ISSN: 2319-3387 Volume 8, Issue 3 www.stmjournals.com

Uranium Content Anomalies in Groundwater of Moga District of Malwa Belt of Punjab (India) for the Assessment of Excess Cancer Risk

Hardev Singh Virk*

Professor of Eminence, Punjabi University, Patiala, Punjab, India

Abstract

Inductively coupled plasma mass spectroscopy (ICPMS) has been used to measure the Uranium (U) content of the ground water samples of Moga district of Malwa belt of Punjab, India. Out of the total number of habitations covered under this survey, 203 were having U content more than 30 ppb (WHO safe limit). Out of 203 habitations, we sorted out 56 with U concentration of more than 100 ppb in groundwater. The U content of ground water of Moga district was used to assess the radiological and chemical risk due to U ingestion to the population of the district. The U content of water samples of the villages selected for our investigation varied from 100.20 to 346.70 ppb (µgl $^{-1}$) with an average value of 166.11 ppb. The excess cancer risk varied from 2.84×10^{-4} to 9.82×10^{-4} , with an average value of 5.49×10^{-4} . The Lifetime Average Daily Dose (LADD) varied from 5.80 to 12.06 (µg kg $^{-1}$ day $^{-1}$) and hazard quotient varied from 1.28 to 4.43, respectively.

Keywords: AERB limit, cancer risk, chemical risk, radiological risk, Uranium (U) content

*Author for Correspondence E-mail: hardevsingh.virk@gmail.com

INTRODUCTION

The World Health Organization (WHO) [1] recommended a reference level of 30 µg 1-1 (ppb) as the permissible limit of uranium (U) in drinking water. The accumulation of U inside the human body results in its chemical and radioactive effects for the two important target organs being the kidneys and lungs [2–4]. The U route to the human body is mainly through drinking water, which contributes about 85% of ingested U and the rest 15% is contributed by the food intake [5]. The transient chemical damage to the kidneys is due to an exposure of about 0.1 mg/kg of body weight of soluble natural U [6]. The natural U is a radioactive heavy metal and its decay products are created as many other radioactive metals or gases in a chain reaction, which can further become a health hazard to the public [7]. U itself is a weak radioactive metal but it may be hazardous to human health if its contamination is high in the drinking water. The assessment of health hazards risk is important, if the concentration of U in water and its extent of getting ingested into the human body is higher than the safe limit provided by the WHO [1].

Punjab is facing a crisis situation due to high levels of U and heavy metals in underground water table of Punjab. More than two dozen reports have been published in The Tribune (www.tribuneindia.com) and other media concerning high toxicity of U in the waters of Punjab setting the alarm bells ringing in the Indian parliament. The author has reported his findings on U contamination and its health hazards in research journals during the past four decades [8-17]. U estimation of the groundwater of Malwa belt of Punjab State and the neighbouring areas in Haryana, Himachal Pradesh, and Jammu has been reported by other workers [18-24]. The present report is based on the data collected by the Punjab Water Supply and Sanitation Department (PWSSD), Mohali, Punjab, India. It is also available on the Ministry of Water Resources, Government of India, and website: www.indiawater.gov.in/IMIS reports. objective of the present investigations was health risk assessment due to natural U in drinking water in Moga district of Punjab, India (Figure. 1).

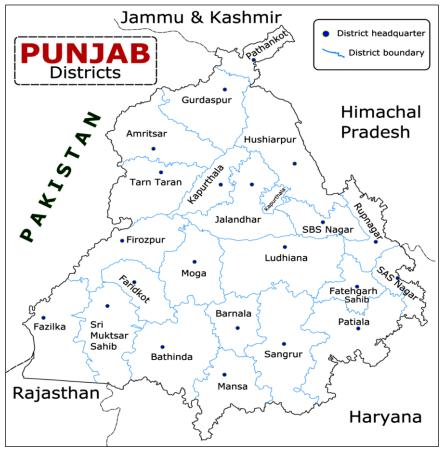


Fig. 1: District Map of Punjab Showing District of Moga in the Malwa Belt.

