zaichick S, Zaichick V. The effect of age and gender on 37 chemical element contents in scalp hair of healthy humans. Biol Trace Elem Res. 2010;134(1):41–54.
- 53. Korelo AM, Zaichick V. Software to optimize the multielement INAA of medical and environmental samples. In: Activation Analysis in Environment Protection; 1993. p. 326–332.
- 54. Zhu H, Wang N, Zhang Y, Wu Q, Chen R, Gao J, et al. Element contents in organs and tissues of Chinese adult men. Health Phys. 2010;98(1):61–73.
- 55. Vlasova ZA. Trace element dynamics in thyroid in connection with age and atherosclerosis. Proceedings of Leningradskii Institute of Medical Doctor Postgraduate Education. 1969;80:135–144.

- 56. Stojsavljević A, Rovčanin B, Krstić D, Borković-Mitić S, Paunović I, Diklić A, et al. Risk assessment of toxic and essential trace metals on the thyroid health at the tissue level: The significance of lead and selenium for colloid goiter disease. Expo Health. 2019;.
- 57. Salimi J, Moosavi K, Vatankhah S, Yaghoobi A. Investigation of heavy trace elements in neoplastic and non-neoplastic human thyroid tissue: A study by proton induced X-ray emissions. Int J Radiat Res. 2004;1(4):211–216.
- Reitblat MA, Kropachyev AM. Some trace elements in thyroid of the Perm Pricam'ya residents. Proceedings of Perm Medical Institute. 1967;78:157–164.
- 59. Reddy SB, Charles MJ, Kumar MR, Reddy BS, Ch A, Raju G, et al. Trace elemental analysis of adenoma and carcinoma thyroid by PIXE method. Nuclear Instruments and Methods in Physics Research Section B: Beam Interactions with. Materials and Atoms. 2002;196(3-4):333–339.
- 60. Ataullachanov IA. Age changes in the content of manganese, cobalt, copper, zinc and iron in the endocrine glands of women. Probl Endocrinol (Mosk). 1969;15(2):98–102.
- 61. Kamenev VF. About trace element contents in thyroid of adults. In: Trace Elements in Agriculture and Medicine; 1963. p. 12–16.
- 62. Boulyga SF, Zhuk IV, Lomonosova EM, Kanash NV, Bazhanova NN. Determination of microelements in thyroids of the inhabitants of Belarus by neutron activation analysis using the k0-method. J Radioanal Nucl Chem. 1997;222(1-2):11–14.
- 63. Boulyga SF, Becker JS, Malenchenko AF, Dietze HJ. Application of ICP-MS for multielement analysis in small sample amounts of pathological thyroid tissue. Microchimica Acta. 2000;134(3-4):215–222.
- 64. Zakutinsky DK, Yud P, Selivanova LN; 1962.

- 65. Kvicala J, Havelka J, Nemec J, Zeman V. Selenium and rubidium changes in subjects with pathologically altered thyroid. Biol Trace Elem Res. 1992;32:253–258.
- 66. Kortev AI, Donthov GI, Lyascheva AP. Bioelements and a human pathology. Sverdlovsk, Russia; 1972.
- Predtechenskaya VC. Nucleic acids and trace elements in thyroid pathology. Proceedings of the Voronezh Medical Faculty. 1975;94:85– 87.
- 68. Kovalev MM. Trace element contents in normal and goitrous glands. Vrach Delo. 1960;12:107–111.
- 69. Błazewicz A, Dolliver W, Sivsammye S, Deol A, Randhawa R, Orlicz-Szczesna G, et al. Determination of cadmium, cobalt, copper, iron, manganese, and zinc in thyroid glands of patients with diagnosed nodular goitre using ion chromatography. J Chromatogr B Analyt Technol Biomed Life Sci. 2010;878(1):34–38.
- Pavlyuchenkova EG, Sorokina AN. Spectrographic determination of chromium in thyroid. In: Trace Elements in Agriculture and Medicine; 1963. p. 23–27.
- Kaya G, Avci H, Akdeniz I, Yaman M. Determination of trace and minor metals in benign and malign human thyroid tissues. Asian J Chem. 2009;21(7):5718–5726.
- Boulyga SF, Petri H, Zhuk IV, Kanash NV, Malenchenko AF. Neutron-activation analysis of trace elements in thyroids. J Radioanal Nucl Chem. 1999;242(2):335–340.
- 73. Stojsavljević A, Rovčanin B, Krstić D, Borković-Mitić S, Paunović I, Kodranov I, et al. Evaluation of trace metals in thyroid tissues: Comparative analysis with benign and malignant thyroid diseases. Ecotoxicol Environ Saf. 2019;183:109479–109479.

