Evaluation of Uranium in Organs of Residents from an Uranium-Rich Region using Teeth as Bioindicators

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Abstract

The Uranium extraction and processing plant of INB (Brazilian Nuclear Industries) is in Caetité, a city located in a region hosting the largest Uranium reserve of the country. The degree of Uranium contamination in the Caetité population was investigated before using teeth as bioindicator, where a quite high Uranium concentration was measured in this region, about 160 times higher than the world-wide average. Radiobiological risks are here evaluated from Uranium burdens in organs as skeleton, kidneys, liver, tissues and blood, which were estimated from transfer coefficients and effective internal doses. This was accomplished by means of calculations with the use of the STATFLUX/ICRP approach, plus a set of Uranium transfer rate parameters as function of individual’s age assuming an uninterrupted exposure over a period of 60 years. It was found that U ingestion rates by residents of Caetité are three orders of magnitude higher than worldwide average, indicating that food and water would exhibit high levels of contamination. Calculated effective internal doses range from a minimum of one to a maximum of three orders of magnitude higher than background doses, for blood and bones respectively. The likelihood that this circumstance could lead to serious health problems as e.g. neoplasia of three orders of magnitude higher than background doses, for blood and bones respectively. The likelihood that this circumstance could lead to serious health problems as e.g. neoplasia is addressed. The methodology presented in this work offers subsidies for further studies on environmental pollution by radionuclides.

Introduction

The Southwest region of the Bahia state in Brazil, encompassing the cities of Caetité, Igaporã and Lagoa Real, hosts the largest Uranium reserve of the country (100 kiloton only in uranium). The INB (Brazilian Nuclear Industries) extraction of Uranium and processing plant is located in Caetité (latitude 13°56'36"S and longitude 42°15'32"W).

As pointed out elsewhere [1-4], wastes from the Caetité plant are categorized mostly as: (a) wastes from the mining operations which include waste rock and drainage waters, and (b) wastes from the industrial process, that is, leached ore and effluents from the milling plant. In fact, contamination of groundwater by the waste ponds is of great concern, where elevated levels of $^{238}\text{U}$ have been observed. This issue has been emphatically dealt with by the International Atomic Energy Agency, covering controllable releases to the environment in normal operation that may result from the mining and processing of ores for the extraction of uranium, and discharges of naturally occurring radioactive material [5]. Recommendations and guidance for performing prospective radiological impact assessments for facilities and activities were presented in IAEA 2018-b[6].

The risk of contamination of food (fruit, vegetables, meat, etc.) and drinking water must therefore be considered in the case of this uranium-rich region. Such a scenario motivated an investigation of Uranium concentrations in teeth of residents (5 to 87 years old) of this region, carried out at this Laboratory using a high-resolution inductively coupled plasma mass spectrometer (ICP-MS) [2]. The highest Uranium concentration in teeth was measured from samples belonging to residents of Caetité (median equal to 16 ppb). However, since Uranium concentrations in teeth and bones are similar, particularly for children and young adults, the Uranium body levels in residents of Caetité are, thus, quite high – actually, at least one order of magnitude higher than the worldwide average for bones [2]. Uranium concentrations in organs like bones and teeth are correlated with the amount of Uranium present in ingested food and water. In this regard, it is a matter of major concern the fact that Caetité has approximately 50,000 inhabitants, where half of them live in small farms, using non-treated water provided by wells and by a creek crossing the mining area.
Teeth are good long-term (chronic) exposure monitors, working as reservoirs for many metals incorporated from the environment [3], and their metabolical similarity with bones, as pointed out above, allows the evaluation of contaminants accumulation in the skeleton [2].

Following uptake through the gastrointestinal tract, Uranium is mostly deposited in the skeleton. In fact, bone is one of the most important biological accumulators of Uranium [7]. As in the case of other metals such as Cd, Pb, Si, Sr, etc., Uranium is also subject to stochastic processes of loss from circulation via excretion, redeposition in bone, or soft tissue uptake [8]. Although accounting for only a small fraction of the loss out of the circulation, soft tissues are quite relevant from a radiological viewpoint since they constitute several important organs.

Therefore, estimates of Uranium burdens also in organs other than teeth and bones are necessary for the evaluation of radiobiological risks to populations of the locality here examined, as well as from any other locality presenting radiological contamination.

Actually, Uranium as an alpha particle emitter is a source of ionizing radiation. Public concern, health risks from low-doses and biological responses could be found elsewhere [9].

In this sense, we present estimates of Uranium accumulation in some important and/or vital organs (skeleton, kidneys, liver, tissues and blood) as function of time, obtained by means of a BIOKINETICS code developed at our Laboratory [4] using as input previously measured data for teeth [2].

Mathematical and procedure details involved in the calculations, as well as the calibration of the whole methodology, were developed at this Laboratory and published elsewhere [4,13].