STUDY AREA AND GROUNDWATER QUALITY Location

The Central Ground Water Board (CGWB) Report throws some light on the nature of soil and geomorphology of the Moga district [25]. It has geographical area of 2071 km². Its boundaries are defined by North Latitude 30°28′30″ and 31°06′15″ and East Longitude 74°54′40″ and 75°24′57″. Moga district is surrounded by Ferozpur and Faridkot in the West, Bathinda and Barnala in the South, Ludhiana in the East and Jalandhar in the North.

Geomorphology and Soil Types [25]

The district forms a part of Indo-Gangetic plain and Sutlej sub-basin of Indus basin. The area as a whole is almost flat with a gentle slope towards the Western and North-westerly direction. The physiography of the district is broadly classified from south to north into four distinct features, i.e., upland plain, sand dune tract, younger flood plain and active flood plain of Sutlej. The Sutlej is an important perennial

river, which forms major drainage of the area and runs parallel to the Northern border of the district. There are two types of soils, viz. Sierozem and Desert soils in Moga district. The sierozem soils are found in major parts of the district and desert soils are found in a relatively smaller area towards the western part of the district.

Ground Water Quality [25]

Aquifer material comprises chiefly of fine-to-medium grained sand. In general, shallow aquifers in the area are unconfined down to depth of 80 m in the district and being semi-confined in the depth range of 100–111 m. These two aquifers are separated by a clay layer almost uniformly present in the district. The aquifer down to depth of 50 m is being tapped by shallow tubewells for purpose of irrigation and drinking. However, few deeper tubewells down to depth of 125 m are being tapped by the Government agencies for drinking purpose and by some farmers for irrigation purpose.



The shallow aquifers show groundwater quality to be marginally saline. Groundwater has higher fluoride concentration than the permissible limit in most of the areas of this district. Iron concentrations have been found in shallow groundwater at some places to be higher than permissible limits in South Eastern, North Western and in areas adjoining river Satluj. By and large, the groundwater is suitable for domestic/irrigation purposes in major part of the district.

MATERIALS AND METHODS

For collection of samples, 20 ml bottles of superior quality plastic were used. The bottles were washed first with soap solution and then with distilled water. These were rinsed with deionised water and dried. Groundwater from the source was allowed to flow freely before collection in plastic bottles. About 10-20 ml of water was collected from the running water source. Samples must be filtered through a 0.45 um capsule filter at the field site to separate the dissolved metal components. Nitric acid (0.5M HNO₃) solubilization is required before the determination of total recoverable U. The preservation and digestion of U in acid is used in order to aid breakdown of complexes and to minimize interferences by polyatoms.

The U analysis of collected water samples has been done using Model 7700 Agilent Series ICP-MS following standard procedure in the Punjab State laboratory set up in Mohali. The method is suitable to measure ions produced by radiofrequency inductively coupled plasma. Analyte species originating in a liquid were nebulized and the resulting aerosol was transported by Argon gas into the plasma torch. The ions produced by high temperatures were entrained in the plasma gas and introduced, by means of an interface, into a mass spectrometer. The ions produced in the plasma were sorted according to their mass-to-charge ratios and quantified with a channel electron multiplier. Interferences must be assessed and valid corrections applied. Interference correction must include compensation for background ions contributed by the plasma gas, reagents, and constituents of the sample matrix.

A mass spectrometer with inductively coupled plasma (ICP) suitable for multi-element and

isotope analysis is required. The spectrometer should be capable of scanning a mass range from 5 m/z (AMU) to 240 m/z (AMU) with a resolution of at least 1 mr/z peak width at 5% of peak height (mr = relative mass of an atom species; z = charge number). The instrument may be fitted with a conventional or extended dynamic range detection system. Most quadrupole ICP-MS, high-resolution ICP-MS and collision cell ICP-MS instrumentation is fit for this purpose. Data analysis was done automatically by the inbuilt system of ICP-MS. In addition to U, data for 40 more trace elements can be retrieved using ICP-MS.

