



Research Article

Comparison of Nineteen Chemical Elements in Thyroid Tissue adjacent to Thyroid Malignant and Benign Nodules using Nuclear Analytical Methods

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Abstract

Background: Thyroid Nodules (TN) are the most common endocrine disorder worldwide. Etiology and pathogenesis of thyroid benign and malignant nodules (TBN and TMN, respectively) are still not enough understood. The present study was performed to clarify the role of some chemical elements (ChEs) in the origination and development of TN. **Methods:** Contents of ChEs such as silver (Ag), calcium (Ca), chlorine (Cl), cobalt (Co), chromium (Cr), copper (Cu), iron (Fe), mercury (Hg), iodine (I), potassium (K), magnesium (Mg), manganese (Mn), sodium (Na), rubidium (Rb), ammonium (Sb), scandium (Sc), selenium (Se), strontium (Sr), and zinc (Zn) were prospectively evaluated in thyroid tissue adjacent to TBN (79 patients) and to TMN (41 patients). Measurements were performed using a combination of non-destructive nuclear analytical methods: X-ray fluorescence and instrumental neutron activation analysis. Results of the study were additionally compared with previously obtained data for the same ChEs in “normal” thyroid tissue. **Results:** It was found that in thyroid tissue adjacent to TMN the mass fractions of I, Rb, and Se were 47%, 79%, and 58%, respectively, higher while mass fractions Cl and Na were 42% and 29%, respectively, lower than in thyroid tissue adjacent to TBN. The common characteristics of thyroid tissue adjacent to TBN and TMN were similar contents of Ca, Cr, Fe, K, Mg, Mn, Sb, Sc, Se, Sr and Zn, as well as elevated levels of Ag, Cl, Co, Cu, Hg, I, Na, and Rb, which overtook those in “normal” thyroid approximately in 32, 2.2, 1.8, 2.2, 41, 1.4, 1.4, and 1.7 times, respectively. **Conclusions:** Role of ChEs in etiology and pathogenesis TBN and TMN is similar and excessive accumulation of Ag, Cl, Co, Cu, Hg, I, Na and Rb in thyroid tissue may be involved in the TN origination and development.

Keywords: Chemical elements; Neutron activation analysis; Thyroid; Thyroid malignant and benign nodules; X-ray fluorescence

Introduction

Thyroid benign and malignant nodules (TBN and TMN, respectively) are the most common endocrine disorder worldwide. Moreover, in some parts of the world, especially those of current or former iodine deficiency, thyroid nodules (TN) are still an endemic disease [1]. Incidence of TBN and TMN has been growing steadily

over the past four decades, despite the use of iodine prophylaxis in many countries [2]. Some factors causing this higher incidence of TN were described in literature [3] and analysis of these data shown intriguing links between the etiologies of TBN and TMN [2,3]. In other words, the factors contributing to increases in the incidence of TBN are the same as those contributing to increases in TMN. However, the current state of knowledge regarding TN demonstrates that the etiology and pathogenesis of TBN and TMN are still not enough understood, because there are many not adequately explored chemicals, which induced thyroid hormone perturbations leading to these diseases.

For over 20th century, there was the dominant opinion that TN is the simple consequence of iodine deficiency [4]. However, it was found that TN is a frequent disease even in those countries and regions where the population is never exposed to iodine shortage. Moreover, it was shown that iodine excess has severe consequences on human health and associated with the presence of TN [5-8]. It was also demonstrated that besides the iodine deficiency and excess many other dietary, environmental, and occupational factors are associated with the TN incidence [3,9-11]. Among these factors a disturbance of evolutionary stable input of many Chemical Elements (ChEs) in human body after industrial revolution plays a significant role in etiology of TN [12].

Besides iodine, many other ChEs have also essential physiological functions [13]. Essential or toxic (goitrogenic, mutagenic, carcinogenic) properties of ChEs depend on tissue-specific need or tolerance, respectively [13]. Excessive accumulation or an imbalance of the ChEs may disturb the cell functions and may result in cellular proliferation, 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 and other ChEs 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 many ChEs content with age in the thyroid of males and females were studied and age- and gender-dependence of some ChEs was observed [25-41]. Furthermore, a significant difference between some ChEs contents in colloid goiter, thyroiditis, thyroid adenoma and cancer in comparison with normal thyroid was demonstrated [42-47].

The present study was performed to clarify the role of some ChEs in the etiology of TBN and TMN. Having this in mind, the aim of this exploratory study was to examine differences in the content of silver (Ag), calcium (Ca), chlorine (Cl), cobalt (Co), chromium (Cr), copper (Cu), iron (Fe), mercury (Hg), iodine (I), potassium (K), magnesium (Mg), manganese (Mn), sodium (Na), rubidium (Rb), ammonium (Sb), scandium (Sc), selenium (Se), strontium (Sr), and zinc (Zn) in thyroid tissue adjacent to TN using a non-destructive energy-dispersive X-Ray fluorescent analysis (EDXRF) combined with instrumental neutron activation analysis with high resolution spectrometry of short and long-lived radionuclides (INAA-SLR and INAA-LLR, respectively) and to compare the levels of these ChEs in two groups of samples (tissue adjacent to TBN and TMN, respectively). Moreover, for understanding a possible role of ChEs in etiology and pathogenesis of TN results of the study were compared with previously obtained data for the same ChEs in "normal" thyroid tissue [42-47].

Material and Methods

All patients suffered from TBN (n=79, mean age M±SD was 44±11 years, range 22-64) and from TMN (n=41, mean age M±SD was 46±15 years, range 16-75) were hospitalized in the Head and Neck Department of the Medical Radiological Research Centre (MRRC), Obninsk. Thick-needle puncture biopsy of suspicious nodules of the thyroid performed on every patient, to permit morphological study of thyroid tissue at these sites and to estimate their trace element contents. In all the cases diagnosis has been confirmed by clinical and morphological results obtained during studies of biopsy and resected materials. Histological conclusions for benign nodules were: 46 colloid goiter, 19 thyroid adenoma, 8 Hashimoto's thyroiditis, and 6 Riedel's Struma, whereas for thyroid malignant tumors were: 25 papillary adenocarcinomas, 8 follicular adenocarcinomas, 7 solid carcinomas, and 1 reticulosarcoma. Samples of visually intact thyroid tissue adjacent to TBN and TMN were taken from resected materials.

"Normal" thyroids for the control group samples were removed at necropsy from 105 deceased (mean age 44±21 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.

