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# Toxicity of rare earth elements: An overview on human health impact

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Rare earth elements (REEs) are metals including the 15 lanthanides together with Yttrium and Scandium. China is the leading country in their exploitation and production (~90%). REEs are necessary for the production of several technological devices. This extended use of REEs has raised concerns about human health safety. In this review, we investigated the hazard of REEs to human health and the main gaps into the knowledge like as the need to develop further focused research activity. We categorized the research papers collected into eight main sections: environmental exposure, association of REEs with health problems, exposure to REEs due to lifestyle, REE exposure through the food chain, Gd contrast agents causing health problems, occupational REE exposure, and cytotoxicity studies of REEs. This review provided information about the exposome of REEs (the exposure of REEs to the human body), the existing research data, and the gaps that require attention and must be further investigated. More than one third of the literature about REE toxicity to human health concerns their cytotoxicity to human cell lines, while hair, blood serum and blood are the most studied matrices. The main results evidenced that REEs can enter human body via several routes, are associated with numerous diseases, can cause ROS production, DNA damage and cell death, and are more toxic to cancer cells than normal cells.

## KEYWORDS

rare earth elements, human health, toxicity, exposome, accumulation

## 1 Introduction

The term rare earth elements (REEs) refers to a group of metals that consists of the 15 lanthanides [Lanthanum (La), Cerium (Ce), Praseodymium (Pr), Neodymium (Nd), Promethium (Pm), Samarium (Sm), Europium (Eu), Gadolinium (Gd), Terbium (Tb), Dysprosium (Dy), Holmium (Ho), Erbium (Er), Thulium (Tm), Ytterbium (Yb), and Lutetium (Lu)] generally along with Scandium (Sc) and Yttrium (Y)] (Wall, 2014). These metals have similar ionic radii and physicochemical properties and are sub-divided into

three groups according to their atomic number and masses (Brown et al., 1990): La, Ce, Pr, Nd, and Pm are called light rare earth elements (LREEs); Sm, Eu, Gd, Tb, Dy, and Ho are called middle rare earth elements (MREEs); Er, Tm, Yb, and Lu are called heavy rare earth elements (HREEs).

China is the leader country in the production of REEs, having provided since recently more than approximately 90% of the global REE supply (Long, 2011). However, China's development in REE production has been decreased during the last years, resulting in production of only 60% of the global REE supply in 2021 (USGS, 2022). REEs are used in several technological applications since the manufacturing of many devices relies on their utilization (Du and Graedel, 2011; US Environmental Protection Agency, 2012; Du and Graedel, 2013; Pagano et al., 2019), with the glass industry being the number one consumer of REE raw materials (Van Gosen et al., 2017).

Human exposures to REEs occur due to anthropogenic activity in many ways. Atmospheric particulates undergo an REE-enrichment by anthropogenic sources and are influenced by strong winds (Wang and Liang 2014). Workers in recycling facilities are exposed to significant amounts of REEs through inhalation, ingestion, and skin contact (Shin et al., 2019). REEs can be transferred to the human body also through the food chain since they are used as feed additives to improve animal health and production without affecting their safety (He et al., 2001; He et al., 2010; Abdelnour et al., 2019), and in pesticides, herbicides, and fertilizers to improve crop yield and quality (Pang et al., 2002). However, unlike animals, an REE accumulation occurs not only in soil (Naccarato et al., 2020) but also in the roots and tops of plants (Xu et al., 2002). As a result, many concerns about human health due to REE exposure have been raised.

Some review papers have investigated the hazards of REEs to human health based on research of REE toxicity both to humans and generally to several biota (Hirano and Suzuki, 1996; Cassee et al., 2011; Pagano et al., 2015a; Pagano et al., 2015b; Rim, 2016; Gwenzi et al., 2018). In a recent review paper, we investigated the environmental species sensitivity distribution of REEs and calculated the environmental hazard concentration of each REE for the 5% (HC5) and the 50% (HC50) of the species, including only research concerning acute toxicity and bioaccumulation of REEs to animals, plants and microorganisms (Albarano et al., 2022).

In the present review, we attempt to investigate the impacts of REEs on human health and find out the main knowledge gaps. Through this literature review, we wish to accomplish three critical issues: i) to gather all the studies that investigated REE toxicity to the human body; ii) to find out the gaps concerning the information about the exposure and toxicity of REEs directly to human health, and iii) to investigate the exposome of REEs directly to human health, i.e. to understand by which routes REEs can enter the human body, its tropism, and the level of toxicity they can exert.