THEORETICAL FORMULATION

Ingestion of U through drinking water results in both—radiological risk (carcinogenic) and chemical risk (noncarcinogenic). The methodology used for the assessment of radiological and chemical risks due to U concentrations in the water samples are described below:

Radiological Risk Assessment Calculation of Excess Cancer Risk

Excess cancer risk from the ingestion of natural U from drinking water has been calculated according to the standard method given by the United States Environmental Protection Agency (USEPA) [26]:

 $ECR = Ac \times R$

Where, 'ECR' is Excess Cancer Risk; 'Ac' is Activity concentration of Uranium (Bql⁻¹) and 'R' is Risk Factor.

The risk factor R (per Bql⁻¹), linked with ingestion of U from drinking water may be estimated by the product of the risk coefficient (r) of U (1.19×10⁻⁹) for mortality and per capita activity intake I. 'I' for U is calculated as product of life expectancy, assumed to be 63.7 years, i.e. 23250 days and daily consumption of water as 4.05 lday⁻¹ [27].

 $I = 4.05 \text{ lday}^{-1} \times 23250 \text{ days}$ Risk Factor (R) = r× I

Chemical Risk Assessment

The chemical toxicity risk for U is defined in terms of Lifetime Average Daily Dose (LADD) of U through drinking water intake. LADD is defined as the quantity of the substance ingested per kg of body weight per day and is given by the following equation [28, 29]:

LADD= C×IR×ED×EFAT×BW ×365

Where, 'C' is the concentration of U (μ gl⁻¹); IR is the water consumption rate (4.05 lday⁻¹); ED is the lifetime exposure duration (63.7 years); EF is the exposure frequency (365 daysy⁻¹); BW is average body weight of the receptor (70 kg); and AT is the Averaging time, i.e., life expectancy (63.7 years).

Calculation of Hazard Quotient

Hazard quotient (HQ) is the measure of the extent of harm produced due to the ingestion of U from the drinking water.

HQ=LADD/RfD

Where, LADD is lifetime average daily dose, and RfD is the reference dose = $4.53 \mu g kg^{-1}day^{-1}$.

RESULTS AND DISCUSSION

Groundwater samples were collected from villages falling under the Moga district of Punjab and analysed for U content using calibrated ICP-MS. U content varied from 33.10 ppb (tubewell in Garden Colony) to 346.70 ppb (tubewell at Chotian Khurd) for all habitations covered under this survey. The variation of U in groundwater for 56 habitations selected (Table 1) for our investigations was 100.20 ppb (tubewell at Khukhurana) to 346.70 ppb (tubewell at Chotian Khurd), with an average value of 166.11 ppb. The safe limit of U in groundwater is fixed to be 60 ppb by Atomic Energy Regulatory Board (AERB) [30] in India, while other agencies fix it in much lower limits of 30 ppb (EPA, USA) [26]; 15 ppb (WHO) [11]; 9 ppb (UNSCEAR) [31] and 1.9 ppb (ICRP) [32]. If the observed data of U content of water are compared with the guidelines of AERB, all 56 samples record higher U content than 60 ppb (Table 1); hence they fail to qualify the safe limit certification of AERB, Government of India.

Radiological Risk

The radiological risk has been calculated due to ingestion of natural U in the drinking water of 56 habitations sorted out for this survey, assuming the consumption rate of 4.05 l/day and lifetime expectancy of 63.7 years for both males and females. The excess cancer risk for the population inhabiting these 56 habitations

has been observed to be in the range of 2.84×10^{-4} – 9.82×10^{-4} , with an average value of 5.49×10⁻⁴. The value of excess cancer risk in the surveyed habitations is much higher than the maximum acceptable level of 1.67×10⁻⁴ according to the AERB guidelines. If we assume lifetime water consumption rate of 4.05 l/day with the present average U content of water, the mean value of excess cancer risk in the surveyed habitations comes out to be 5.49×10⁻⁴, which works out to be nearly 5.5 per 10,000 people. According to the Cancer Registry of Government of India, national average of cancer risk is 80 cancers per million population; while for Punjab it is slightly higher with 90 cancers per million. It is much higher for Malwa belt of Punjab [13–16]. Our investigation revealed that for Moga district in the Malwa belt of Punjab, it has assumed alarming proportions at 550 cancers per million.