The Ethical Committees of MRRC approved all the studies. 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. Informed consent was obtain from all individual participants included in the study.

All tissue samples obtained from tumors and visually intact tissue adjacent to tumors were divided into two portions using a titanium scalpel to prevent contamination by ChEs of stainless steel [48]. One used for morphological study while the other intended for ChEs analysis. After the samples intended for ChEs analysis were weighed, they were freeze-dried and homogenized [49]. To determine contents of the ChE 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 used as standards. Ten sub-samples of Certified Reference Material (CRM) of the International Atomic Energy Agency (IAEA) IAEA H-4 (animal muscle) and IAEA HH-1 (human hair) weighing about 100 mg were treated and analyzed in the same conditions as thyroid samples to estimate the precision and accuracy of results.

The content of Cu, Fe, Rb, Sr, and Zn were determined by EDXRF. Details of the relevant facility for this method, source with

¹⁰⁹Cd radionuclide, methods of analysis and the results of quality control were presented in our earlier publications concerning the EDXRF of ChE contents in human thyroid [25,26] and prostate tissue [51].

The content of Br, Ca, Cl, I, K, Mg, Mn and Na were determined by INAA-SLR using a horizontal channel equipped with the pneumatic rabbit system of the WWR-c research nuclear reactor (Branch of Karpov Institute, Obninsk). Details of used neutron flux, nuclear reactions, radionuclides, gamma-energies, spectrometric unit, sample preparation and measurement were presented in our earlier publications concerning the INAA-SLR of ChE contents in human thyroid [27,28], prostate [52,53], and scalp hair [54].

In a few days after non-destructive INAA-SLR all thyroid samples were repacked and used for INAA-LLR. A vertical channel of the WWR-c research nuclear reactor (Branch of Karpov Institute, Obninsk) was applied to determine the content of Ag, Co, Cr, Fe, Hg, Rb, Sb, Sc, Se, and Zn by INAA-LLR. Details of used neutron flux, nuclear reactions, radionuclides, gamma-energies, spectrometric unit, sample preparation and measurement were presented in our earlier publications concerning the INAA-LLR

of ChE contents in human thyroid [29,30], scalp hair [54], and prostate [55].

A dedicated computer program for INAA-SLR and INAA-LLR mode optimization was used [56]. All thyroid samples for ChEs analysis were prepared in duplicate and mean values of ChEs contents were used in final calculation. Mean values of ChE contents were used in final calculation for the Fe, Rb, and Zn mass fractions measured by two methods. Using Microsoft Office Excel software, a summary of the statistics, including, arithmetic mean, standard deviation of mean, standard error of mean, minimum and maximum values, median, percentiles with 0.025 and 0.975 levels was calculated for ChEs contents in two groups of tissue adjacent to TBN and TMN. Data for “normal” thyroid were taken from our previous publications [42-47]. The difference in the results between two groups of samples “adjacent to TBN” and “adjacent to TMN”, as well as between “normal” and “adjacent to TBN and TMN combined” was evaluated by the parametric Student’s *t*-test and non-parametric Wilcoxon-Mann-Whitney *U*-test.

Results

All thyroid tissue samples investigated and used in the present study are included in Table 1.

Thyroid tissue	n	Age of patient/individuals, years	
		M±SD	Range
Thyroid tissue adjacent to thyroid benign nodules	79	44±11	22 - 64
Thyroid tissue adjacent to thyroid malignant nodules	41	46±15	16-75
Thyroid tissue of “normal” glands	105	44±21	2-87

M: Arithmetic Mean; SD: Standard Deviation

Table 1: Tissue samples investigated and used in the present study.

Table 2 presents 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, Ca, Cl, Co, Cr, Cu, Fe, Hg, I, K, Mg, Mn, Na, Rb, Sb, Sc, Se, Sr, and Zn mass fraction in thyroid intact tissue samples of two groups “adjacent to TBN” and “adjacent to TMN”.

Tissue	Element	Mean	SD	SEM	Min	Max	Median	P 0.025	P 0.975
TATBN	Ag	0.474	0.662	0.130	0.0210	3.31	0.282	0.0516	2.07
	Ca	1532	1700	380	418	6466	994	442	6312
	Cl	9203	6033	1384	2881	23731	8161	3294	22429
	Co	0.0728	0.0979	0.0170	0.0051	0.594	0.0525	0.0086	0.219
	Cr	0.575	0.618	0.108	0.0180	3.14	0.401	0.0596	2.19
	Cu	10.2	7.9	4.0	3.60	20.4	8.35	3.65	19.8
	Fe	213	140	24	41.5	620	171	58.2	557

	Hg	1.36	0.96	0.17	0.0140	4.68	1.21	0.268	4.25
	I	2158	1436	214	343	7912	1917	527	5441
	K	6793	4044	862	3406	18255	5607	3500	18077
	Mg	316	275	59	15.0	987	292	15.0	890
	Mn	1.77	1.66	0.36	0.100	5.83	1.10	0.100	5.67
	Na	10850	5541	1209	4663	31343	9642	5548	23981
	Rb	10.5	4.2	0.7	4.10	20.0	9.80	4.74	19.4
	Sb	0.131	0.174	0.030	0.0076	0.757	0.0759	0.0269	0.749
	Sc	0.0058	0.0147	0.0020	0.0002	0.0654	0.0002	0.0002	0.0468
	Se	1.95	0.87	0.15	0.647	4.34	1.65	0.906	3.66
	Sr	6.28	5.17	2.59	1.30	13.5	5.15	1.54	12.9
	Zn	121	118	20	34.2	669	91.3	43.0	401
TATMN	Ag	0.503	0.450	0.103	0.079	2.00	0.303	0.0984	1.53
	Ca	862	560	140	81.0	1909	672	149	1822
	Cl	5339	22512	581	2526	11767	4922	2595	10201
	Co	0.0707	0.0581	0.0120	0.0152	0.205	0.0455	0.0170	0.201
	Cr	0.556	0.468	0.094	0.0512	1.58	0.457	0.0589	1.56
	Cu	8.08	3.15	1.58	4.90	12.1	7.65	5.01	11.9
	Fe	244	137	27	95.2	752	213	104	591
	Hg	2.19	1.92	0.38	0.0160	7.78	1.43	0.158	6.50
	I	3183	1673	301	563	8240	2982	853	7766
	K	5717	2525	652	2097	12681	5429	2466	10953
	Mg	339	407	105	15.0	1412	199	15.0	1287
	Mn	1.72	1.63	0.41	0.410	6.78	1.15	0.429	5.54
	Na	7671	2597	649	3865	14373	7434	4169	13009
	Rb	18.8	17.0	3.3	5.00	67.0	11.9	5.69	65.6
	Sb	0.247	0.416	0.085	0.0069	1.77	0.0634	0.0159	1.38
	Sc	0.0059	0.0134	0.0030	0.0002	0.0539	0.0002	0.0002	0.0442
	Se	3.08	1.67	0.33	0.704	6.91	2.56	0.942	6.89
	Sr	1.16	0.29	0.14	0.83	1.40	1.20	0.84	1.40
	Zn	109	55	11	20.4	272	109	29.1	213
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.									