## 2 Data collection and comparison

We used the following main keywords for searching activities: rare earth elements, human, toxicity, accumulation. The literature search was made up to 28<sup>th</sup> July 2022 using SCOPUS, PubMed and Google Scholar as the main search engines. We found 135 articles mainly about accumulation due to REE mining, risk assessment to human health, cytotoxicity to human cell lines. They were categorized into seven sections: environmental exposure, association of REEs with health problems, exposure to REEs due to daily habits, REE exposure through the food chain, Gd contrast agents causing health problems, occupational exposure to REEs, cytotoxicity studies of REEs. Table 1 is demonstrating a schematic summary of this review paper. The relative abundance (%) of papers for each section studied in response to REEs are shown in Figure 1.

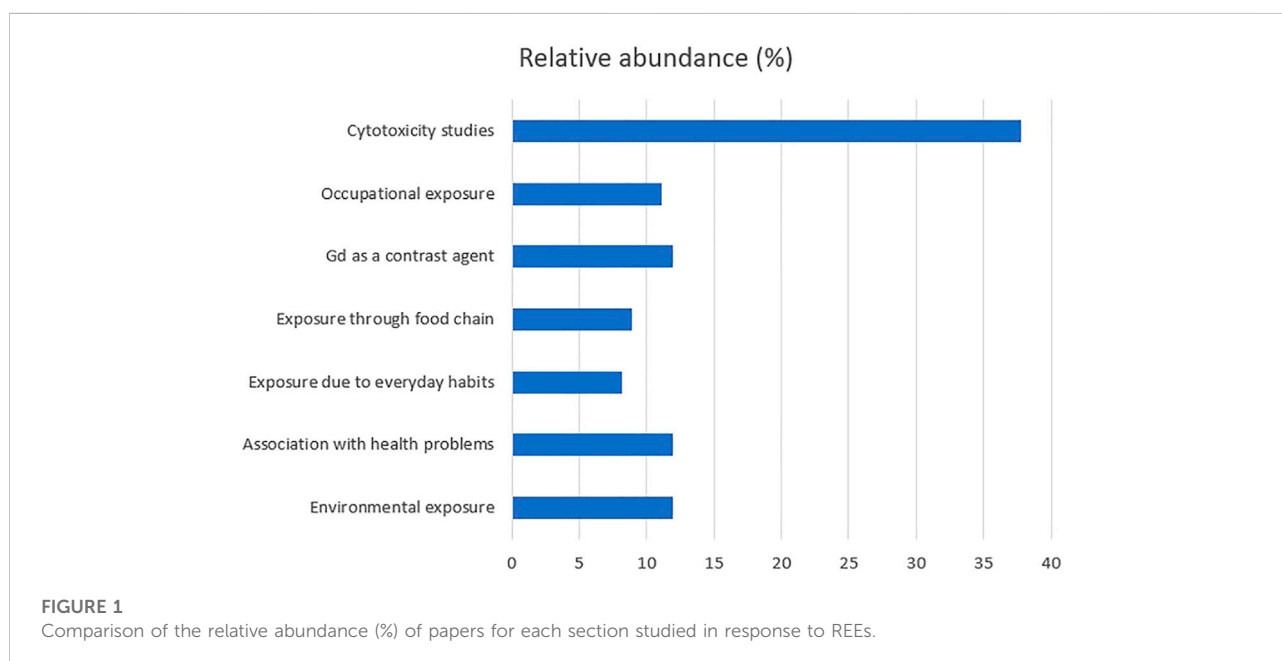
Comparing the sections studied about REE toxicity, 37.8% of papers included cytotoxicity studies, followed by a 11.9% for environmental exposure, association of REEs with health problems and Gd-based contrast agents. The comparison among the analyzed biological matrices is shown in Figure 2. Hair was the most analyzed matrix (19%), followed by blood serum (16%) and blood (13%). Comparing the percentage of the articles considering the investigated REEs, Ce was present in 65.9% of the papers, La in 48.9%, Gd in 42.2%, while Nd in 37%. The complete data is shown in Figure 3. Comparing the different types of Ce for which we found information, more than half of them (52%) were reports in which Ce levels were traced in biological matrices, and studies about the cytotoxicity of CeO<sub>2</sub> nanoparticles (NPs) were 24%. Comparison of the relative abundance (%) of each type of Ce studied is shown in Figure 4.