Chemical Toxicity Risk

Uranium is a radioactive heavy metal, so it has health impacts due to its both radioactive and chemical nature. If we take into account chemical toxicity of U, the kidneys are the most important target organ. The chemical toxicity of U dominates over its radiological toxicity on the kidney in general at lower exposure levels [33]. The chemical toxicity has been estimated from the value of LADD and HQ. HQ has been estimated by comparing the value of the calculated LADD with the reference dose level of 4.53 µg kg⁻¹day⁻¹. The reference level has been calculated for the maximum contamination level of U in water of 60 ppb (μ gl⁻¹). The variation in the values of LADD and HOs have been observed from 5.80 μg/kg/day to 12.06 μg/kg/day and from 1.28 to 4.43, respectively (Table 1).

HQ higher than one is an indication of chemical toxicity risk factor due to ingested U via the drinking water. For overall risk assessment, both radiological and chemical toxicity are to be taken into consideration. It has become possible to estimate age-dependent ingestion and inhalation doses due to intake of U to the exposed population in the area [24].

Volume 8, Issue 3 ISSN: 2319-3387



Table 1: Uranium Content in Groundwater of Moga District and Corresponding Risk Factors (WHO limit 30 ppb; AERB limit 60 ppb).

Excess Cancer LADD (µg Sr. Location Source of Depth Uranium Uranium $kg^{-1} day^{-1}$ Conc. (ppb) Conc. (Bq l-1) risk * 10⁻⁴ No. Groundwater (m) Quotient 100.20 2.84 1 Khukhurana Tubewell 131.4 2.53 5.80 1.28 2.55 1.29 Bare Wala Tubewell 108.0 101.00 2.86 5.84 3 Budh Singh Wala Tubewell 108.0 101.30 2.56 2.87 5.86 1.29 2.61 2.92 5.97 4 Sadda Singh Wala Tubewell 131.06 103.10 1.32 2.93 5 Badhni Kalan <u>Tub</u>ewell 118.87 103.30 2.61 5.98 1.32 5.98 2.93 S.C.Basti Null Null 103.35 2.61 1.32 6 Sukha Nand Null Null 103.35 2.93 5.98 1.32 7 2.61 Tubewell 2.95 8 RamuwalaKalan Null 104.00 2.63 6.02 1.33 Tubewell 119.0 104.10 2.95 6.02 1.33 9 Minian 2.63 10 Bode Null 107.90 2.73 3.06 1.38 Null 6.24 2.78 1.41 11 Nangal Null Null 110.13 3.12 6.37 2.83 3.17 Tubewell 128.2 112.00 1.43 Vadda Ghar I 6.48 112.89 2.85 3.20 6.53 1.44 13 Dhurkot Kalan Null Null 2.88 14 Vadda Ghar II Tubewell 128.02 114.00 3.23 6.60 1.46 15 Langiana Khurd Tubewell 137.5 123.00 3.11 3.48 7.12 1.57 1.57 16 Abadi of Langiana Purana Tubewell 137.5 123.00 3.11 3.48 7.12 123.10 3.49 1.57 17 Bare Wala Tubewell 108.0 3.11 7.12 18 Baghele Wala Tubewell Null 123.60 3.12 3.50 7.15 1.58 19 Chotian Khurd Tubewell 153.0 123.60 3.12 3.50 7.15 1.58 3.15 1.59 20 Nihal Garh 118.87 124.50 3.53 7.20 Tubewell 124.50 1.59 21 Longiwind 118.87 3.53 7.20 Tubewell 3.15 22 Tehsil Complex Tubewell 180.0 