



## RESEARCH ARTICLE

## COMPARISON OF LEVELS OF TWENTY CHEMICAL ELEMENTS IN NORMAL THYROID TISSUE AND HYPERTROPHIC THYROID TISSUE

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### Abstract

**Background:** Goiter can appear as a palpable or visible enlargement of the thyroid gland at the base of the neck. If the goiter is accompanied by hypothyroidism or hyperthyroidism, it may be accompanied by symptoms of the underlying disorder, and nodular goiter (NG) is a health problem of international importance. The aim of this exploratory study was to assess whether there were significant changes in thyroid tissue levels of twenty chemical elements (ChE) Al, B, Ba, Br, Ca, Cl, Cu, Fe, I, K, Li, Mg, Mn, Na, P, S, Si, Sr, V, and Zn are present in the goitrous transformed thyroid.

**Methods:** Thyroid tissue level of twenty ChE was prospectively evaluated in 46 patients with NG and 105 healthy populations. The measurements were performed using a combination of non-destructive and destructive methods: instrumental neutron activation analysis and inductively coupled plasma atomic emission spectrometry, respectively. Tissue samples were divided into two parts. One was used for morphological study while the other was for ChE analysis.

**Results:** It was found that contents of Al, B, Br, Cl, Cu, Fe, Li, Mg, Mn, Na, P, S, Si, V, and Zn are significantly higher while the I levels are lower in NG than in normal tissues.

**Conclusion:** There are considerable changes in ChE contents in goitrous tissue of thyroid. Thus, it is reasonable to assume that the levels of these ChE in thyroid tissue can be used as NG markers. However, this topic needs additional studies.

**Keywords:** Biomarkers for goiter diagnosis, chemical elements, inductively coupled plasma atomic emission spectrometry, instrumental neutron activation analysis, intact thyroid thyroid nodular goiter.

### INTRODUCTION

At least 10% of the world's population is affected by goiter detected during examination and palpation and most thyroid lesions are nodular goiter (NG)<sup>1</sup>. However, the use of NG ultrasound can be detected in approximately 70% of the general population<sup>2</sup>. NG is also known as endemic nodular goitre, simple goitre, nodular hyperplasia, nontoxic uninodular goitre or multinodular goiter<sup>3</sup>. NG are benign lesions; however, during clinical examination, they can mimic malignancies. NG can be hyper functioning, hypofunctioning, and functioning normally. Euthyroid NG is defined as localized enlargement of the thyroid gland without concomitant disruption of thyroid function<sup>3</sup>.

For more than twenty centuries, there has been a prevailing view that NG is a minor consequence of iodine (I) deficiency. However, NG has been found to be a frequent disease even in those countries and regions where the population is never exposed to I

deficiency<sup>4</sup>. Moreover, it was shown that I excess has severe consequences on human health and associated with the presence of thyroidal dysfunctions and autoimmunity, NG and diffuse goiter, benign and malignant tumors of gland<sup>5-8</sup>. It was also demonstrated that besides the iodine deficiency and excess many other dietary, environmental, and occupational factors are associated with the NG incidence<sup>9-11</sup>. Among them a disturbance of evolutionary stable input of many chemical elements (ChE) in human body after industrial revolution plays a significant role in etiology of thyroidal disorders<sup>12</sup>. Besides iodine involved in thyroid function, ChE has basic physiological functions such as maintaining and regulating cell function, regulating genes, activating or inhibiting enzymatic reactions, and regulating membrane function<sup>13</sup>. The essential or toxic (goitrogenic, mutagenic, and carcinogenic) properties of ChE depend on the tissue-specific need or tolerance, respectively<sup>13</sup>. Excessive accumulation or an imbalance of the ChE may disturb the cell functions and may result in cellular

degeneration, death, benign or malignant transformation<sup>13-15</sup>. 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 I and other ChE contents in the normal and pathological thyroid<sup>16-22</sup>. Level of I in the normal thyroid was investigated in relation to age, gender and some non-thyroidal diseases<sup>23-24</sup>. After that, variations of ChE content with age in the thyroid of males and females were studied and age- and gender-dependence of some ChE was observed<sup>25-41</sup>. Furthermore, a significant difference between some ChE contents in normal and cancerous thyroid was demonstrated<sup>42-47</sup>.

So far, the pathogenesis of NG has to be considered as multifactorial. The present study was performed to clarify the role of twenty ChE in maintaining thyroid growth and goitrogenesis. With this in mind, our aim was to assess the aluminum (Al), boron (B), barium (Ba), bromine (Br), calcium (Ca), chlorine (Cl), copper (Cu), iron (Fe), I, potassium (K), lithium (Li), magnesium (Mg), manganese (Mn), sodium (Na), phosphorus (P), sulfur (S), silicon (Si), strontium (Sr), vanadium (V), and zinc (Zn) mass fraction contents in NG tissue using a combination of non-destructive and destructive methods: instrumental neutron activation analysis with high resolution spectrometry of short-lived radionuclides (INAA-SLR) and inductively coupled plasma atomic emission spectrometry (ICPAES), respectively. A further aim was to compare the levels of these twenty ChE in the goitrous thyroid with those in normal gland of apparently healthy persons.

## SUBJECTS AND METHODS

### Samples

All patients suffered from NG (n=46, mean age  $M \pm SD$  was  $48 \pm 12$  years, range 30-64) were hospitalized in the Head and Neck Department of the Medical Radiological Research Centre and informed consent was taken from the subjects. Thick-needle 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 ChE 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  $44 \pm 21$  years, range 2-87), who had died suddenly. Samples were obtained within 48 hours after a sudden death. The majority of deaths were due to trauma. Histological examination was used in the control group to match the age criteria, as well as to confirm the absence of micro-nodules and underlying cancer.

### Sample preparation, instrumentation and analytical methods

All tissue samples were divided into two portions using a titanium scalpel<sup>48</sup>. One was used for morphological study while the other was intended for ChE analysis. After the samples intended for ChE analysis were

weighed, they were freeze-dried and homogenized<sup>49</sup>. The pounded samples weighing about 5-10 mg (for biopsy) and 100 mg (for resected materials) were used for ChE measurement by INAA-SLR. The samples for INAA-SLR were sealed separately in thin polyethylene films washed beforehand with acetone and rectified alcohol. The sealed samples were placed in labeled polyethylene ampoules. 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).

After INAA-SLR investigation the thyroid samples were taken out from the polyethylene ampoules and used for ICP-AES. The samples were decomposed in autoclaves. Sample aliquots were used to determine the Al, B, Ba, Ca, Cu, Fe, K, Li, Mg, Mn, Na, P, S, Si, Sr, V, and Zn mass fractions by ICP-AES using the Spectrometer ICAP-61 (Thermo Jarrell Ash, USA). The determination of the ChE content in aqueous solutions was made by the quantitative method using calibration solutions (High Purity Standards, USA) of 0.5 and 10 mg/L of each element. The calculations of the ChE content in the probe were carried out using software of a spectrometer (ThermoSPEC, version 4.1). Information detailing with the NAA-SLR and ICP-AES methods used and other details of the analysis were presented in our earlier publications concerning ChE contents in human thyroid, scalp hair, and prostate<sup>33,34,50-55</sup>.

### Standards and certified reference material

To determine contents of the ChE by comparison with a known standard, biological synthetic standards (BSS) prepared from phenol-formaldehyde resins were used<sup>56</sup>. In addition to BSS, aliquots of commercial, chemically pure compounds were also used as standards. Ten sub-samples of certified reference material (CRM) IAEA H-4 (animal muscle) and five sub-samples of CRM of the Institute of Nuclear Chemistry and Technology (INCT, Warszawa, Poland) INCT-SBF-4 Soya Bean Flour, INCT-TL-1 Tea Leaves, and INCT-MPH-2 Mixed Polish Herbs were treated and analyzed in the same conditions that thyroid samples to estimate the precision and accuracy of results. More details about the quality controls of ChE contents in human thyroid were presented in our earlier publications<sup>33,34</sup>.