Table 2: Some statistical parameters of Ag, Br, Ca, Cl, Co, Cr, Cu, Fe, Hg, I, K, Mg, Mn, Na, Rb, Sb, Sc, Se, Sr, and Zn mass fraction (mg/kg, dry mass basis) in thyroid tissue adjacent to thyroid benign (TATBN) and malignant (TATMN) nodules.

The ratios of means and the comparison of mean values of Ag, Ca, Cl, Co, Cr, Cu, Fe, Hg, I, K, Mg, Mn, Na, Rb, Sb, Sc, Se, Sr, and Zn mass fractions in pair of sample groups such as “adjacent to TBN” and “adjacent to TMN” is presented in Table 3.

Element	Thyroid tissue adjacent to thyroid nodules				Ratio
	TATBN	TATMN	Student's t-test <i>p</i> £	U-test <i>p</i>	TATMN/TATBN
Ag	0.474±0.130	0.503±0.103	0.864	>0.05	1.06
Ca	1532±380	862±140	0.111	>0.05	0.56
Cl	9203±1384	5339±581	0.017	≤0.01	0.58
Co	0.0728±0.0170	0.0707±0.0120	0.918	>0.05	0.97
Cr	0.575±0.108	0.556±0.094	0.898	>0.05	0.97
Cu	10.2±4.0	8.08±1.58	0.648	>0.05	0.79
Fe	213±24	244±27	0.389	>0.05	1.15
Hg	1.36±0.17	2.19±0.38	0.057	>0.05	1.62
I	2158±214	3183±301	0.0074	≤0.01	1.47
K	6793±862	5717±652	0.326	>0.05	0.84
Mg	316±59	339±105	0.851	>0.05	1.07
Mn	1.77±0.36	1.72±0.41	0.921	>0.05	0.97
Na	10850±1209	7671±649	0.028	≤0.01	0.71
Rb	10.5±0.7	18.8±3.3	0.022	≤0.01	1.79
Sb	0.131±0.030	0.247±0.085	0.208	>0.05	1.89
Sc	0.0058±0.0020	0.0059±0.0030	0.964	>0.05	1.02
Se	1.95±0.15	3.08±0.33	0.0033	≤0.01	1.58
Sr	6.28±2.59	1.16±0.14	0.142	>0.05	0.18
Zn	121±20	109±11	0.595	>0.05	0.90

M: Arithmetic Mean; Sem: Standard Error Of Mean; Statistically Significant Values Are In Bold.

Table 3: Differences between mean values (M±SEM) of Ag, Br, Ca, Cl, Co, Cr, Cu, Fe, Hg, I, K, Mg, Mn, Na, Rb, Sb, Sc, Se, Sr, and Zn mass fraction (mg/kg, dry mass basis) in thyroid tissue adjacent to thyroid benign (TATBN) and malignant (TATMN) nodules.

Table 4 depicts 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, Ca, Cl, Co, Cr, Cu, Fe, Hg, I, K, Mg, Mn, Na, Rb, Sb, Sc, Se, Sr, and Zn mass fraction in thyroid tissue adjacent “TTA” to TN (two groups “adjacent to TBN” and “adjacent to TMN” combined).

Tissue	Element	Mean	SD	SEM	Min	Max	Median	P 0.025	P 0.975
TTA	Ag	0.486	0.576	0.086	0.0210	3.31	0.297	0.0709	1.93
	Ca	1234	1348	225	81.0	6466	918	239	6182
	Cl	7498	5079	871	2526	23731	5456	2688	21344
	Co	0.072	0.083	0.011	0.0051	0.594	0.0467	0.0115	0.202
	Cr	0.567	0.554	0.073	0.018	3.14	0.429	0.0566	1.88
	Cu	9.13	5.68	2.01	3.60	20.4	7.65	3.72	19.0
	Fe	224	138	18	41.5	752	186	68.6	581
	Hg	1.72	1.50	0.20	0.0140	7.78	1.30	0.117	5.37
	I	2577	1608	184	343	8240	2400	554	7646
	K	6357	3508	577	2097	18255	5429	3046	17950
	Mg	325	329	54	15.0	1412	247	15.0	1092
	Mn	1.75	1.62	0.27	0.100	6.78	1.11	0.100	5.93
	Na	9475	4735	778	3865	31343	8283	4583	18091
	Rb	14.1	12.3	1.6	4.10	67.0	10.6	4.90	54.5
	Sb	0.180	0.303	0.040	0.0069	1.77	0.075	0.0136	0.971
	Sc	0.0058	0.0141	0.0020	0.0002	0.0654	0.0002	0.0002	0.0490
	Se	2.43	1.38	0.18	0.647	6.91	2.12	0.828	6.45
	Sr	3.71	4.36	1.54	0.83	13.5	1.40	0.860	12.2
	Zn	110	68	8.7	20.4	344	101	34.0	314

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.

Table 4: Some statistical parameters of Ag, Br, Ca, Cl, Co, Cr, Cu, Fe, Hg, I, K, Mg, Mn, Na, Rb, Sb, Sc, Se, Sr, and Zn mass fraction (mg/kg, dry mass basis) in in thyroid tissue adjacent (TTA) to thyroid benign and malignant nodules (combined).

The ratios of means and the comparison of mean values of Ag, Ca, Cl, Co, Cr, Cu, Fe, Hg, I, K, Mg, Mn, Na, Rb, Sb, Sc, Se, Sr, and Zn mass fractions in pair of sample groups such as normal thyroid tissue “NT” and “TTA” is presented in Table 5.