124.70 $3.1\overline{5}$ 3.53 7.21 1.59 23 Kothe Kartar Singh Handpump Null 129.90 3.28 3.68 7.52 1.66 24 Mahlan Kalan Tubewell 107.0 130.50 3.30 3.70 7.55 1.67 7.56 25 | Chotian Kalan **Fubewell** 137.16 130.60 3.30 3.70 1.67 7.56 3.30 3.70 26 Saho Ke Tubewell 86.0 130.60 1.67 7.75 27 Kore Wala Khurd 3.39 3.80 1.71 134.00 Tubewell Null 134.30 28 Nathe Wala I 3.40 3.80 132.65 7.77 1.72 Tubewell 29 Nathe Wala II 148.5 134.30 3.40 3.80 7.77 1.72 Tubewell 30 Budh Singh Wala Tubewell 65.0 138.60 3.50 3.93 8.02 1.77 3.95 1.78 31 Ransih Kalan Tubewell 114.3 139.60 3.53 8.08 32 Kalie Wala I 140.60 3.55 3.98 1.80 Tubewell 106.0 8.13 33 Kalie Wala II Tubewell Null 144.50 3.65 4.09 1.85 8.36 34 Bambiha Bhai Null Null 148.30 3.75 4.20 8.58 1.89 3.75 35 Droli Bhai Tubewell Null 148.30 4.20 8.58 1.89 36 SinghaWala 148.50 1.90 Handpump 60.96 3.75 4.21 8.59 85.0 37 Jita Singh Wala 4.59 9.37 2.07 Tubewell 162.00 4.10 4.70 2.12 38 Rattian Tubewell Null 166.00 4.20 9.60 2.23 39 Abadi on Talwandi Road Tubewell 118.87 174.90 4.42 4.95 10.12 2.23 40 Saido Ke **Fubewell** 118.87 174.90 4.42 4.95 10.12 2.24 4.97 175.50 4.44 10.15 41 Sangatpura Tubewell 89.0 2.25 Ganji Gulab Singh 107.0 175.90 4.45 4.98 10.18 42 Tubewell 2.28 178.80 4.52 5.06 10.34 43 Kahan Singh Wala Tubewell Null 2.37 180.0 185.70 4.69 5.26 10.74 44 Ransih Khurd Tubewell 2.39 45 Ganji Gulab Singh Tubewell Null 187.00 4.73 5.30 10.82 95.0 187.90 5.32 2.40 46 Harijan Basti Tubewell 4.75 10.87 47 Khotte Tubewell 95.0 187.90 4.75 5.32 10.87 2.40 48 Takhtupura Tubewell 114.3 188.20 4.76 5.33 10.89 2.40 5.72 2.58 49 Kore Wala Kalan Tubewell Null 202.00 5.11 11.69 2.96 50 Ganji Gulab Singh RO Raw Water 232.00 5.86 6.57 13.42 Null 51 Ganji Gulab Singh 107.0 246.10 6.22 6.97 14.24 3.14 Tubewell 3.18 52 Daya Kalan Tubewell 180.0 248.70 6.29 7.04 14.39 53 Rau Ke Kalan Tubewell 107.0 251.30 6.35 7.12 14.54 3.21 251.30 14.54 3.21 54 Harijan Basti Tubewell 107.0 6.35 7.12 7.79 Thraj Tubewell 116.5 275.00 6.95 15.91 3.51 9.82 56 Chotian Khurd Tubewell Null 346.70 8.76 20.06 4.43

^{*}Cancer risk is the likelihood, or chance, of getting cancer.

We say "excess cancer risk" because we have a "background risk" of about one in four chances of getting cancer. In other words, in a million people, it is expected that 250,000 individuals would get cancer from a variety of causes. If we say that there is a "one in a million" excess cancer risk from a given exposure to a contaminant, we mean that if one million people are exposed to a carcinogen at a certain concentration over their lifetime, then one cancer above the background chance, or the 250,000th cancer, may appear in those million persons from that particular exposure [34].