### Computer programs and statistic

A dedicated computer program for INAA mode optimization was used<sup>56</sup>. All thyroid samples were prepared in duplicate, and mean values of ChE contents were used. Mean values of ChE contents were used in final calculation for the Br, Fe, Rb, and Zn mass fractions measured by two methods. 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 ChE contents. The difference in the results between two groups (normal and NG) was evaluated by the parametric Student's *t*-test and non-parametric Wilcoxon-Mann-Whitney *U*-test. All studies were approved by the Ethical Committees of the Medical Radiological Research Centre (MRRC), Obninsk

(Reference number 115050610007, year2017). 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.

**Table 1: Some statistical parameters of Al, B, Ba, Br, Ca, Cl, Cu, Fe, I, K, Li, Mg, Mn, Na, P, S, Si, Sr, V, and Zn mass fraction (mg/kg, dry mass basis) in normal and goitrous thyroid.**

Tissue	Element	Mean	SD	SEM	Min	Max	Median	P 0.025	P 0.975
Normal n=105	Al	10.5	13.4	1.8	0.800	69.3	6.35	1.19	52.9
	B	0.476	0.434	0.058	0.200	2.30	0.300	0.200	1.73
	Ba	1.12	1.15	0.15	0.0480	5.00	0.680	0.0838	4.48
	Br	14.9	11.0	1.2	1.90	54.1	11.6	2.56	49.3
	Ca	1682	999	106	373	5582	1454	444	4183
	Cl	3400	1452	174	1030	6000	3470	1244	5869
	Cu	4.08	1.22	0.14	0.500	7.15	4.10	1.57	6.41
	Fe	223	95	10	52.0	489	210	72.8	432
	I	1841	1027	107	114	5061	1695	230	4232
	K	6418	2625	290	1914	15293	5948	2947	13285
	Li	0.0208	0.0155	0.0022	0.0015	0.0977	0.0178	0.0041	0.0487
	Mg	296	134	16	66.0	930	284	95.8	541
	Mn	1.28	0.56	0.07	0.470	4.04	1.15	0.537	2.23
	Na	6928	1730	175	3686	13453	6835	3974	10709
	P	4290	1578	207	496	8996	4221	1360	7323
	S	8259	2002	263	644	11377	8399	3662	11208
	Si	50.8	46.9	6.2	5.70	180	36.0	7.11	174
	Sr	3.81	2.93	0.34	0.100	12.6	2.90	0.365	11.3
	V	0.102	0.039	0.005	0.0200	0.250	0.100	0.0440	0.192
Zn	94.8	39.6	4.2	7.10	215	88.5	34.9	196	
Goiter n=46	Al	27.1	24.7	5.3	6.60	95.1	20.5	6.92	85.2
	B	1.71	1.19	0.26	0.90	5.00	1.00	0.95	5.00
	Ba	1.43	1.75	0.37	0.18	8.20	0.96	0.238	5.79
	Br	36.3	31.3	6.99	8.0	131	26.6	8.95	110
	Ca	1422	834	164	288	4333	1272	362	3219
	Cl	9117	3866	1223	4226	16786	8259	4504	15869
	Cu	8.51	7.15	1.60	2.90	34.8	5.95	3.00	26.2
	Fe	337	321	51	62.0	1350	199	65.0	1214
	I	1310	1433	221	29.0	8260	974	107	3713
	K	6610	2233	430	3353	12222	6110	3395	10984
	Li	0.0281	0.0117	0.0030	0.0073	0.0541	0.0259	0.0089	0.0530
	Mg	356	119	23	63.0	612	371	149	559
	Mn	1.77	1.13	0.23	0.450	5.50	1.60	0.516	4.12
	Na	11782	4342	836	7229	28481	10697	7279	20921
	P	5181	1798	383	2890	9637	5030	2919	8827
	S	10961	2091	446	5591	14970	10719	6824	14579
	Si	81.3	57.3	12.5	7.80	182	69.9	12.0	178
	Sr	5.87	8.42	1.59	0.93	32.0	2.26	1.11	31.5
	V	0.152	0.074	0.016	0.043	0.370	0.150	0.056	0.310
Zn	120.5	50.8	7.8	47.0	264	113	49.1	257	

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.

## RESULTS

Table 1 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 Al, B, Ba, Br, Ca, Cl, Cu, Fe, I, K, Li, Mg, Mn, Na, P, S, Si, Sr, V, and Zn mass fraction in normal and goitrous thyroid. The comparison of our results with published data for Al, B, Ba, Br, Ca, Cl, Cu, Fe, I, K, Li, Mg, Mn, Na, P, S, Si, Sr, V, and Zn mass fraction in normal and goitrous thyroid<sup>58-91</sup> is shown in Table 2. The ratios of means and the difference between mean values of Al, B, Ba, Br, Ca, Cl, Cu, Fe, I, K, Li, Mg, Mn, Na, P, S,

Si, Sr, V, and Zn mass fractions in normal and goitrous thyroid are presented in Table 3.

## DISCUSSION

### Precision and accuracy of results

As was shown before<sup>33,34</sup>, good agreement of our results for the Al, B, Ba, Br, Ca, Cl, Cu, Fe, I, K, Mg, Mn, Na, P, S, Sr, V, and Zn mass fractions with the certified values of CRM IAEA H-4, INCT-SBF-4, INCT-TL-1, and INCT-MPH-2 as well as the similarity of the means of the Ca, K, Mg, Mn, and Na mass fractions in the normal human thyroid determined by both INAA-SLR and ICP-AES methods demonstrates acceptable precision and accuracy of the results

obtained in the study and presented in Tables 1, Table 2 and Table 3. The mean values and all selected statistical parameters were calculated for twenty ChE (Al, B, Ba, Br, Ca, Cl, Cu, Fe, I, K, Li, Mg, Mn, Na, P, S, Si, Sr, V, and Zn) mass fractions (Table 1). The

mass fraction of Al, B, Ba, Br, Ca, Cl, Cu, Fe, I, K, Li, Mg, Mn, Na, P, S, Si, Sr, V, and Zn were measured in all, or a major portion of normal and goitrous tissue samples.

**Table 2: Median, minimum and maximum value of means Al, B, Ba, Br, Ca, Cl, Cu, Fe, I, K, Li, Mg, Mn, Na, P, S, Si, Sr, V, and Zn contents in the normal and goitrous thyroid according to data from the literature in comparison with our results (mg/kg, dry mass basis).**