Element	Thyroid tissue				Ratio
	NT	TTA	Student’s t-test <i>p</i> £	U-test <i>p</i>	TTA/NT
Ag	0.0151±0.0016	0.486±0.086	0.0000019	≤ 0.01	32.2
Ca	1711±109	1234±225	0.062	>0.05	0.72
Cl	3400±174	7498±871	0.000049	≤ 0.01	2.21
Co	0.0399±0.0030	0.0720±0.0110	0.0056	≤ 0.01	1.80

Cr	0.539±0.032	0.567±0.073	0.724	>0.05	1.05
Cu	4.23±0.18	9.13±2.01	0.045	≤0.01	2.16
Fe	223±10	224±18	0.953	>0.05	1.00
Hg	0.0421±0.0041	1.72±0.20	0.0000001	≤0.01	40.9
I	1841±107	2577±184	0.00076	≤0.01	1.40
K	6071±306	6357±577	0.663	>0.05	1.05
Mg	285±17	325±54	0.476	>0.05	1.14
Mn	1.35±0.07	1.75±0.27	0.161	>0.05	1.30
Na	6702±178	9475±778	0.0013	≤0.01	1.41
Rb	8.16±0.49	14.1±1.6	0.00062	≤0.01	1.73
Sb	0.111±0.008	0.180±0.040	0.094	>0.05	1.62
Sc	0.0046±0.0008	0.0058±0.0020	0.523	>0.05	1.26
Se	2.32±0.14	2.43±0.18	0.608	>0.05	1.05
Sr	4.55±0.37	3.71±1.54	0.614	>0.05	0.82
Zn	105.1±4.3	110±8.7	0.587	>0.05	1.05

M: Arithmetic Mean; SEM: Standard Error Of Mean; Statistically Significant Values are in Bold.

Table 5: Differences between mean values (M±SEM) of Ag, Br, Ca, Cl, Co, Cr, Cu, Fe, Hg, I, K, Mg, Mn, Na, Rb, Sb, Sc, Se, Sr, and Zn mass fraction (mg/kg, dry mass basis) in normal thyroid (NT) and thyroid tissue adjacent to thyroid benign and malignant nodules (TTA)

Discussion

As it was shown before [25-30,51-55] good agreement of the Ag, Ca, Cl, Co, Cr, Cu, Fe, Hg, I, K, Mg, Mn, Na, Rb, Sb, Sc, Se, Sr and Zn contents in CRM IAEA H-4 and IAEA HH-1 (human hair) samples determined by EDXRF, INAA-SLR and INAA-LLR with the certified data of these CRMs indicates acceptable accuracy of the results obtained in the study of “adjacent to TBN”, “adjacent to TMN”, “NT” and “TTA” groups of thyroid tissue samples presented in Tables 1-4.

From Table 2, it is observed that in thyroid tissue adjacent to TMN the mass fraction of I, Rb, and Se is 47%, 79%, and 58%, respectively, higher while mass fractions Cl and Na 42% and 29%, respectively, lower than in thyroid tissue adjacent to TBN. In a general sense Ag, Ca, Co, Cr, Cu, Fe, Hg, K, Mg, Mn, Sb, Sc, Sr and Zn contents found in the “adjacent to TBN” and “adjacent to TMN” groups of thyroid tissue samples were similar (Table 2). It allowed combine data obtained for two groups for the purposes of finding a common ChEs composition of TTA to TN and improving statistical characteristics of results for this group of samples (Table 3).

From obtained results it was found that the common characteristics of thyroid tissue adjacent to TBN and TMN were

elevated levels of Ag, Cl, Co, Cu, Hg, I, Na, and Rb, which overdrove those in “normal” thyroid approximately in 32, 2.2, 1.8, 2.2, 41, 1.4, 1.4, and 1.7 times, respectively (Table 4). Thus, if we accept the ChEs contents in “normal” thyroid glands as a norm, we have to conclude that with a nodular transformation the Ag, Cl, Co, Cu, Hg, I, Na and Rb the contents in thyroid intact tissue adjacent to TN significantly changed.

Characteristically, elevated or reduced levels of ChEs observed in thyroid nodules are discussed in terms of their potential role in the initiation and promotion of these thyroid lesions. In other words, using the low or high levels of the ChEs in affected thyroid tissues researchers try to determine the role of the deficiency or excess of each ChEs in the etiology and pathogenesis of thyroid diseases. In our opinion, abnormal levels of some ChEs in TN could be and cause, and also effect of thyroid tissue transformation. From the results of such kind studies, it is not always possible to decide whether the measured decrease or increase in ChEs level in pathologically altered tissue is the reason for alterations or vice versa. According to our opinion, investigation of ChEs contents in thyroid tissue adjacent to TN and comparison obtained results with ChEs levels typical of “normal” thyroid gland may give additional useful information on the topic because these data show conditions of tissue in which TN were originated and developed.

Silver

Ag is a TE with no recognized trace metal value in the human body [57]. Food is the major intake source of Ag and this metal authorized as a food additive (E174) in the EU [58]. Another source of Ag is contact with skin and mucosal surfaces because Ag is widely used in different applications (e.g., jewelry, wound dressings, or eye drops) [59]. Ag in metal form and inorganic Ag compounds ionize in the presence of water, body fluids or tissue exudates. The silver ion Ag^+ is biologically active and readily interacts with proteins, amino acid residues, free anions and receptors on mammalian and eukaryotic cell membranes [60]. 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 [61]. Experimental studies shown that Ag nanoparticles may affect thyroid hormone metabolism [62]. More detailed knowledge of the Ag toxicity can lead to a better understanding of the impact on human health, including thyroid function.

Chlorine and sodium

Cl and Na are ubiquitous, extracellular electrolytes essential to more than one metabolic pathway. In the body, Cl and Na mostly present as sodium chloride. Therefore, as usual, there is a correlation between Na and Cl contents in tissues and fluids of human body. Because Cl is halogen like I, in the thyroid gland the biological behavior of chloride has to be similar to the biological behavior of iodide. The main source of natural Cl for human body is salt in food and chlorinated drinking water. Environment (air, water and food) polluted by artificial nonorganic Cl-contained compounds, for example such as sodium chlorate ($NaClO_3$), and organic Cl-contained compounds, for example such as polychlorinated biphenyls (PCBs) and dioxin, is other source. There is a clear association between using chlorinated drinking water, levels $NaClO_3$, PCBs and dioxin in environment and thyroid disorders, including cancer [63-67]. Thus, on the one hand, the accumulated data suggest that Cl level in thyroid tissue might be responsible for TMNs development. However, on the other hand, It is well known that Cl and Na mass fractions in human tissue samples depend mainly on the extracellular water volume [68]. TN and thyroid tissues adjacent to nodules can be more vascularized than normal thyroid. Because blood is extracellular liquid, it is possible to speculate that more intensive vascularization could be the reason for elevated levels of Cl and Na in thyroid tissue adjacent to TB and TMN. If that is the only case, the equilibrium between Cl and Na increases has to be, however, in comparison with “normal” thyroid the change of Cl level in adjacent tissue is significantly higher than change of Na level. Thus, it is possible to assume that an excessive accumulation of Cl in thyroid tissue is involved in TBN and TMN etiology.