- 74. Zagrodzki P, Nicol F, Arthur JR, Słowiaczek M, Walas S, Mrowiec H, et al. Selenoenzymes, laboratory parameters, and trace elements in different types of thyroid tumor. Biol Trace Elem Res. 2010;134(1):25–40.
- 75. Katoh Y, Sato T, Yamamoto Y. Determination of multielement concentrations in normal human organs from the Japanese. Biol Trace Elem Res. 2002;90(1-3):57–70.
- Schroeder HA, Tipton IH, Nason AP. Trace metals in man: strontium and barium. J Chron Dis. 1972;25(9):491–517.
- 77. Zaichick V;.
- Zaichick V, Zaichick S. A search for losses of chemical elements during freeze-drying of biological materials. J Radioanal Nucl Chem. 1997;218(2):249–253.
- Zaichick V. Losses of chemical elements in biological samples under the dry aching process. Trace Elements in Medicine. 2004;5(3):17–22.
- Lansdown AB. Critical observations on the neurotoxicity of silver. Crit Rev Toxicol. 2007;37(3):237–250.
- Lansdown AB. Silver in health care: antimicrobial effects and safety in use. Curr Probl Dermatol. 2006;33:17–34.
- 82. Drake PL, Hazelwood KJ. Exposure-related health effects of silver and silver compounds: a review. Ann Occup Hyg. 2005;49(7):575–585.
- 83. Prescott E, Netterstrøm B, Faber J, Hegedüs L, Suadicani P, Christensen JM. Effect of occupational exposure to cobalt blue dyes on the thyroid volume and function of female plate painters. Scand J Work Environ Health. 1992;18(2):101–104.
- Yu R. Cobalt toxicity, an overlooked cause of hypothyroidism. J Endocrinol Thyroid Res. 2017;1(3):1–4.
- 85. Simonsen LO, Harbak H, Bennekou P. Cobalt metabolism and toxicology–a brief update. Sci Total Environ. 2012;432:210–215.

- 86. Järup L. Hazards of heavy metal contamination. Br Med Bull. 2003;68:167–182.
- Nigam A, Priya S, Bajpai P, Kumar S. Cytogenomics of hexavalent chromium (Cr 6+) exposed cells: a comprehensive review. Indian J Med Res. 2014;139(3):349–370.
- 88. Zhitkovich A. Chromium in drinking water: sources, metabolism, and cancer risks. Chem Res Toxicol. 2011;24(10):1617–1629.
- 89. Ding SZ, Yang YX, Li XL, Michelli-Rivera A, Han SY, Wang L, et al. Epithelialmesenchymal transition during oncogenic transformation induced by hexavalent chromium involves reactive oxygen speciesdependent mechanism in lung epithelial cells. Toxicol Appl Pharmacol. 2013;269(1):61–71.
- Manz DH, Blanchette NI, Paul BT, Torti FM, Torti SV. Iron and cancer: recent insights. Ann N Y Acad Sci. 2016;1368(1):149–161.
- Torti SV, Manz DH, Paul BT, Blanchette-Farra N, Torti FM. Iron and Cancer. Annu Rev Nutr. 2018;38:97–125.
- 92. Selby JV, Friedman GD. Epidemiologic evidence of an association between body iron stores and risk of cancer. Int J Cancer. 1988;41:677–682.
- 93. Meneghini R. Iron homeostasis, oxidative stress, and DNA damage. Free Radic Biol Med. 1997;23:783–792.
- 94. Clarkson TW, Magos L. The toxicology of mercury and its chemical compounds. Crit Rev Toxicol. 2006;36:609–662.
- 95. Hazelhoff MH, Bulacio RP, Torres AM. Gender related differences in kidney injury induced by mercury. Int J Mol Sci. 2012;13:10523– 10536.
- 96. Clarkson TW, Magos L. The toxicology of mercury and its chemical compounds. Crit Rev Toxicol. 2006;36:609–662.

- 97. Malandrino P, Russo M, Ronchi A, Minoia C, Cataldo D, Regalbuto C, et al. Increased thyroid cancer incidence in a basaltic volcanic area is associated with non-anthropogenic pollution and biocontamination. Endocrine. 2016;53(2):471–479.
- 98. Abnoos H, Fereidoni M, Mahdavi-Shahri N, Haddad F, Jalal R. Developmental study of mercury effects on the fruit fly (Drosophila melanogaster). Interdiscip Toxicol. 2013;6(1):34–40.
- 99. Crespo-López ME, Macêdo GL, Pereira SI, Arrifano GP, Picanço-Diniz DL, Nascimento JLD, et al. Mercury and human genotoxicity: critical considerations and possible molecular mechanisms. Pharmacol Res. 2009;60(4):212– 220.
- 100. Zaichick S, Zaichick V, Karandashev V, Nosenko S. Accumulation of rare earth elements in human bone within the lifespan. Metallomics. 2011;3(2):186–194.
- 101. Horovitz CT, Toxicology C, Scandium P, Yttrium. New York, NY, USA: Springer; 2000.
- 102. Roohani N, Hurrell R, Kelishadi R, Schulin R. Zinc and its importance for human health: An integrative review. J Res Med Sci. 2013;18:144–157.
- 103. To PK, Do MH, Cho JH, Jung C. Growth modulatory role of zinc in prostate cancer and

application to cancer therapeutics. Int J Mol Sci. 2020;21(8):2991–2991.

- 104. Barber RG, Grenier ZA, Burkhead JL. Copper toxicity is not just oxidative damage: zinc systems and insight from Wilson disease. Biomedicines. 2021;9(3):316–316.
- 105. Alam S, Kelleher SL. Cellular mechanisms of zinc dysregulation: a perspective on zinc homeostasis as an etiological factor in the development and progression of breast cancer. Nutrients. 2012;4(8):875–903.
- 106. Zaichick V, Zaichick S, Wynchank S. Intracellular zinc excess as one of the main factors in the etiology of prostate cancer. J Anal Oncol. 2016;5(3):124–131.
- 107. Jouybari L, Kiani F, Akbari A, Sanagoo A, Sayehmiri F, Aaseth J, et al. A meta-analysis of zinc levels in breast cancer. J Trace Elem Med Biol. 2019;56:90–99.

How to cite this article: Zaichick V. Trace Element Contents in Thyroid of Patients with Diagnosed Nodular Goiter Investigated by Instrumental Neutron Activation Analysis. Journal of Medical Research and Health Sciences. 2021;1405–1417. https://doi.o rg/10.52845/JMRHS/2021-4-8-5