Tissue	El	Published data [Reference]			This work
		Median of means (n)*	Min of means M or M±SD, (n)**	Max of means M or M±SD, (n)**	
Normal	Al	33.6 (12)	0.33 (-) <sup>58</sup>	420 (25) <sup>59</sup>	10.5±13.4
	B	0.151 (2)	0.084 (3) <sup>60</sup>	0.46 (3) <sup>60</sup>	0.476±0.434
	Ba	0.67 (7)	0.0084 (83) <sup>61</sup>	≤5.0 (16) <sup>62</sup>	1.12±1.15
	Br	18.1 (11)	5.12 (44) <sup>63</sup>	284±44 (14) <sup>64</sup>	16.3±11.6
	Ca	1600 (17)	840±240 (10) <sup>65</sup>	3800±320 (29) <sup>65</sup>	1663±999
	Cl	6800 (5)	804±80 (4) <sup>66</sup>	8000 (-) <sup>67</sup>	3400±1452
	Cu	6.0 (61)	0.16 (83) <sup>61</sup>	220±22 (10) <sup>66</sup>	3.93±1.43
	Fe	252 (21)	56 (120) <sup>68</sup>	3360 (25) <sup>59</sup>	223±95
	I	1888 (95)	159±8 (23) <sup>69</sup>	5772±2708 (50) <sup>70</sup>	1841±1027
	K	4300 (17)	46.4±4.8 (4) <sup>66</sup>	6090 (17) <sup>62</sup>	6418±2625
	Li	6.3 (2)	0.092 (-) <sup>71</sup>	12.6 (180) <sup>72</sup>	0.0208±0.0154
	Mg	390 (16)	3.5 (-) <sup>58</sup>	1520 (20) <sup>73</sup>	296±134
	Mn	1.62 (40)	0.076 (83) <sup>61</sup>	69.2±7.2 (4) <sup>66</sup>	1.28±0.56
	Na	8000 (9)	438 (-) <sup>74</sup>	10000±5000 (11) <sup>75</sup>	6928±1730
	P	2860 (10)	16 (7) <sup>76</sup>	7520 (60) <sup>63</sup>	4290±1578
	S	11000 (3)	4000 (-) <sup>67</sup>	11800 (44) <sup>63</sup>	8259±2002
	Si	16.0 (3)	0.97 (-) <sup>58</sup>	143±6 (40) <sup>77</sup>	50.8±46.9
	Sr	0.61 (9)	0.055 (83) <sup>61</sup>	46.8±4.8(4) <sup>66</sup>	3.81±2.93
	V	0.065 (6)	0.0124 (2) <sup>78</sup>	18±2 (4) <sup>66</sup>	0.102±0.039
	Zn	110 (56)	2.1(-) <sup>58</sup>	820±204 (14) <sup>64</sup>	94.8±39.7
Goiter	Al	3.84 (6)	2.45 (123) <sup>79</sup>	840 (25) <sup>59</sup>	27.1±24.7
	B	-	-	-	1.71±1.19
	Ba	4.92 (1)	4.92±4.56 (51) <sup>80</sup>	4.92±4.56 (51) <sup>80</sup>	1.43±1.75
	Br	480 (4)	9 (5) <sup>81</sup>	777 (1) <sup>82</sup>	36.3±31.3
	Ca	3168 (8)	600 (1) <sup>81</sup>	9200 (1) <sup>81</sup>	1422±834
	Cl	-	-	-	9117±3866
	Cu	10.0 (33)	0.84 (1) <sup>72</sup>	353 (101) <sup>83</sup>	8.51±7.15
	Fe	390 (5)	128±52 (13) <sup>84</sup>	4848±3056 (11) <sup>64</sup>	337±321
	I	770 (44)	52 (1) <sup>85</sup>	2800 (4) <sup>86</sup>	1310±1433
	K	3725 (4)	276 (75) <sup>87</sup>	6030±620 (-) <sup>88</sup>	6610±2233
	Li	-	-	-	0.0281±0.0117
	Mg	834 (4)	588±388 (13) <sup>84</sup>	1616 (70) <sup>73</sup>	356±119
	Mn	2.64 (21)	0.352 (130) <sup>89</sup>	34.9 (101) <sup>90</sup>	1.77±1.13
	Na	3360 (1)	3360 (25) <sup>59</sup>	3360 (25) <sup>59</sup>	11782±4342
	P	8200 (1)	8200±280 (-) <sup>88</sup>	8200±280 (-) <sup>88</sup>	5181±1798
	S	10300 (1)	10300±340 (-) <sup>88</sup>	10300±340 (-) <sup>88</sup>	10961±2091
	Si	64 (1)	45 (122) <sup>88</sup>	114 (122) <sup>88</sup>	81.3±57.3
	Sr	1.45 (2)	1.26 (25) <sup>59</sup>	1.64 (51) <sup>80</sup>	5.87±8.42
	V	3.92 (1)	3.92±8.84 (51) <sup>80</sup>	3.92±8.84 (51) <sup>80</sup>	0.152±0.074
	Zn	146 (25)	22.4 (130) <sup>89</sup>	1236±560 (2) <sup>91</sup>	120.5±50.8

El- element, M- arithmetic mean, SD- standard deviation, (n)\*- number of all references, (n)\*\*- number of samples.

**Comparison with published data**

The means obtained for Al, B, Ba, Br, Ca, Cl, Cu, Fe, I, K, Mg, Mn, Na, P, S, Si, Sr, V, and Zn mass fraction, as shown in Table 2, agree well with the medians of mean values reported by other researches for the human thyroid, including samples received from persons who died from different non-thyroid diseases. The mean obtained for Li is two orders of magnitude lower than the median of previously reported data.

Moreover, it is outside the range of previously reported means. A number of values for ChE mass fractions were not expressed on a dry mass basis by the authors of the cited references. Hence we calculated these values using published data for water 75%<sup>92</sup> and ash 4.16% on dry mass basis<sup>93</sup> contents in thyroid of adults. In goitrous tissues our results for Al, Br, Ca, Cu, Fe, I, Mn, Si, and Zn contents were within the range of published means, while means for K and Sr were some

higher median of previously reported means and also higher the upper level of the range of these means (Table 2). Only one published article on Ba<sup>79</sup>, Na<sup>59</sup>, P<sup>88</sup>, S<sup>88</sup>, Si<sup>88</sup>, and V<sup>79</sup> contents in the goitrous tissue samples was found in the literature. The mean obtained in the present study for S content in the goitrous tissue agreed well with early published data, while means for

Ba and P were some lower and the mean for Na was some higher. The obtained mean for V content in the goitrous tissue was more than one order of magnitude lower than the only reported result. No published data referring B, Cl, and Li contents of goitrous thyroid tissue were found.

**Table 3: Differences between mean values (M±SEM) of Al, B, Ba, Br, Ca, Cl, Cu, Fe, I, K, Li, Mg, Mn, Na, P, S, Si, Sr, V, and Zn mass fraction (mg/kg, dry mass basis) in normal and goitrous thyroid.**

Element	Thyroid tissue		Student's t-test <i>p</i> ≤	U-test <i>p</i>	Ratio Goiter to Norm
	Norm n=105	Goiter n=46			
Al	10.5±1.8	27.1±5.3	<b>0.0057</b>	≤0.01	2.58
B	0.476±0.058	1.71±0.26	<b>0.00013</b>	≤0.01	3.59
Ba	1.12±0.15	1.43±0.37	0.446	>0.05	1.28
Br	14.9±1.2	36.3±6.99	<b>0.0067</b>	≤0.01	2.44
Ca	1682±106	1422±164	0.188	>0.05	0.84
Cl	3400±174	9117±1223	<b>0.0011</b>	≤0.01	2.68
Cu	4.08±0.14	8.51±1.60	<b>0.012</b>	≤0.01	2.09
Fe	223±10	337±51	<b>0.034</b>	≤0.01	1.51
I	1841±107	1310±221	<b>0.035</b>	≤0.01	0.71
K	6418±290	6610±430	0.713	>0.05	1.03
Li	0.0208±0.0022	0.0281±0.0030	<b>0.037</b>	≤0.01	1.35
Mg	296±16	356±23	<b>0.037</b>	≤0.01	1.20
Mn	1.28±0.07	1.77±0.23	<b>0.048</b>	≤0.01	1.38
Na	6928±175	11782±836	<b>0.000041</b>	≤0.01	1.70
P	4290±207	5181±383	<b>0.049</b>	≤0.05	1.21
S	8259±263	10961±446	<b>0.000074</b>	≤0.01	1.33
Si	50.8±6.2	81.3±12.5	<b>0.037</b>	≤0.01	1.60
Sr	3.81±0.34	5.87±1.59	0.216	>0.05	1.54
V	0.102±0.005	0.152±0.016	<b>0.0072</b>	≤0.01	1.49
Zn	94.8±4.2	120.5±7.8	<b>0.0053</b>	≤0.01	1.27

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

The range of means of Al, B, Ba, Br, Ca, Cl, Cu, Fe, I, K, Li, Mg, Mn, Na, P, S, Si, Sr, V, and Zn level reported in the literature for normal and for goitrous thyroid varies widely (Table 2). This can be explained by a dependence of ChE on many factors, including the region of the thyroid, from which the sample was taken, age, gender, ethnicity, gland mass, and the NG stage. Not all these factors were strictly controlled in cited studies. Another and, in our opinion, the main reason for the inter-observer discrepancy can be attributed to the accuracy of the analytical techniques, sample preparation methods, and the inability to take standardized samples from affected tissues. It was insufficient quality control of results in these studies. In many reported papers 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 by use of these sample preparation methods some quantities of certain ChE are lost as a result of this treatment That concern not only such volatile halogen as Br, but also other ChE investigated in the study<sup>94-96</sup>.