Results and discussion

Biokinetics Model outcome

Bone and teeth are nearly indistinguishable regarding metal incorporation (as e.g. lead and uranium). The younger the individual is, the greater is the similarity between teeth and bones. Moreover, cortical bone constitutes about 80% of the whole skeleton, and it is the bone structure presenting the highest similarity with teeth [14,15]. In this sense, and for the purposes of the present work, we consider the Uranium concentration in the skeleton of residents from the studied location (Caetité City) equal to the Uranium concentration in their teeth, as measured at this Laboratory as function of the individuals age [2] (Figure 1).

Two continuous curves for skeleton were generated by our STATFLUX/ICRP routine (see Material and Methods above) assuming Uranium ingestion rates (a free fitting parameter in the present approach) equal to 2mg-U/day (curve-1 in figure 1) and 12mg-U/day(curve-2 in figure 1), which provided the best set of curves encompassing the scattered data points for teeth. This wide amplitude of the U ingestion rate is closely related to the heterogeneous dispersion of Uranium in the environment.

Actually, uranium worldwide average ingestion is 1 and 2 μg-U/day for food and water, respectively [13]. The fact that our estimates for U ingestion are nearly three orders of magnitude higher strongly suggests that contamination of food (fruit, vegetables, meat, etc.) and drinking water is a fact in this uranium-rich region (the circumstances were discussed in the Introduction).

The rationale of the present study is straightforward.

Figure 1: Data points: Uranium concentrations in teeth from residents of the studied location (Caetité city) as function of their age [2]. Continuous curves: outcomes of biokinetics model calculations for the skeleton (this work), where curve-1 and curve-2 were obtained assuming Uranium ingestion rates of 2mg-U/day and 12 mg-U/d, respectively.
From experimentally obtained U concentration in teeth/bone, plus our STATFLUX/ICRP routine, it was possible infer that U ingestion rates of Caetité residents ranges, which ranges from 2 mg-U/day to 12 mg-U/day – this ingestion rate interval is by itself a quite important and concerning result. Next, by running the STATFLUX/ICRP routine and inputting this same U ingestion rate interval at a chronic ingestion regime, we got estimates of Uranium concentrations in organs of great radiobiological relevance as function of the individuals age (Figure 2).

Transfer coefficients

In fact, food intake is one of the major routes for human exposure to environmental pollutants. This circumstance represents a significant contribution to the mean annual effective dose due to natural sources [16]. Measurement of radionuclide concentrations present in foodstuff, intended for assessment of dose imparted by intake of feed products, was also previously performed at this Laboratory [13].

It is quite important, in this context, the obtaining of a radiobiological parameter which combines Uranium concentration, in a given organ, with the U ingestion rate. In this regard, the so-called transfer coefficient ($T_c$) measures the incorporation efficiency of a contaminant as a function of its daily intake (ingestion rate).

It is defined by

$$T_c = \frac{C_e}{A} \text{ (kg}^{-1}\text{.d)}$$

(1)

where $C_e$ (mg-U/kg-organ) is, in the present study, the equilibrium concentration of Uranium in the organ after a prolonged, chronic daily ingestion, and $A$ (mg-U.d$^{-1}$) is the daily administered amount of Uranium [17].

Taking into account the present results (Figure 1), it is reasonable to consider that the equilibrium concentrations $C_e$ are reached when the individual age is equal or over 60 years, which corresponds to the asymptotic outcome of the present STATFLUX/ICRP approach (curves in Figure 1). By entering these data in Eqn. 1 we obtained $T_c$ for skeleton (Table 1), kidneys, liver and blood (Table 2).

It is well established that $T_c$ is a decreasing function of $A$ which saturates at high ingestion rates, while steeply increasing for much smaller $A$ (see e.g. Figure 2 in Arruda-Neto et al., 2004-b) [18]. Quite consistently in this regard, $T_c = 4.7 \text{ kg}^{-1}\text{.d}$ for $A = 0.7 \text{ mg-U/d}$ (worldwide average for humans), while is an order of magnitude smaller (0.42 kg$^{-1}\text{.d}$) for $A \geq 2$ mg-U/d (present case study – Caetité) – Table 1. In this sense, the fact that $T_c (A = 2 \text{ mg-U/d}) \approx T_c (A = 12 \text{ mg-U/d})$

within calculation uncertainties, is additional indication that Uranium ingestion rates by Caetité residents are quite high. This $T_c$ saturation was also observed for the kidneys simulation. Transfer coefficients for kidneys, liver and blood are presented in table 2. While $T_c$ for kidneys is of the same order of those for bones (Table 1), for blood they are quite small (this issue is retaken in paragraph 3.3.3).

Additionally, the Uranium $T_c$ for bones of the residents of Caetité compares well with those obtained with Beagle dogs [19]. In fact, earlier results suggest that the Beagle may be an appropriate experimental animal from which we can extrapolate data to humans with reference to the percentage of U, Th, and Pu found in their organs [20].