CONCLUSIONS

- (i). The concentration of U in groundwater samples collected from the tubewells, handpumps and RO raw water of 56 villages of Moga district was found to be much higher than the safe limit of 60 ppb recommended by the AERB, India.
- (ii). Moga district has second highest number of habitations in Punjab, next only to Fazilka, where U contamination of groundwater has reached alarming levels.
- (iii). The cancer risk due to presence of U in groundwater was found to be among the highest for the districts of Punjab with an estimated number of 550 cancers per million population.
- (iv). It will be of interest to study the epidemiological effects of U in groundwater on the inhabitants of Moga district of Punjab, India.

ACKNOWLEDGEMENT

Author is obliged to thank the authorities of PWSSD for supply of uranium in groundwater data of Moga district.

REFERENCES

- 1. World Health Organization. *Guidelines for drinking-water quality, 4th Ed.* Geneva, Switzerland: WHO; 2011.
- 2. World Health Organization. *Life in the 21st century: A vision for all.* Geneva, Switzerland: WHO; 1998.
- 3. Agency for Toxic Substances and Diseases Registry. *Toxicological profile for Radium*. Atlanta, Georgia: US Department of Health and Human Services, ATSDR; 1990.
- 4. Bleise A, Danesi PR, Burkart W. Properties, use and health effects of depleted Uranium (DU): a general

- overview. *J Environ Radioact.* 2003; 64: 93–112p.
- 5. Cothern CR, Lappenbusch WL. Occurrence of uranium in drinking water. *Health Physics*. 1983; 45: 89–99p.
- 6. Tanner AB. Radon migration in the ground, a supplementary review. In: Gesell TF, Lowder WM (Eds.). *The Natural Radiation Environment III*. Springfield, V.A.: National Technical Information Services, CONF-780422; 1980. 5–56p.
- 7. Somogyi G. *Technical Reports Series No.* 310. Vienna: IAEA; 1990. 1: 229p.
- 8. Virk HS, Kaur H. Estimation of uranium in plant and water samples. *Curr Sci.* 1979; 48: 293–5p.
- 9. Virk HS. Uranium Content Anomalies in Groundwaters of Fazilka District of Punjab (India) for the Assessment of Excess Cancer Risk. *Research & Reviews: Journal of Oncology and Hematology*. 2017; 6(2): 21–6p.
- 10. Virk HS. A Crisis Situation Due to uranium and heavy metal contamination of ground waters in Punjab state, India: a preliminary report. *Research & Reviews: A Journal of Toxicology*. 2017; 7(2): 6–11p.
- 11. Virk HS. Uranium anomalies in groundwater of Sangrur district of Punjab (India) for cancer risk assessment. *Current Science*. 2017; 113(9): 1661–3p.
- 12. Virk HS. Uranium content anomalies in groundwaters of Ferozepur district of Punjab (India) and the corresponding risk factors. *Research & Reviews: Journal of Oncology and Hematology*. 2018: 6(3); 18–24p.
- 13. Virk HS. Uranium Content Anomalies in Groundwater of Barnala District of Malwa Belt of Punjab (India) for the Assessment of Excess Cancer Risk. Research & Reviews: Journal of Oncology and Hematology. 2019; 8(1): 19–26p.
- 14. Virk HS. Uranium Content Anomalies in Groundwater of Patiala District of Punjab (India) for the Assessment of Excess Cancer Risk. *Research & Reviews: Journal of Oncology and Hematology*. 2019; 8(2): 13–19p.
- 15. Virk HS. Assessment of Excess Cancer Risk due to Uranium Content Anomalies in Groundwaters of Bathinda District of Malwa Belt of Punjab (India). *International Journal of Science and Research*. 2019: 8(3); 1228–32p.