#### Effect of goitrous transformation on ChE contents

From Table 3, it is observed that in goitrous tissue the mass fraction of Al, B, Br, Cl, and Cu are approximately 2.6, 3.6, 2.4, 2.7, and 2.1 times, respectively, higher and also mass fractions of Fe, Li, Mg, Mn, Na, P, S, Si, V, and Zn are almost in 51%,

35%, 20%, 38%, 70%, 21%, 33%, 60%, 49%, and 27% respectively, higher than in normal tissues of the thyroid. In contrast, the mass fraction of I is 29% significantly lower. Thus, if we accept the ChE contents in thyroid glands in the control group as a norm, we have to conclude that with a goitrous transformation the levels of Al, B, Br, Cl, Cu, Fe, Li, Mg, Mn, Na, P, S, Si, V, and Zn in thyroid tissue significantly increased, whereas the level of I some decreased.

#### Role of ChE in goitrous transformation of the thyroid

Characteristically, elevated or reduced levels of ChE observed in goitrous tissues are discussed in terms of their potential role in the initiation and promotion of thyroid goiter. In other words, using the low or high levels of the ChE in goitrous tissues researchers try to determine the goitrogenic role of the deficiency or excess of each ChE in investigated organ. In our opinion, abnormal levels of many ChE 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 ChE level in pathologically altered tissue is the reason for alterations or vice versa.

#### Aluminum

The trace element Al is not described as essential, because there is no biochemical function directly associated with it. At this point in our knowledge, there

is no doubt that AI overload negatively affects human health, including thyroid function<sup>97</sup>.

#### **Boron**

Trace element B is known to influence the activity of many enzymes<sup>98</sup>. Numerous studies have demonstrated beneficial effects of B on human health, including anti-inflammatory stimulus-reduces levels of inflammatory biomarkers, such as high-sensitivity C-reactive protein (hs-CRP) and tumor necrosis factor  $\alpha$  (TNF- $\alpha$ ); as well as raises levels of antioxidant enzymes, such as superoxide dismutase (SOD), catalase, and glutathione peroxidase<sup>98</sup>. Why B content in goitrous thyroid is higher than normal level and how an excess of B acts on thyroid are still to be cleared.

#### **Bromine**

This is one of the most abundant and ubiquitous of the recognized trace elements in the biosphere. The inorganic bromide is the ionic form of Br that exerts therapeutic as well as toxic effects. An enhanced intake of bromide could interfere with the metabolism of I at the whole-body level. In the thyroid gland the biological behavior of bromide is more similar to the biological behavior of iodide<sup>99</sup>. In our previous studies, we found a significant age-related increase of Br content in human thyroid<sup>27,28,30-33</sup>. Therefore, a goitrogenic and, probably, carcinogenic effect of excessive Br levels in the thyroid of old females was assumed. On the one hand, elevated levels of Br in NG tissues, observed in the present study, supports this conclusion. But, on the other hand, bromide compounds, especially potassium bromide (KBr), sodium bromide (NaBr), and ammonium bromide (NH<sub>4</sub>Br), are frequently used as sedatives in Russia<sup>101</sup>. It may be the reason for elevated levels of Br in specimens of patients with NG.

#### **Chlorine**

Cl is a ubiquitous, extracellular electrolyte essential to more than one metabolic pathway. Cl exists in the ionic form (chloride) in the human body. In the body, it is mostly present as sodium chloride. Therefore, as usual, there is a correlation between Na and Cl contents in tissues and fluids of human body. It is well known that Cl mass fractions in samples depend mainly on the extracellular water volume, including the blood volumes, in tissues<sup>102</sup>. NG tissues are predominantly highly vascularized lesions<sup>103</sup>. Thus, it is possible to speculate that thyroid goiters are characterized by an increase of the mean value of the Cl mass fraction because the level of goiter vascularization is higher than that in normal thyroid tissue.

#### **Coper**

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-monooxygenase, extracellular superoxide dismutase, lysyl oxidase, peptidyl glycine amidating monooxygenase, 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<sup>104</sup>. Thus, Cu accumulation in thyroid parenchyma with age may be involved in oxidative stress, dwindling gland function, and increasing risk of goiter/cancer<sup>25,26,31,34</sup>. The significantly elevated level of Cu in goitrous thyroid, 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 the thyroid goiter etiology and metabolism.

#### **Iron**

It is well known that Fe as a trace element 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<sup>104</sup>. 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<sup>25,26,29-34</sup>. Most experimental and epidemiological data support the hypothesis that Fe overload is a risk factor for benign and malignant tumors<sup>106</sup>. This goitrogenic and oncogenic effect could be explained by an overproduction of ROS and free radicals<sup>107</sup>.

#### **Iodine**

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. Goitrous transformation is accompanied by a partial loss of tissue-specific functional features, which leads to a significant reduction in I content associated with functional characteristics of the human thyroid tissue.

#### **Lithium**

The results of lifelong Li-poor nutrition of animals show that Li is essential to the fauna, and thus, to humans as well<sup>108</sup>. Li-poor nutrition has a negative influence on some enzyme activity, mainly the enzymes of the citrate cycle, glycolysis, and of nitrogen metabolism<sup>108</sup>. On the other hand, Li is widely used in medicine as a mood-stabilizing drug. Because of the active transport of Na<sup>+</sup>/I<sup>-</sup> ions, Li is accumulated in the thyroid gland at a concentration 3-4 times higher than that in the plasma. It can inhibit the formation of colloid in thyrocytes, change the structure of thyroglobulin, weaken the iodination of tyrosines, and disrupt their coupling<sup>109</sup>. In addition, it reduces the clearance of free thyroxine in the serum, thereby indirectly reducing the activity of 5-deiodinase type 1 and 2 and reducing the deiodination of these hormones in the liver<sup>108</sup>. All these actions may cause the development of goiter.

#### **Magnesium**

Mg is abundant in the human body. This element is essential for the functions of more than 300 enzymes (e.g. alkaline phosphatases, ATP-ases, phosphokinases, the oxidative phosphorylation pathway). It plays a crucial role in many cell functions such as energy metabolism, protein and DNA syntheses, and cytoskeleton activation. Moreover, Mg is involved in

the thyroid function and plays a central role in determining the clinical picture associated with thyroid disease<sup>110</sup>. The higher Mg levels in NG than do normal tissues, possibly is a result of the high Mg requirement of growing cells.

#### **Manganese**

Trace element Mn is a cofactor for numerous enzymes, playing many functional roles in living organisms. The Mn containing enzyme, manganese superoxide dismutase (Mn-SOD), is the principal antioxidant enzyme which neutralizes the toxic effects of reactive oxygen species. It has been speculated that Mn interferes with thyroid hormone binding, transport, and activity at the tissue level<sup>111</sup>. However, an overall comprehension of Mn homeostasis and physiology, which is not yet acquired, is mandatory to establish Mn exact role in the thyroid goiter etiology and metabolism.