Burden in the organs

Radiological impacts and risk of radiation effects in humans and populations were recently updated by the International Atomic Energy Agency [6]. In this regard, radiological burdens related to Uranium could be evaluated from its concentrations in the organs, while estimates of effective doses allows assessments on impacts and risk as outlined below.

| $A$ (mg-U/d) | 0.7 | 2 | 5 | 12 | 60 | References |
|-------------|-----|---|---|----|----|------------|
| Caetité     | --- | 0.35 ± 0.10 | --- | 0.42 ± 0.10 | --- | this work  |
| Worldwide average | 4.7 | --- | --- | --- | --- | Fermin. et al. 2006 (Table 3) |
| Wistar rats | --- | 0.6 ± 0.1 | 0.2 ± 0.05 | --- | --- | Arruda-Neto, et al. 2006-b |
| Beagle dogs | --- | --- | --- | 0.31 ± 0.08 | 0.14 ± 0.05 | Arruda-Neto et al. 2004-a |

| $A$ (mg-U/d) | 2 | 12 |
|-------------|---|---|
| kidneys     | 0.21 ± 0.04 | 0.18 ± 0.03 |
| liver       | 0.08 ± 0.03 | 0.8 ± 0.2 |
| blood       | (8 ± 2) x 10$^{-4}$ | (6 ± 2) x 10$^{-4}$ |
As detailed calculated elsewhere [18], the average dose imparted by Uranium to a target tissue, only by its emitted alpha particles, is (see eqn. 19 in Arruda-Neto, et al. 2004-b) [18].

\[ D_\alpha = 3.17 \times 10^{-6} x M_e (\text{mg}) x \Delta t (s) \ (\text{Gy} – \text{Gray}) \] (2)

where \( M_e \) is the Uranium mass embedded in the tissue and \( \Delta t \) is the exposure time.

The effective dose from alpha particles in one kilogram of soft tissue for one year of exposure is (see eqns. 20 and 21 in Arruda-Neto, et al. 2004-b) [18]

\[ E_\alpha = 4.2 \times \text{ppm(U)} \ (\text{Sv}/(y.\text{kg}) – \text{Sievert per year per kg}) \] (3)

Results are shown in table 3, which could be compared to the natural background (= 2 mSv/y). It should be emphasized that results obtained from eqn. 3 are lower limits estimative since it was not considered there activities from the U daughters.

**Bones**: One should be much more concerned with the results for U-accumulation in bone, since there are now evidences that through prolonged ingestion Uranium is accumulated and distributed in both bone and bone marrow as experimentally demonstrated elsewhere [18], also indicating that doses delivered to the bone stem cells (in the bone marrow) are intense enough as to induce neoplasia in individuals submitted to chronic ingestion of uranium, after a prolonged period of exposition.

It was estimated, in the present case study, that bones of Caetité city residents are annually exposed to internal effective doses two to three orders of magnitude higher than the natural background (Table 3). It is worth to recall that bones are the organ storing about 80% of the whole Uranium incorporation.

**Kidneys and liver**: The internal effective doses in kidneys and liver are very high too, but our calculations revealed an aggravating circumstance, in the sense that Uranium concentrations in these organs saturate at ages of 10 and 20 years, respectively (all results shown in tables 1 to 3 refer to 60 years old residents). This is of great concern too because, in this case, severe radiological burdens would be inflicted to children and young adults.

Also quite harmful, particularly for the youngest residents of Caetité, is the nephrotoxicity of Uranium and its compounds as uranyl salts – see e.g. the summary document ATSDR [21]. A compilation of Uranium burdens in kidney published by Chen, et al. [22] shows that after 25 years of chronic ingestion, the Uranium burden in kidneys reaches its final equilibrium for all age groups.

**Blood**: This organ presented the smallest Uranium concentrations, but in relative terms its internal effective doses are up to one order of magnitude higher than the natural background (Table 3). Moreover, and much more alarming, the calculations indicate that Uranium transfer to blood saturate quite swiftly at ages between 2 and 3 years (Figure 2).

Although these predictions could be of use for dose evaluation, it is important to point out that Uranium concentration in blood and soft tissues is not well described in literature. The majority of data feeding ICRP models were obtained from acute injection studies, and because of this, results from ICRP model calculations for chronic regime must be analyzed and used cum grano salis. The present calculations, for instance, assume that flow parameters and organ’s transfer coefficients are constant during a 60 years life span, and that they do not depend on the physiological tolerance to long exposures.

**Conclusion**

Estimates of U ingestion by residents of Caetité are three orders of magnitude higher than worldwide average ingestion, a circumstance strongly suggesting that food and water from this Uranium-rich region would exhibit high levels of contamination.

Uranium transfer coefficients of bones and kidneys saturate for the estimated ingestion range (2 – 12 mg-U/d), a strong indication that Uranium ingestion rates by Caetité residents are very high.

Effective internal doses are from one (in blood) to three (in skeleton and kidneys) orders of magnitude higher than the background doses. Such a circumstance could lead to serious health problems as neoplasia.

As a byproduct, this work could subside further studies toward the identification of the driving factors in environmental pollution, associated with possible risks to the public health.

**Acknowledgement**

The authors acknowledge the Brazilian Agencies FAPESP and CNPq for partially supporting this work.

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