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 [69,70]. Co is genotoxic and carcinogenic, mainly caused by oxidative DNA damage by reactive oxygen species, perhaps combined with inhibition of DNA repair [71]. In our previous studies, it was found that a significant age-related increase of Co content in female thyroid [29]. Therefore, a goitrogenic and, probably, carcinogenic effect of excessive Co level in the thyroid of old females was assumed. Elevated level of Co in TBN and TMN, observed in the present study, supports this conclusion.

Copper

Cu is a ubiquitous element in the human body, which plays many roles at different levels. Various Cu-enzymes (such as amine oxidase, ceruloplasmin, cytochrome-c oxidase, dopamine-monoxygenase, extracellular superoxide dismutase, lysyl oxidase, peptidylglycineamidating monoxygenase, Cu/Zn superoxide dismutase, and tyrosinase) mediate the effects of Cu deficiency or excess. Cu excess can have severe negative impacts. Cu generates oxygen radicals and many investigators have hypothesized that excess copper might cause cellular injury via an oxidative pathway, giving rise to enhanced lipid peroxidation, thiol oxidation, and, ultimately, DNA damage [72-74]. Thus, Cu accumulation in thyroid parenchyma with age may be involved in oxidative stress, dwindling gland function, and increasing risk of TBN and TMN [25,26,31-34]. The significantly elevated level of Cu in thyroid tissue adjacent to TBN and TMN, observed in the present study, supports this speculation. However, an overall comprehension of Cu homeostasis and physiology, which is not yet acquired, is mandatory to establish Cu exact role in TBN and TMN etiology and metabolism.

Mercury

In the general population, potential sources of Hg exposure include the inhalation of this metal vapor in the air, ingestion of contaminated foods and drinking water, and exposure to dental amalgam through dental care [75]. Hg is one of the most dangerous environmental pollutants [76]. 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. Many experimental and occupational studies of Hg in different chemical states shown significant alterations in thyroid hormones metabolism and thyroid gland parenchyma [77,78]. Moreover, Hg was classified as certain or probable carcinogen by the International Agency for Research on Cancer [79]. For example, in Hg polluted area thyroid cancer incidence was almost 2 times higher than in adjacent control areas [80].

Iodine

To date, it was well established, that iodine excess has severe consequences on human health and associated with the presence of TBN and TMN [4-8,81-84]. In the present study, elevated level of I in thyroid tissue adjacent to TBN and TMN was found in comparison with “normal” thyroid. Thus, on the one hand, it is likely that elevated level of I in thyroid tissue might be involved in the TN origination and development. On the other hand, however, elevated level of I in thyroid tissue adjacent to TN may explain by unusually intensive work of this tissue. Compared to other soft tissues, the human thyroid gland has higher levels of I, because this element plays an important role in its normal functions, through the production of thyroid hormones (thyroxin and triiodothyronine) which are essential for cellular oxidation, growth, reproduction, and the activity of the central and autonomic nervous system. As was shown in our previous study, TBN and, particularly, TMN transformation of thyroid gland is accompanied by a significant loss of tissue-specific functional features, which leads to a significant reduction in I content associated with functional characteristics of the human thyroid tissue [43-47]. Because the affected part of gland reduced productions of thyroid hormones, the rest “intact” part of thyroid tries to compensate thyroid hormones deficiency and work more intensive than usual.

Rubidium

There is very little information about Rb effects on thyroid function. Rb as a monovalent cation Rb^+ is transferred through membrane by the $Na^+K^+-ATPase$ pump like K^+ and concentrated in the intracellular space of cells. Thus, Rb seems to be more intensively concentrated in the intracellular space of cells. The source of Rb elevated level in thyroid tissue adjacent to TN may be Rb environment overload. The excessive Rb intake may result a replacement of medium potassium by Rb, which effects on iodide transport and iodoaminoacid synthesis by thyroid [85]. The source of Rb increase in thyroid tissue adjacent to TN may be not only the excessive intake of this TE in organism from the environment, but also changed $Na^+K^+-ATPase$ or $H^+K^+-ATPase$ pump membrane transport systems for monovalent cations, which can be stimulated by endocrin system, including thyroid hormones [86]. It was found also that Rb has some function in immune response [87] and that elevated concentration of Rb could modulate proliferative responses of the cell, as was shown for bone marrow leukocytes [88]. These data partially clarify the possible role of Rb in etiology and pathogenesis of TBN and TMN.

Selenium

The high level of Se content found just in thyroid tissue adjacent to TMN cannot be regarded as pure chance. The seleno-protein characterized as Se-dependent glutathione peroxidase (Se-GSH-Px) is involved in protecting cells from peroxidative damage.

This enzyme may reduce tissue concentration of free radicals and hydroperoxides. It is particular important for the thyroid gland, because thyroidal functions involve oxidation of iodide, which is incorporated into thyroglobulin, the precursor of the thyroid hormones. For oxidation of iodide thyroidal cells produce a specific thyroid peroxidase using of physiologically generated hydrogen-peroxide (H_2O_2) as a cofactor [89]. It follows that the thyroid parenchyma must continuously exposed to a physiological generation of H_2O_2 and in normal conditions must be a balance between levels of Se (as Se-GSH-Px) and H_2O_2 . The elevated level of Se in thyroid tissue adjacent to TMN was accompanied excessive accumulation of Ag, Co, Hg, I and Rb in comparison with “normal” values for these elements. Moreover, contents of Ag, Co, Hg, I and Rb in adjacent tissue were higher than in malignant nodules [47]. Thus, it might be assumed, that the elevated level of Se is reaction of adjacent tissue on an increase in concentration of free radicals and hydroperoxides in thyroid gland and that this increase preceded the TMN origination and development.