#### **Sodium**

Knowledge concerning ion regulation in many normal and abnormal cell processes has had a rapid development. It was found, among other regulations, that sodium-calcium exchange is associated with the cytoskeleton and the cell membrane. A hypothesis was eventually established that a wide variety of pathological phenomena ranging from acute cell death to chronic processes, such as neoplasia, all have a common series of cellular reactions<sup>112</sup>. Furthermore, iodide (I<sup>-</sup>), an essential constituent of the thyroid hormones, is actively transported into the thyroid via the Na<sup>+</sup>/I<sup>-</sup> symporter (NIS), a key plasma membrane glycoprotein<sup>113</sup>. In addition, Na is mainly an extracellular electrolyte and its elevated level in NG might link with a higher goiter vascularization in comparison with the normal thyroid (see *Chlorine*).

#### **Phosphorus**

P is necessary for several, various biological roles in the signal transduction of cells and energy exchange of human body. About 80–90% of P is founded in teeth and bones in the form of hydroxyapatite. Thyroid hormones play an important role in homeostasis of Ca and P levels by their direct action on bone turnover and, as a consequence, Ca and P metabolism is frequently disturbed in thyroid dysfunction with a significant increase in the P serum levels<sup>114</sup>. The elevated level of P in serum results the higher content of this element in NG tissue, because the goiter vascularization is higher in comparison with the normal thyroid. Besides, the elevated level of thyroid phospholipids in NG is common<sup>115</sup>.

#### **Sulfur**

Proteins contain between 3 and 6% of sulfur amino acids. Sulfur amino acids contribute substantially to the maintenance and integrity of the cellular systems by influencing the cellular redox state and the capacity to detoxify toxic compounds, free radicals and reactive oxygen species (ROS)<sup>115</sup>. ROS are generated during normal cellular activity and may exist in excess in some pathophysiological conditions, such as inflammation. Therefore exploring fundamental aspects of sulfur metabolism such as the antioxidant effects of sulfur-containing amino acids<sup>116</sup> may help elucidate the mechanism by which the S content increases in NG.

Thus, it might be assumed that the elevated S level in goitrous thyroid reflects an increase in concentration of ROS in goiter tissue.

#### **Silicon**

Si as a trace element is essential to some specific biological functions in humans<sup>118</sup>. For example, Si is necessary for the association between cells and one or more macromolecules such as osteonectin, which affects cartilage composition and ultimately cartilage calcification<sup>119</sup>. However, an association between the disorders of thyroid function and the Si excess in the diets was found<sup>120</sup>. An increase in the thyrotropin (TSH) level in rats was observed after Si-treatment, without statistically significant differences in thyroid hormones concentrations between the test and control groups of animals<sup>121</sup>.

#### **Vanadium**

V complexes are cofactors for several enzymes that maintaining hemostasis in health and pathology. For example, V compounds normalized blood pressure, ischemia and the metabolism of the thyroid<sup>122</sup>. However, all V compounds have been considered toxic and a goitrogenic and carcinogenic role of V on the thyroid was proposed<sup>123</sup>. V compounds promote the induction and perpetuation of an inflammatory reaction in the thyroid<sup>123</sup>. Thus, the elevated V level in thyroid may be a cause of the gland dysfunctions, NG and cancer.

#### **Zinc**

Zn is active in more than 300 proteins and over 100 DNA-binding proteins, including the tumor suppressor protein p53, a Zn-binding transcription factor acting as a key regulator of cell growth and survival upon various forms of cellular stress. p53 is mutated in half of human tumors and its activity is tightly regulated by metals and redox mechanisms. On the other hand, excessive intracellular Zn concentrations may be harmful to normal metabolism of cells<sup>124</sup>. By now much data has been obtained related both to the direct and indirect action of intracellular Zn on the DNA polymeric organization, replication and lesions, and to its vital role for cell division<sup>125,126</sup>. Other actions of Zn have been also described. They include its action as a potent anti-apoptotic agent<sup>127-131</sup>. All these facts allowed us to speculate that age-related overload Zn content in female thyroid, as was found in our previous study<sup>25,29,31,33</sup>, is probably one of the factors in etiology of thyroid goiter and malignant tumors. Therefore, the elevated Zn level in NG in comparison with normal level, detected in this study, supports our hypothesis.

Our findings show that mass fraction of Al, B, Br, Cl, Cu, Fe, I, Li, Mg, Mn, Na, P, S, Si, V, and Zn are significantly different in NG as compared to normal thyroid tissues (Tables 6). Thus, it is plausible to assume that levels of these ChE in thyroid tissue can be used as NG markers.

#### **Limitations of the study**

This study has several limitations. Firstly, analytical techniques employed in this study measure only twenty ChE (Al, B, Ba, Br, Ca, Cl, Cu, Fe, I, K, Li, Mg, Mn, Na, P, S, Si, Sr, V, and Zn) mass fractions. Future studies should be directed toward using other analytical methods which will extend the list of ChE 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 ChE contents in NG group using differentials like gender, histological types of goiter, 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 Al, B, Br, Cl, Cu, Fe, I, Li, Mg, Mn, Na, P, S, Si, V, and Zn level alteration and shows the necessity to continue ChE research of goitrous thyroid.

## CONCLUSIONS

In this work, ChE measurements in tissue samples from normal and NG colloid thyroid were performed using two useful analytical methods: non-destructive neutron activation analysis with high-resolution short-lived radionuclide spectrometry and inductively coupled plasma atomic emission spectrometry. The combination of these methods has been shown to be a suitable analytical tool for the determination of twenty ChEs in tissue samples from healthy and affected human thyroid, including needle biopsy samples. It was observed that the content of goitrous tissues of Al, B, Br, Cl, Cu, Fe, Li, Mg, Mn, Na, P, S, Si, V and Zn increased significantly while the level of I decreased in comparison with normal thyroid tissues. In our opinion, the presented study data strongly suggest that ChE plays an important role in thyroid health and the etiology of colloidal NG. It was assumed that the differences in ChE levels in affected thyroid tissue could be used as colloidal NG markers.

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## AUTHOR'S CONTRIBUTION

**Zaichick V:** Writing original draft, review, literature survey, editing, methodology, data curation.

## DATA AVAILABILITY

Data will be made available on reasonable request.

## CONFLICT OF INTEREST

None to declare.

## REFERENCES

1. Carlé A, Krejbjerg A, Laurberg P. Epidemiology of nodular goitre. Influence of iodine intake. *Best Pract Res Clin Endocrinol Metab* 2014; 28(4):465-479. <https://doi.org/10.1016/j.beem.2014.01.001>
2. Kant R, Davis A, Verma V. Thyroid nodules: Advances in evaluation and management. *Am Fam Physician* 2020; 102(5):298-304. PMID: 32866364
3. Hoang VT, Trinh CT. A review of the pathology, diagnosis and management of colloid goitre. *Eur Endocrinol. Eur Endocrinol* 2020; 16(2):131-135. <https://doi.org/10.17925/EE.2020.16.2.131>
4. 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. <https://doi.org/10.1053/beem.2000.0104>
5. Zaichick V. Iodine excess and thyroid cancer. *J Trace Elem Exp Med* 1998; 11(4):508-509.
6. Zaichick V, Iljina T. Dietary iodine supplementation effect on the rat thyroid 131I blastomogenic action. In: *Die Bedeutung der Mengen- und Spurenelemente*. 18. Arbeitstangung. Jena: Friedrich-Schiller-Universität; 1998: 294-306.
7. 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 (2013-2015). *Nutrients* 2019; 11(11):2757. <https://doi.org/10.3390/nu11112757>
8. Vargas-Uricoechea P, 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; 6239243. <https://doi.org/10.1155/2019/6239243>
9. Stojavljević A, Rovčanin B, Krstić D, *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. <https://doi.org/10.1016/j.jtemb.2019.06.009>
10. 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):200903. <https://doi.org/10.5696/2156-9614-10.27.200903>
11. Liu M, Song J, Jiang Y, *et al*. A case-control study on the association of mineral elements exposure and thyroid tumor and goiter. *Ecotoxicol Environ Saf* 2021; 208:111615. <https://doi.org/10.1016/j.ecoenv.2020.111615>
12. 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 ml U/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. <https://doi.org/10.1016/j.bbacli.2017.03.003>
14. Beyersmann D, Hartwig A. Carcinogenic metal compounds: recent insight into molecular and cellular mechanisms. *Arch Toxicol* 2008; 82(8):493-512. <https://doi.org/10.1007/s00204-008-0313-y>
15. Martinez-Zamudio R, Ha HC. Environmental epigenetics in metal exposure. *Epigenetics* 2011; 6(7):820-827. <https://doi.org/10.4161/epi.6.7.16250>
16. Zaichik VE, RaibukhinYuS, 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. PMID: 5449249
17. Zaichik VE, Matveenko EG, Vtiurin BM, Medvedev VS. Intrathyroid iodine in the diagnosis of thyroid cancer. *Vopr Onkol* 1982;28(3):18-24.PMID: 7064415
18. Zaichick V, Tsyb AF, Vtyurin BM. Trace elements and thyroid cancer. *Analyst* 1995; 120(3):817-821. <https://doi.org/10.1039/an9952000817>
19. Zaichick VYe, Choporov Yu Ya. Determination of the natural level of human intra-thyroid iodine by instrumental neutron activation analysis. *J Radio 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.