- 16. Virk HS, Jakhu R, Bangotra P. Natural uranium content in ground waters of Mohali and Fatehgarh districts of north Punjab (India) for the assessment of excess cancer risk. *Global J Human-Social Science*. 2016; 16(4): 12–17p.
- 17. Virk HS. Measurement of Concentration of Natural Uranium in Ground Waters of Bathinda District (S. Punjab) for the Assessment of Annual Effective Dose. *Global J of Human-Social Science*. 2016; 16(5): 25–9p.
- 18. Singh S, Malhotra R, Kumar J, *et al.* Uranium analysis of geological samples, water and plants from Kulu Area, Himachal Pradesh, India. *Radiat Meas.* 2001; 34: 427–31p.
- 19. Kumar M, Kumar A, Singh S, *et al.* Uranium content measurement in drinking water samples using track etch technique. *Radiat Meas.* 2003; 36: 479–81p.
- 20. Mehra R, Singh S, Singh K. Uranium studies in water samples belong to Malwa region in Punjab by track etching technique. *Radiat Meas*. 2007; 42(3): 441–5p.
- 21. Kaur S, Mehra R. Toxicological risk assessment of protracted ingestion of uranium in groundwater. *Environ Geochem Health*. 2019; 41(2): 681–98p. Available from: https://doi.org/10.1007/s10653-018-0162-4.
- 22. Bajwa BS, Kumar S, Singh S, *et al.* Uranium and other heavy toxic elements distribution in the drinking water samples of SW-Punjab, India. *J Radiat Res and Appl Sci.* 2017; 10(1): 13–19p. doi:10.1016/j.jrras.2015.01.002.
- 23. Rohit M, Anamica KM, Parveen M. A comparative study of uranium concentration using two different analytical techniques and assessment of physicochemical parameters in groundwater. *J Radiat Nucl Appl.* 2018; 3(3): 149–56p.
- 24. Kaur M, Kumar A, Mehra R, et al. Age-dependent ingestion and inhalation doses due to intake of uranium and radon in water samples of Shiwalik Himalayas of Jammu and Kashmir, India. Environ Monit Assess. 2019; 191(4): 224p. Available from: https://doi.org/10.1007/s10661-019-7361-z.
- 25. Report of Central Ground Water Board, Ministry of Water Resources, Government

- of India, North Western Region, Chandigarh. India: Moga District; 2013.
- 26. United States Environmental Protection Agency. *National primary drinking water regulations, radionuclides. Final Rule.* Washington, DC: USEPA; 2000.
- 27. *Human Development Report (HDR)*. Mumbai, India: National Resource Centre for Urban Poverty and All India Institute of Local Self Government; 2009.
- 28. Lee JS, Chon HT, Kim KW. Human risk assessment of As, Cd, Cu and Zn in the abandoned metal mine site. *Environ Geochem and Health*. 2005; 27: 185–91p.
- 29. Health Canada. *Uranium in drinking water.*Document for Public Comment Prepared by Federal Provincial Subcommittee on Drinking Water. Ottawa, ON, Canada: Health Canada; 1999.
- 30. Atomic Energy Regulatory Board (AERB). *Drinking water specifications in India*. Government of India: Department of Atomic Energy; 2004.
- 31. United Nations Scientific Committee on the Effects of Atomic Radiation. *Ionizing Radiation: Sources and Biological Effects*. NewYork, USA: UNSCEAR; 1982.
- 32. International Commission on Radiological Protection (ICRP). *Annals of the ICRP* 23(2). Pergamon Press, Oxford: ICRP Publication: 1993.
- 33. Cantaluppi C, Degetto S. Civilian and military uses of depleted uranium: Environment and health problem. *Ann Chim.* 2000; 90: 665–76p.
- 34. National Research Council. Appendix A: "Radiation as a Carcinogen". In: analysis of cancer risks in populations near nuclear facilities: phase 1. National research council. 2012. Washington, DC: The National Academies Press; 2012. doi: 10.17226/13388.

Cite this Article

Hardev Singh Virk. Uranium Content Anomalies in Groundwater of Moga District of Malwa Belt of Punjab (India) for the Assessment of Excess Cancer Risk. Research & Reviews: Journal of Oncology and Hematology. 2019; 8(3): 18–24p.