Limitations

This study has several limitations. Firstly, analytical techniques employed in this study measure only nineteen ChE (Ag, Ca, Cl, Co, Cr, Cu, Fe, Hg, I, K, Mg, Mn, Na, Rb, Sb, Sc, Se, Sr and Zn) mass fractions. Future studies should be directed toward using other analytical methods which will extend the list of ChEs investigated in thyroid tissue adjacent to TN. Secondly, the sample size of TBN and TMN group was relatively small and prevented investigations of ChEs contents in this group using differentials like gender, functional activity of nodules, stage of disease and dietary habits of patients with TN. Lastly, generalization of our results may be limited to Russian population. Despite these limitations, this study provides evidence on some ChEs level alteration in thyroid tissue adjacent to TN and shows the necessity to continue ChEs research of TN.

Conclusion

In this work, ChEs analysis was carried out in the thyroid tissue adjacent to TBN and TMN using a combination of nuclear analytical methods. It was shown that a combination of three methods such as EDXRF, INAA-SLR and INAA-LLR is an adequate analytical tool for the non-destructive determination of nineteen ChE (Ag, Ca, Cl, Co, Cr, Cu, Fe, Hg, I, K, Mg, Mn, Na, Rb, Sb, Sc, Se, Sr, and Zn) content in the tissue samples of human thyroid in norm and pathology. I found that in thyroid tissue adjacent to TMN the mass fraction of I, Rb, and Se is 47%, 79%, and 58%, respectively, higher while mass fractions Cl and Na 42% and 29%, respectively, lower than in thyroid tissue adjacent to TBN. The common characteristics of thyroid tissue adjacent to TBN and TMN were elevated levels of Ag, Cl, Co, Cu, Hg, I, Na, and Rb, which overdrew those in “normal” thyroid approximately

in 32, 2.2, 1.8, 2.2, 41, 1.4, 1.4, and 1.7 times, respectively, and similar contents of Ca, Cr, Fe, K, Mg, Mn, Sb, Sc, Se, Sr and Zn. Thus, from results obtained, it was possible to conclude that the role of ChEs in etiology and pathogenesis TBN and TMN is similar and excessive accumulation of Ag, Cl, Co, Cu, Hg, I, Na and Rb in thyroid tissue may be involved in the TN origination and development.

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References

1. Verburg FA, Reiners C (2010) The association between multinodular goiter and thyroid cancer. *Minerva Endocrinol* 35: 187-192.
2. Simsir IY, Cetinkalp S, Kabalak T (2020) Review of factors contributing to nodular goiter and thyroid carcinoma. *Med Princ Pract* 29: 1-5.
3. Foster JR, Tinwell H, Melching-Kollmuss S (2021) A review of species differences in the control of, and response to, chemical-induced thyroid hormone perturbations leading to thyroid cancer. *Arch Toxicol* 95:807-836.
4. Barrea L, Gallo M, Ruggeri RM, Di Giacinto P, Sesti F, et al, (2021) Nutritional status and follicular-derived thyroid cancer: An update. *Crit Rev Food Sci Nutr* 61:25-59.
5. Zaichick V (1998) Iodine excess and thyroid cancer. *J Trace Elem Exp Med* 11:508-509.
6. Zaichick V, Iljina T (1998) Dietary iodine supplementation effect on the rat thyroid ¹³¹I blastomogenic action. In: *Die Bedeutung der Mengen- und Spurenelemente*. 18. Arbeitstangung. Jena: Friedrich-Schiller-Universität 294-306.
7. Kim K, Cho SW, Park YJ, Lee KE, Lee D-W et al., (2021) Association between iodine intake, thyroid function, and papillary thyroid cancer: A case-control study. *Endocrinol Metab (Seoul)* 36:790-799.
8. Vargas-Uricoechea P, Pinzón-Fernández MV, Bastidas-Sánchez BE, Jojoa-Tobar E, Ramírez-Bejarano LE et al., (2019) Iodine status in the colombian population and the impact of universal salt iodization: a double-edged sword? *J Nutr Metab*.
9. Stojavljević A, Rovčanin B, Krstić D, Borković-Mitić S, Paunović I et al., (2019) 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* 12: 255-264.
10. Fahim YA, Sharaf NE, Hasani IW, Ragab EA, Abdelhakim HK (2020) Assessment of thyroid function and oxidative stress state in foundry workers exposed to lead. *J Health Pollut* 10: 200903.
11. Liu M, Song J, Jiang Y, Lin Y, Peng J et al., (2021) A case-control study on the association of mineral elements exposure and thyroid tumor and goiter. *Ecotoxicol Environ Saf* 208:111615.
12. Zaichick V (2006) Medical elementology as a new scientific discipline. *J Radioanal Nucl Chem* 269:303-309.
13. Moncayo R, Moncayo H (2017) 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* 7:115-119.
14. Beyersmann D, Hartwig A (2008) Carcinogenic metal compounds: recent insight into molecular and cellular mechanisms. *Arch Toxicol* 82:493-512.
15. Martinez-Zamudio R, Ha HC (2011) Environmental epigenetics in metal exposure. *Epigenetics* 6:820-827.
16. Zaichick V, Raibukhin YuS, Melnik AD, Cherkashin VI (1970) Neutron-activation analysis in the study of the behavior of iodine in the organism. *Med Radiol (Mosk)* 15:33-36.
17. Zaichick V, Matveenko EG, Vtiurin BM, Medvedev VS (1982) Intrathyroid iodine in the diagnosis of thyroid cancer. *Vopr Onkol* 28:18-24.
18. Zaichick V, Tsyb AF, Vtyurin BM (1995) Trace elements and thyroid cancer. *Analyst* 120:817-821.
19. Zaichick V, Choporov YuYa (1996) Determination of the natural level of human intra-thyroid iodine by instrumental neutron activation analysis. *J Radioanal Nucl Chem* 207:153-161.
20. Zaichick V (1998) In vivo and in vitro application of energy-dispersive XRF in clinical investigations: experience and the future. *J Trace Elem Exp Med* 11:509-510.
21. Zaichick V, Zaichick S (1999) Energy-dispersive X-ray fluorescence of iodine in thyroid puncture biopsy specimens. *J Trace Microprobe Tech* 17:219-232.
22. Zaichick V (2000) Relevance of, and potentiality for in vivo intrathyroidal iodine determination. *Ann N Y Acad Sci* 904:630-632.
23. Zaichick V, Zaichick S (1997) Normal human intrathyroidal iodine. *Sci Total Environ* 206:39-56.
24. Zaichick V (1999) Human intrathyroidal iodine in health and non-thyroidal disease. In: *New aspects of trace element research* (Eds: M.Abdulla, M.Bost, S.Gamon, P.Arnaud, G.Chazot). London: Smith-Gordon; and Tokyo: Nishimura 114-119.
25. Zaichick V, Zaichick S (2017) Age-related changes of some trace element contents in intact thyroid of females investigated by energy dispersive X-ray fluorescent analysis. *Trends Geriatr Healthc* 1:31-38.
26. Zaichick V, Zaichick S (2017) Age-related changes of some trace element contents in intact thyroid of males investigated by energy dispersive X-ray fluorescent analysis. *MOJ Gerontol Ger* 1:133-140.
27. Zaichick V, Zaichick S (2017) 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* 1:5.1.
28. Zaichick V, Zaichick S (2017) 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* 1:1002.
29. Zaichick V, Zaichick S (2017) Age-related changes of Ag, Co, Cr, Fe, Hg, Rb, Sb, Sc, Se, and Zn contents in intact thyroid of females investigated by neutron activation analysis. *J Gerontol Geriatr Med* 3:015.
30. Zaichick V, Zaichick S (2017) Age-related changes of Ag, Co, Cr, Fe, Hg, Rb, Sb, Sc, Se, and Zn contents in intact thyroid of males investigated by neutron activation analysis. *Curr Trends Biomedical Eng Biosci* 4:555644.