21. Zaichick V, Zaichick S. Energy-dispersive X-ray fluorescence of iodine in thyroid punctures 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.  
<https://doi.org/10.1111/j.1749-6632.2000.tb06530.x>
23. Zaichick V, Zaichick S. Normal human intrathyroidal iodine. *Sci Total Environ* 1997; 206(1):39-56.  
[https://doi.org/10.1016/s0048-9697\(97\)00215-5](https://doi.org/10.1016/s0048-9697(97)00215-5)
24. Zaichick V. Human intrathyroidal iodine in health and non-thyroidal disease. In: *New aspects of trace element research*. London: Smith-Gordon; and Tokyo:Nishimura; 1999:114-119.
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(1):31-38. <https://doi.org/10.36959/452/579>
26. Zaichick V, Zaichick S. Age-related changes of some trace element contents in intact thyroid of males investigated by energy dispersive X-ray fluorescent analysis. *MOJ Gerontol Ger* 2017; 1(5):00028.  
<https://doi.org/10.15406/mojgg.2017.01.00028>
27. 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; 1:5.1.
28. 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.
29. Zaichick V, Zaichick S. 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* 2017;3:015.  
<https://doi.org/10.24966/GGM-8662/100015>
30. Zaichick V, Zaichick S. 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* 2017;4(4):555644.  
<https://doi.org/10.19080/CTBEB.2017.04.555644>
31. Zaichick V, Zaichick S. Effect of age on chemical element contents in female thyroid investigated by some nuclear analytical methods. *Micro Medicine* 2018;6(1):47-61.
32. Zaichick V, Zaichick S. Neutronactivation 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.  
<https://doi.org/10.32474/OAJBEB.2018.02.000144>
33. 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. <https://doi.org/10.16966/2576-5833.114>
34. 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.  
<https://doi.org/10.36876/smggr.1014>
35. 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.  
<https://doi.org/10.1007/s40520-018-0906-0>
36. 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.
37. Zaichick V, Zaichick S. Possible role of inadequate quantities of intra-thyroidal bromine, calcium and magnesium in the etiology of female subclinical hypothyroidism. *IntGyn and Women's Health* 2018;1(3):IGWHC.MS.ID.000113.  
<https://doi.org/10.32474/IGWHC.2018.01.000113>
38. 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):000108.  
<https://doi.org/10.23880/whsj-16000108>
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. *Gynaecol Perinatol* 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 Scient Med Sci* 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 J Cancer Res Rev* 2018; 2: 5.  
<https://doi.org/10.28933/ajocrr-2017-12-2801>
43. 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.  
<https://doi.org/10.31532/CancerStud.2.1.002>
45. Zaichick V, Zaichick S. Twenty chemical element contents in normal and cancerous thyroid. *Int J Hematol Blo Dis* 2018;3(2):1-13. <https://doi.org/10.1007/s10552-008-9260-4>
46. 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.  
<https://doi.org/10.20517/2394-4722.2018.52>
47. Zaichick V, Zaichick S. Fifty trace element contents in normal and cancerous thyroid. *Acta Scientific Cancer Biol* 2018;2(8):21-38.
48. Zaichick V, Zaichick S. Instrumental effect on the contamination of biomedical samples in the course of sampling. *The J Analyt Chem* 1996;51(12):1200-1205.
49. Zaichick V, Tsislyak YuV. A simple device for biosample lyophilic drying. *Lab Delo* 1978; 2: 109-110.
50. 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.  
<https://doi.org/10.1007/s12011-009-8456-0>
51. Zaichick V, Nosenko S, Moskvina I. The effect of age on 12 chemical element contents in intact prostate of adult men investigated by inductively coupled plasma atomic emission spectrometry. *Biol Trace Elem Res* 2012; 147:49-58.  
<https://doi.org/10.1007/s12011-011-9294-4>
52. Zaichick V, Zaichick S. NAA-SLR and ICP-AES Application in the assessment of mass fraction of 19 chemical elements in pediatric and young adult prostate glands. *Biol Trace Elem Res* 2013; 156:357-366.  
<https://doi.org/10.1007/s12011-013-9826-1>
53. Zaichick V, Zaichick S. Determination of trace elements in adults and geriatric prostate combining neutron activation with inductively coupled plasma atomic emission spectrometry. *Open J Biochem* 2014; 1(2):16-33.  
<https://doi.org/10.4236/ajac.2014.511079>
54. Zaichick S, Zaichick V. 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* 2011;288:197-202.  
<https://doi.org/10.1007/s10967-010-0927-4>
55. Zaichick V, Zaichick S. 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. *J Appl Radiat Isot* 2013; 82:145-151.  
<https://doi.org/10.1016/j.apradiso.2013.07.035>