## 3 Environmental exposure

Some studies showed that REE mining affected the residents in the areas near the mining sites by analyzing the REE levels in human samples. Urine REE concentrations of people living near Baiyun Obo mining area were significantly higher than in residents outside the mining area (Hao et al., 2015; Liang et al., 2018). Other studies have investigated the REE-pollution through analyses of hair samples. In the study of Wei et al. (2013), scalp hair of people living near a mining site in Inner Mongolia contained significantly higher REE levels compared to people from the control area. In a similar study, the hair of children living in Shang and Liao villages, near a mining site, contained higher REE levels versus children from reference areas (Tong et al., 2004). As reported by Yu et al. (2007), REE blood levels were higher in people living near the mining site of Xunwu. They had some effects on the peripheral blood mononuclear cells, such as increasing telomerase activity, promoting the diploid DNA replication, and increasing the

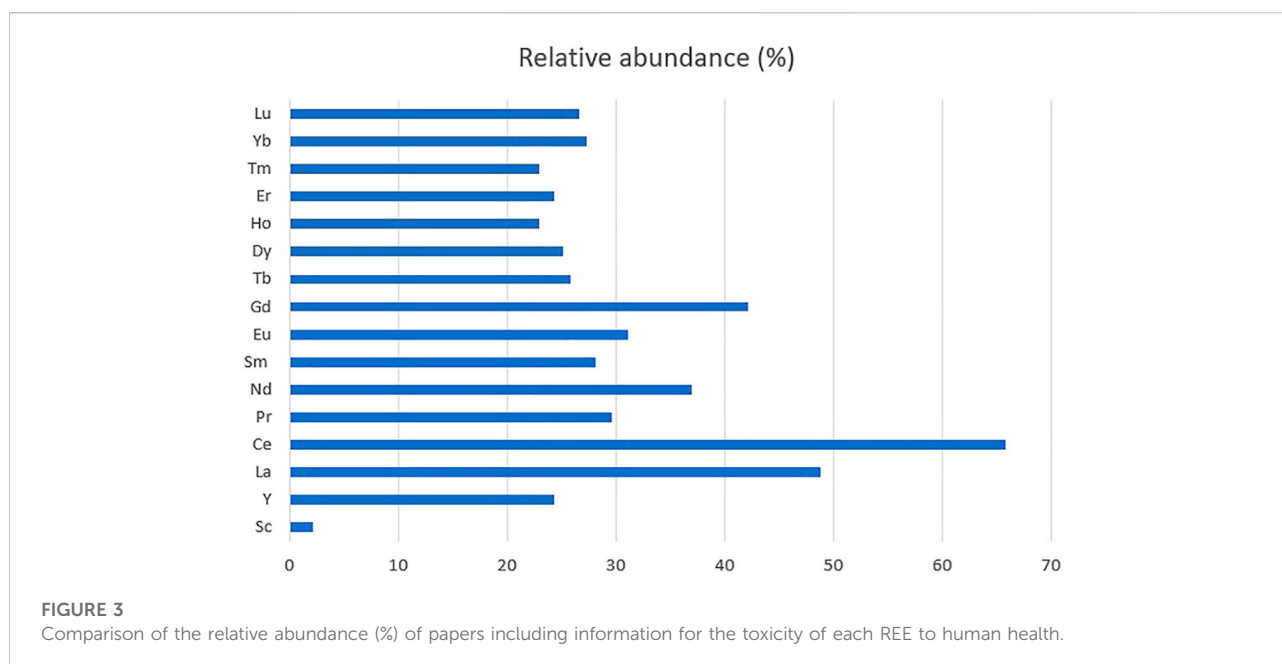
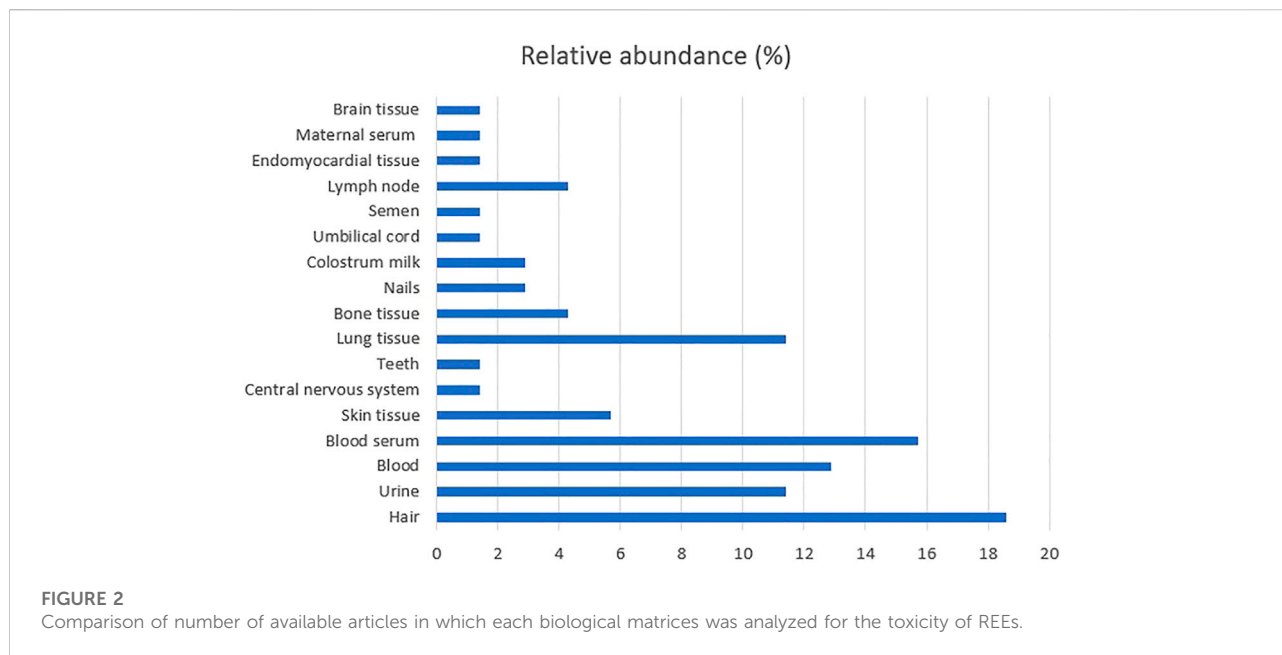
**TABLE 1** Schematic summary table demonstrating the different sections that each REE has been studied about at least once, and the relative abundance (%) of papers including information for the toxicity of each REE to human health.