31. Zaichick V, Zaichick S (2018) Effect of age on chemical element contents in female thyroid investigated by some nuclear analytical methods. *MicroMedicine* 6:47-61.
32. Zaichick V, Zaichick S (2018) 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* 2:202-212.
33. Zaichick V, Zaichick S (2018) 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* 3:1-10.
34. Zaichick V, Zaichick S (2018) Association between age and twenty chemical element contents in intact thyroid of males. *SM Gerontol Geriatr Res* 2:1014.
35. Zaichick V, Zaichick S (2018) Associations between age and 50 trace element contents and relationships in intact thyroid of males. *Aging Clin Exp Res* 30:1059-1070.
36. Zaichick V, Zaichick S (2018) Possible role of inadequate quantities of intra-thyroidal bromine, rubidium and zinc in the etiology of female subclinical hypothyroidism. *EC Gynaecology* 7:107-115.
37. Zaichick V, Zaichick S (2018) 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*.
38. Zaichick V, Zaichick S (2018) Possible role of inadequate quantities of intra-thyroidal cobalt, rubidium and zinc in the etiology of female subclinical hypothyroidism. *Womens Health Sci J*.
39. Zaichick V, Zaichick S (2018) 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* 2:340-355.
40. Zaichick V, Zaichick S (2018) Investigation of association between the high risk of female subclinical hypothyroidism and inadequate quantities of twenty intra-thyroidal chemical elements. *Clin Res: Gynecol Obstet* 1:1-18.
41. Zaichick V, Zaichick S (2018) 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* 2:23-37.
42. Zaichick V (2021) Comparison between Twenty Chemical Element Contents in Colloid Nodular Goiter of Different Histology. *Archives of Clinical Case Studies and Case Reports* 2:243-251.
43. Zaichick V (2021) Determination of twenty chemical element contents in normal and goitrous thyroid using X-ray fluorescent and neutron activation analysis. *World Journal of Advanced Research and Reviews* 11:130-146.
44. Zaichick V (2021) Evaluation of Twenty Chemical Element Contents in Thyroid Adenomas using X-Ray Fluorescent and Neutron Activation Analysis. *Journal of Cellular & Molecular Oncology*.
45. Zaichick V (2021) Comparison of Nineteen Chemical Element Contents in Normal Thyroid and Thyroid with Riedel's Struma. *Journal of Medical Research and Health Sciences* 4: 1529-1538.
46. Zaichick V (2021) Evaluation of Twenty Chemical Elements in Thyroid with Hashimoto's thyroiditis using X-Ray Fluorescent and Neutron Activation Analysis. *Journal of Medical Research and Health Sciences* 2:1500-1510.
47. Zaichick V, Zaichick S (2018) Twenty Chemical Element Contents in Normal and Cancerous Thyroid. *Int J Hematol Blo Dis* 3:1-13.
48. Zaichick V, Zaichick S (1996) Instrumental effect on the contamination of biomedical samples in the course of sampling. *The Journal of Analytical Chemistry* 51:1200-1205.
49. Zaichick V, Zaichick S (1997) A search for losses of chemical elements during freeze-drying of biological materials. *J Radioanal Nucl Chem* 218:249-253.
50. Zaichick V (1995) Applications of synthetic reference materials in the medical Radiological Research Centre. *Fresenius J Anal Chem* 352:219-223.
51. Zaichick S, Zaichick V (2011) The Br, Fe, Rb, Sr, and Zn contents and interrelation in intact and morphologic normal prostate tissue of adult men investigated by energy-dispersive X-ray fluorescent analysis. *X-Ray Spectrom* 40:464-469.
52. Zaichick S, Zaichick V (2011) INAA application in the age dynamics assessment of Br, Ca, Cl, K, Mg, Mn, and Na content in the normal human prostate. *J Radioanal Nucl Chem* 288:197-202.
53. Zaichick V, Zaichick S (2013) The effect of age on Br, Ca, Cl, K, Mg, Mn, and Na mass fraction in pediatric and young adult prostate glands investigated by neutron activation analysis. *Appl Radiat Isot* 82:145-151.
54. Zaichick S, Zaichick V (2010) The effect of age and gender on 37 chemical element contents in scalp hair of healthy humans. *Biol Trace Elem Res* 134:41-54.
55. Zaichick S., Zaichick V (2011) The effect of age on Ag, Co, Cr, Fe, Hg, Sb, Sc, Se, and Zn contents in intact human prostate investigated by neutron activation analysis. *Appl Radiat Isot* 69: 827-833.
56. Korelo AM, Zaichick V (1993) Software to optimize the multielement INAA of medical and environmental samples. In: *Activation Analysis in Environment Protection*. Dubna, Russia: Joint Institute for Nuclear Research, Pg: 326-332.
57. Lansdown AB (2007) Critical observations on the neurotoxicity of silver. *Crit Rev Toxicol* 37:237-250.
58. De Vos S, Waegeneers N, Verleysen E, Smeets K, Mast J (2020) Physico-chemical characterisation of the fraction of silver (nano) particles in pristine food additive E174 and in E174-containing confectionery. *Food Addit Contam Part A Chem Anal Control Expo Risk Assess* 37:1831-1846.
59. Hadrup N, Sharma AK, Loeschner K (2018) Toxicity of silver ions, metallic silver, and silver nanoparticles materials after in vivo dermal and mucosal surface exposure: A review. *Regul Toxicol Pharmacol* 98:257-267.
60. Lansdown AB (2006) Silver in health care: antimicrobial effects and safety in use. *Curr Probl Dermatol* 33:17-34.
61. Drake PL, Hazelwood KJ (2005) Exposure-related health effects of silver and silver compounds: a review. *Ann Occup Hyg* 49:575-585.
62. Katarzyńska-Banasik D, Grzesiak M, Kowalik K, Sechman A (2021) Administration of silver nanoparticles affects ovarian steroidogenesis and may influence thyroid hormone metabolism in hens (*Gallus domesticus*). *Ecotoxicol Environ Saf* 208:111427.
63. Leko MB, Gunjača I, Pleić N, Zemunik T (2021) Environmental factors affecting thyroid-stimulating hormone and thyroid hormone levels. *Int J Mol Sci* 22:6521.