56. Zaichick V. Applications of synthetic reference materials in the Medical Radiological Research Centre. *Fresenius J Anal Chem* 1995; 352:219-223. <https://doi.org/10.1007/BF00322330>
57. Korelo AM, Zaichick V. Software to optimize the multielement INAA of medical and environmental samples. In: *Activation Analysis in Environment Protection*. Dubna, Russia: Joint Institute for Nuclear Research; 1993: 326-332.
58. Korteve AI, Dontsov GI, Lyascheva AP. Bioelements and a human pathology. Sverdlovsk, Russia: Middle-Ural publishing-house; 1972.
59. Kamenev VF. About trace element contents in thyroid of adults. In: *Trace Elements in Agriculture and Medicine*. Ulan-Ude, Russia: Buryatia publishing-house; 1963:12-16.
60. Tipton IH, Cook MJ. Trace elements in human tissue. Part II. Adult subjects from the United States. *Health Phys* 1963; 9(2):103-145. <https://doi.org/10.1097/00004032-196302000-00002>
61. Reitblat MA, Kropachyev AM. Some trace elements in thyroid of the Perm Pricam'ya residents. *Proceedings of Perm Medical Institute* 1967; 78:157-164.
62. Forssen A. Inorganic elements in the human body. I. Occurrence of Ba, Br, Ca, Cd, Cs, Cu, K, Mn, Ni, Sn, Sr, Y and Zn in the human body. *Ann Med Exp Biol Fenn* 1972; 50(3):99-162. PMID: 5081903
63. Zhu H, Wang N, Zhang Y, *et al.* Element contents in organs and tissues of Chinese adult men. *Health Phys* 2010; 98(1):61-73. <https://doi.org/10.1097/HP.0b013e3181bad921>
64. 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.
65. 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 k $\alpha$ -method. *J Radioanal Nucl Chem* 1997; 222(1-2): 11-14. <https://doi.org/10.1007/BF02034238>
66. Reddy SB, Charles MJ, Kumar MR, *et al.* Trace elemental analysis of adenoma and carcinoma thyroid by PIXE method. *Nucl Instrum Methods Phys Res B: Beam Interactions with Materials and Atoms* 2002; 196(3-4): 333-339. [https://www.researchgate.net/publication/223853105\\_Trace\\_elemental\\_analysis\\_of\\_adenoma\\_and\\_carcinoma\\_thyroid\\_by\\_PIXE\\_method](https://www.researchgate.net/publication/223853105_Trace_elemental_analysis_of_adenoma_and_carcinoma_thyroid_by_PIXE_method)
67. Woodard HQ, White DR. The composition of body tissues. *Brit J Radiol* 1986; 708: 1209-1218. [https://doi.org/10.1016/S0168-583X\(02\)01292-2](https://doi.org/10.1016/S0168-583X(02)01292-2)
68. 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. PMID: 5807109
69. Neimark II, Timoshnikov VM. Development of thyroid cancer in persons living in the endemic goiter area. *Probl Endocrinol (Mosk)* 1978;24(3):28-32. PMID: 674124
70. Zabala J, Carrion N, Murillo M, *et al.* Determination of normal human intrathyroidal iodine in Caracas population. *J Trace Elem Med Biol* 2009; 23(1): 9-14. <https://doi.org/10.1016/j.jtemb.2008.11.002>
71. Zakutinsky DK, Parfyenov YuD, Selivanova LN. *Handbook of the toxicology of radioactive isotopes*. Moscow: State Publishing House of Medical Literature; 1962.
72. Remiz AM. Endemic goiter and trace elements in Kabardino-Balkaria ASSR. In: *The 5<sup>th</sup> meeting of surgeons of Northern Caucasia*. Rostov-on-Don: 1962:276-278.
73. Li AA. Level of some macro- and trace element contents in blood and thyroid of patients with endemic goiter in Kalinin region. PhD thesis. Kalinin, Russia: Kalinin medical institute; 1973.
74. 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.
75. Soman SD, Joseph KT, Raut SJ, Mulay CD, Parameshwaran M, Panday VK. Studies of major and trace element content in human tissues. *Health Phys* 1970; 19(5):641-656.
76. Novikov GV, Vlasova ZA. Some organism functions in connection with the iodine content in diet and feed of experimental animals. In: *Role of Trace Elements in Agriculture and Medicine*. Leningrad: Nauka; 1970;2:6-7.
77. Bredikhin LM, Soroka VP. Trace element metabolism in patients with goiter during therapy. *Vrach Delo*. 1969; 51(6): 81-84. PMID: 5821671
78. Byrne AR. Vanadium in foods and in human body fluids and tissues. *Sci Total Environ* 1978; 10: 17-30. [https://doi.org/10.1016/0048-9697\(78\)90046-3](https://doi.org/10.1016/0048-9697(78)90046-3)
79. Ianchur NM, Elenevskaia NS, But-Gusaim AM, Nikhamkin LI. The content of manganese, aluminum, copper and zinc in the blood and the thyroid gland of patients with goiter. *Klin Khir* 1967; 4:27-30. PMID: 4888317
80. Antonova MV, Elinova VG, Voitekhovskaya YaV. Some trace element contents in thyroid and water in endemic goiter region. *Zdravookhranenie BSSR* 1966; 9:42-44.
81. Maeda K, Yokode Y, Sasa Y, Kusuyama H, Uda M. Multielemental analysis of human thyroid glands using particle induced X-ray emission (PIXE). *Nucl Instrum Methods Phys Res B* 1987; 22(1-3):188-190. [https://doi.org/10.1016/0168-583X\(87\)90323-5](https://doi.org/10.1016/0168-583X(87)90323-5)
82. Turetskaia ES. Studies on goitrous thyroid glands for iodine and bromine content. *Probl Endocrinol Gormonoter* 1961;7(2):75-80. PMID: 13778682
83. Aingorn NM, Chartorizhskaya NA. Comparative characteristics of trace element contents under thyroid disorders. In: *Trace Elements in Agriculture and Medicine*. Ulan-Ude, Russia: Buryatiapublishing-house; 1966:113-114.
84. Kaya G, Avci H, Akdeniz I, Yaman M. Determination of trace and minor metals in benign and malignant human thyroid tissues. *Asian J Chem* 2009; 21(7):5718-5726.
85. Dimitriadou A, Suvanik R, Fraser TR, Pearson J.D. Endemic goiter in Thailand. A study contrasting these iodine-deficient goiters with sporadic non-toxic goiters seen in London. *J Endocrinol* 1966; 34(1):23-39. <https://doi.org/10.1677/joe.0.0340023>
86. Braasch JW, Abbert A, Keating FR, Black BM. A note of the iodinated constituents of normal thyroids and of exophthalmic goiters. *J Clin Endocrinol Metab* 1955; 15(4):732-738. <https://doi.org/10.1210/jcem-15-6-732>
87. Bolkvadze AI. Contents of electrolytes (K, Na, Ca, I and F) in thyroid and blood under different forms of thyroid pathology. PhD thesis. Tbilisi: Tbilisi medical institute; 1970.
88. Borodin AE, Sokolova II, Gogolev VG, Makarova MYa. About goitrous thyroid chemical composition. In: *Goiter in Amur region*. Blagoveshchensk: Khabarovsk publishing-house; 1967:21-29.
89. Stojsavljević A, Rovčanin B, Krstić D, *et al.* Evaluation of trace metals in thyroid tissues: Comparative analysis with benign and malignant thyroid diseases. *Ecotoxicol Environ Saf* 2019;183:109479. <https://doi.org/10.1016/j.ecoenv.2019.109479>
90. Petrov IC, Alyab'ev GA, Dmitrichenko MM. Contents of iodine, manganese, and cobalt in thyroid and blood in the local residents and migrants of Irkutsk region. In: *Trace elements in agriculture and medicine*. Ulan-Ude, Russia: Buryatia publishing-house; 1968:648-651.
91. Zagrodzki P, Nicol F, Arthur JR, *et al.* Selenoenzymes, laboratory parameters, and trace elements in different types of thyroid tumor. *Biol Trace Elem Res* 2010; 134(1): 25-40. <https://doi.org/10.1007/s12011-009-8454-2>
92. 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. <https://doi.org/10.1385/BTER:90:1-3:57>
93. Schroeder HA, Tipton IH, Nason AP. Trace metals in man: strontium and barium. *J Chron Dis* 1972; 25(9):491-517. [https://doi.org/10.1016/0021-9681\(72\)90150-6](https://doi.org/10.1016/0021-9681(72)90150-6)
94. Zaichick V. Sampling, sample storage and preparation of biomaterials for INAA in clinical medicine, occupational and environmental health. In: *Harmonization of health-related*