64. Schwartz GG, Klug MG (2019) Thyroid Cancer Incidence Rates in North Dakota are Associated with Land and Water Use. *Int J Environ Res Public Health* 16:3805.
65. National Toxicology Program (2005) Toxicology and carcinogenesis studies of sodium chlorate (Cas No. 7775-09-9) in F344/N rats and B6C3F1 mice (drinking water studies). *Natl Toxicol Program Tech Rep Ser* 517:1-255.
66. Parazzini F, Esposito G, Tozzi L, Tozzi S (2017) Epidemiology of endometriosis and its comorbidities. *Eur J Obstet Gynecol Reprod Biol* 209:3-7.
67. Sokal A, Jarmakiewicz-Czaja S, Tabarkiewicz J, Filip R (2021) Dietary intake of endocrine disrupting substances presents in environment and their impact on thyroid function. *Nutrients* 13:867.
68. Zaichick V (1998) X-ray fluorescence analysis of bromine for the estimation of extracellular water. *J Appl Radiat Isot* 49:1165-1169.
69. Leyssens L, Vinck B, Van Der Straeten C, Wuyts F, Maes L (2017) Cobalt toxicity in humans—A review of the potential sources and systemic health effects. *Toxicology* 387:43-56.
70. Yu R (2017) Cobalt Toxicity, An overlooked Cause of Hypothyroidism. *J Endocrinol Thyroid Res* 1:1-4.
71. Simonsen LO, Harbak H, Bennekou P (2012) Cobalt metabolism and toxicology—a brief update. *Sci Total Environ* 432:210-215.
72. Li Y, Trush MA (1993) DNA damage resulting from the oxidation of hydroquinone by copper: role for a Cu(II)/Cu(I) redox cycle and reactive oxygen generation. *Carcinogenesis* 14:1303-1311.
73. Becker TW, Krieger G, Witte I (1996) DNA single and double strand breaks induced by aliphatic and aromatic aldehydes in combination with copper (II). *Free Radic Res* 24:325-332.
74. Glass GA, Stark AA (1997) Promotion of glutathione-gamma-glutamyl transpeptidase-dependent lipid peroxidation by copper and ceruloplasmin: the requirement for iron and the effects of antioxidants and antioxidant enzymes. *Environ Mol Mutagen* 29:73-80.
75. Kim SA, Kwon YM, Kim S, Joung H (2016) Assessment of dietary mercury intake and blood mercury levels in the Korean population: Results from the Korean National Environmental Health Survey 2012–2014. *Int J Environ Res Public Health* 13:877.
76. Clarkson TW, Magos L (2006) The toxicology of mercury and its chemical compounds. *Crit Rev Toxicol* 36:609-662.
77. Correia MM, Chammas MC, Zavariz JD, Arata A, Martins LC et al., (2020) Evaluation of the effects of chronic occupational exposure to metallic mercury on the thyroid parenchyma and hormonal function. *Int Arch Occup Environ Health* 93:491-502.
78. Hu O, Han X, Dong G, Yan W, Wang X et al., (2021) Association between mercury exposure and thyroid hormones levels: A meta-analysis. *Environ Res* 196:110928.
79. Järup L (2003) Hazards of heavy metal contamination. *Br Med Bull* 68:167-182.
80. Malandrino P, Russo M, Ronchi A, Minoia C, Cataldo D (2016) Increased thyroid cancer incidence in a basaltic volcanic area is associated with non-anthropogenic pollution and biocontamination. *Endocrine* 53:471-479.
81. Kant R, Davis A, Verma V (2020) Thyroid nodules: Advances in evaluation and management. *Am Fam Physician* 102:298-304.
82. Leung AM, Braverman LE (2014) Consequences of excess iodine. *Nat Rev Endocrinol* 10:136-142.
83. Lee J-H, Hwang Y, Song R-Y, Yi JW, Yu HW et al., (2017) Relationship between iodine levels and papillary thyroid carcinoma: A systematic review and meta-analysis. *Head Neck* 39:1711-1718.
84. Aakre I, Evensen LT, Kjellefold M, Dahl L, Henjum S et al., (2020) Iodine status and thyroid function in a group of seaweed consumers in Norway. *Nutrients* 12: 3483.
85. Haibach H, Greer MA (1973) Effect of replacement of medium potassium by sodium, cesium or rubidium on in vitro iodide transport and iodoamino acid synthesis by rat thyroid. *Proc Soc Exp Biol Med* 143: 114-117.
86. York DA, Bray GA, Yukimura Y (1978) An enzymatic defect in the obese (ob/ob) mouse: Loss of thyroid-induced sodium- and potassium-dependent adenosinetriphosphatase. *Proc Natl Acad Sci USA* 75: 477-481.
87. Jones JM, Yeralan O, Hines G, Maher M, Roberts DW et al., (1990) Effects of lithium and rubidium on immune responses of rats. *Toxicology Letters* 52: 163-168.
88. Petrini M, Vaglini F, Carulli G, Azzarà A, Ambrogi F et al., (1990) Rubidium is a possible supporting element for bone marrow leukocyte differentiation. *Haematologica* 75: 27-31.
89. Aaseth J, Frey H, Glatte E, Norheim G, Ringstad J et al., (1990) Selenium concentrations in the human thyroid gland. *Biol Trace Elem Res* 24: 147-152.