- environmental measurements using nuclear and isotopic techniques. Vienna: IAEA; 1997:123-133.
95. 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. <https://doi.org/10.1007/BF02039345>
96. Zaichick V. Losses of chemical elements in biological samples under the dry aching process. *Trace Elements in Medicine* 2004; 5(3):17-22.
97. Krewski D, Yokel RA, Nieboer E, *et al.* Human health risk assessment for aluminium, aluminium oxide, and aluminium hydroxide. *J Toxicol Environ Health Part B* 2007; 10:1-269. <https://doi.org/10.1080/10937400701597766>
98. Naghii MR, Mofid M, Asgari AR, Hedayati M, Daneshpour MS. Comparative effects of daily and weekly boron supplementation on plasma steroid hormones and proinflammatory cytokines. *J Trace Elem Med Biol* 2011; 25:54-58. <https://doi.org/10.1016/j.jtemb.2010.10.001>
99. Pizzorno L. Nothing boring about boron. *Integr Med (Encinitas)* 2015; 14:35-48. PMID: 26770156
100. Pavelka S. Radiometric determination of thyrotoxic effects of some xenobiotics. *Rad Applic* 2016; 1(2):155-158. <https://doi.org/10.21175/RadJ.2016.02.029>
101. Maschkovsky MD. The sedatives. In: *The Medicaments*. 15<sup>th</sup> ed. Moscow: Novaya Volna; 2005:72-86.
102. Zaichick V. X-ray fluorescence analysis of bromine for the estimation of extracellular water. *J Appl Radiat Isot* 1998; 49(12): 1165-1169. [https://doi.org/10.1016/s0969-8043\(97\)10118-x](https://doi.org/10.1016/s0969-8043(97)10118-x)
103. Lyshchik A, Moses R, Barnes SI, *et al.* Quantitative analysis of tumor vascularity in benign and malignant solid thyroid nodules. *J Ultrasound Med* 2007; 26(6): 837-846. <https://doi.org/10.7863/jum.2007.26.6.837>
104. Li Y, Trush MA. DNA damage resulting from the oxidation of hydroquinone by copper: role for a Cu(II)/Cu(I) redox cycle and reactive oxygen generation. *Carcinogenesis* 1993; 14(7): 1303-1311. <https://doi.org/10.1093/carcin/14.7.1303>
105. Torti SV, Manz DH, Paul BT, Blanchette- Farra N, Torti FM. Iron and Cancer. *Annu Rev Nutr* 2018; 38: 97-125. <https://doi.org/10.1146/annurev-nutr-082117-051732>
106. Selby JV, Friedman GD. Epidemiologic evidence of an association between body iron stores and risk of cancer. *Int J Cancer* 1988; 41: 677-682. <https://doi.org/10.1002/ijc.2910410507>
107. Meneghini R. Iron homeostasis, oxidative stress, and DNA damage. *Free Radic Biol Med* 1997; 23: 783-792. [https://doi.org/10.1016/s0891-5849\(97\)00016-6](https://doi.org/10.1016/s0891-5849(97)00016-6)
108. Anke M, Arnold W, Schäfer U, Müller R. Recent progress in exploring the essentiality of the ultratrace element lithium to the nutrition of animals and man. *Biomed Res Trace Elem* 2005; 16(3): 169-176. <https://doi.org/10.11299/brte.16.169>
109. Czarnywojtek A, Zgorzalewicz-Stachowiak M, Czarnocka B, *et al.* Effect of lithium carbonate on the function of the thyroid gland: mechanism of action and clinical implications. *J Physiol Pharmacol* 2020; 71(2): 191-199. <https://doi.org/10.26402/jpp.2020.2.03>
110. Chandra A.K. Effects of magnesium on cytomorphology and enzyme activities in thyroid of rats. *Indian J Exp Biol* 2014; 52: 787-792. PMID: 25141541
111. Soldin OP, Aschner M. Effects of manganese on thyroid hormone homeostasis. *Neurotoxicol* 2007; 28: 951-956. <https://doi.org/10.1016/j.neuro.2007.05.003>
112. Trump BF, Berezsky IK, Phelps PC. Sodium and calcium regulation and the role of the cytoskeleton in the pathogenesis of disease: A review and hypothesis. *Scan Electron Microsc* 1981; (Pt 2): 434-454. PMID: 7034180
113. Ravera S, Reyna-Neyra A, Ferrandino G, Amzel M, Carrasco N, The sodium/iodide symporter (NIS): Molecular physiology and preclinical and clinical applications. *Annu Rev Physiol* 2017; 79: 261-289. <https://doi.org/10.1146/annurev-physiol-022516-034125>
114. Ashmaik AS, Gabra HM, Elzein AOM, Shrif NEMA, Hassan EE. Assessment of serum levels of calcium and phosphorous in Sudanese patients with hypothyroidism. *Asian J Biomed Pharm* 2013; 3(25): 21-26.
115. Stelmach H, Moško P, Jaroszewicz L, Piotrowski Z, Puchalski Z. Phospholipids of human thyroid gland. *Acta Physiol Hung* 1993; 81(3): 263-267. PMID: 8197881
116. Townsend DM, Tew KD, Tapiero H. Sulfur containing amino acids and human disease. *Biomed Pharmacother* 2004; 58: 47-55. <https://doi.org/10.1016/j.biopha.2003.11.005>
117. Atmaca G. Antioxidant effects of sulfur-containing amino acids. *Yonsei Med J* 2004; 45: 776-788. <https://doi.org/10.3349/ymj.2004.45.5.776>
118. Pérez-Granados AM, Vaquero MP. Silicon, aluminium, arsenic and lithium: essentiality and human health implications. *J Nutr Health Aging* 2002; 6(2): 154-162. PMID: 12166372
119. Nielsen FH. Nutritional requirements for boron, silicon, vanadium, nickel, and arsenic: current knowledge and speculation. *FASEB J* 1991; 5(12): 2661-2667. PMID: 1916090
120. Semenov VD, Suslikov VL. Role of nutrition in the development of functional shifts in the thyroid. *Vopr Pitan* 1983; 3: 65-68. PMID: 6225246
121. Najda J, Gmiński J, Drózd M, Zych F. The influence of inorganic silicon (Si) on pituitary-thyroid axis. *Biol Trace Elem Res* 1993; 37(2-3): 101-106. <https://doi.org/10.1007/BF02783785>
122. Gruzewska K, Michno A, Pawelczyk T, Bielarczyk H. Essentiality and toxicity of vanadium supplements in health and pathology. *J Physiol Pharmacol* 2014; 65(5): 603-611. PMID: 25371519
123. Fallahi P, Foddiss R, Elia G, *et al.* Vanadium pentoxide induces the secretion of CXCL9 and CXCL10 chemokines in thyroid cells. *Oncol Rep* 2018; 39(5): 2422-2426. <https://doi.org/10.3892/or.2018.6307>
124. Bozym RA, Chimienti F, Giblin LJ, *et al.* Free zinc ions outside a narrow concentration range are toxic to a variety of cells *in vitro*. *Exp Biol Med* (Maywood) 2010; 235(6): 741-750. <https://doi.org/10.1258/ebm.2010.009258>
125. Matusik RJ, Kreis C, McNicol P, *et al.* Regulation of prostatic genes: role of androgens and zinc in gene expression. *Biochem Cell Biol* 1986; 64: 601-607. <https://doi.org/10.1139/o86-083>
126. Blok LJ, Grossmann ME, Perry JE, Tindall DJ. Characterization of an early growth response gene, which encodes a zinc finger transcription factor, potentially involved in cell cycle regulation. *Mol Endocrinol* 1995; 9(11): 1610-1620. <https://doi.org/10.1210/mend.9.11.8584037>
127. Zezerov EG. Hormonal and molecular biological factors in pathogenesis of prostate cancer. *Vopr Onkol* 2001; 47(2):174-181. PMID: 11383453
128. Truong-Tran AQ, Ho LH, Chai F, Zalewski PD. Cellular zinc fluxes and the regulation of apoptosis/gene-directed cell death. *J Nutr* 2000; 130(5S Suppl): 1459S-1466S. <https://doi.org/10.1093/jn/130.5.1459S>
129. Kontargiris E, Vadalouka A, Ragos V, Kalfakakou V. Zinc inhibits apoptosis and maintains NEP downregulation, induced by Ropivacaine, in HaCaT cells. *Biol Trace Elem Res* 2012; 150: 460-466. <https://doi.org/10.1007/s12011-012-9492-8>
130. Liang D, Yang M, Guo B, *et al.* Zinc inhibits H<sub>2</sub>O<sub>2</sub>-induced MC3T3-E1 cells apoptosis via MAPK and PI3K/AKT pathways. *Biol Trace Elem Res* 2012; 148: 420-429. <https://doi.org/10.1007/s12011-012-9387-8>
131. Zhang X, Liang D, Guo B, Yang L, Wang L, Ma J. Zinc inhibits high glucose-induced apoptosis in peritoneal mesothelial cells. *Biol Trace Elem Res* 2012; 150: 424-432. <https://doi.org/10.1007/s12011-012-9473-y>