Element	Sections studied	Articles percentage (%)
Sc	Environmental exposure, Food chain	2.2
Y	Environmental exposure, Health disorders, Lifestyle, Food chain, Occupational exposure, Cytotoxicity studies	24.4
La	Environmental exposure, Health disorders, Lifestyle, Food chain, Occupational exposure, Cytotoxicity studies	48.9
Ce	Environmental exposure, Health disorders, Lifestyle, Food chain, Occupational exposure, Cytotoxicity studies	65.9
Pr	Environmental exposure, Health disorders, Lifestyle, Food chain, Occupational exposure, Cytotoxicity studies	29.6
Nd	Environmental exposure, Health disorders, Lifestyle, Food chain, Occupational exposure, Cytotoxicity studies	37.0
Sm	Environmental exposure, Health disorders, Lifestyle, Food chain, Occupational exposure, Cytotoxicity studies	28.1
Eu	Environmental exposure, Health disorders, Lifestyle, Food chain, Occupational exposure, Cytotoxicity studies	31.1
Gd	Environmental exposure, Health disorders, Lifestyle, Food chain, Gd contrast agents, Occupational exposure, Cytotoxicity studies	42.2
Tb	Environmental exposure, Health disorders, Lifestyle, Food chain, Occupational exposure, Cytotoxicity studies	25.9
Dy	Environmental exposure, Health disorders, Lifestyle, Food chain, Occupational exposure, Cytotoxicity studies	25.2
Ho	Environmental exposure, Health disorders, Lifestyle, Food chain, Occupational exposure, Cytotoxicity studies	23.0
Er	Environmental exposure, Health disorders, Lifestyle, Food chain, Occupational exposure, Cytotoxicity studies	24.4
Tm	Environmental exposure, Health disorders, Lifestyle, Food chain, Occupational exposure, Cytotoxicity studies	23.0
Yb	Environmental exposure, Health disorders, Lifestyle, Food chain, Occupational exposure, Cytotoxicity studies	27.4
Lu	Environmental exposure, Health disorders, Lifestyle, Food chain, Occupational exposure, Cytotoxicity studies	26.7



percentage of cells in the S-phase and the G2/M phase. The blood and hair of people living near a large-scale mining site in southwest Fujian Province, China, were found enriched with REEs but much more with LREEs (Li et al., 2014). A correlation between the concentration of REEs in soil and human samples was found in Hezhang County, China, where REEs levels (mostly La, Ce, and Nd) in the soil near a smelting area were higher than

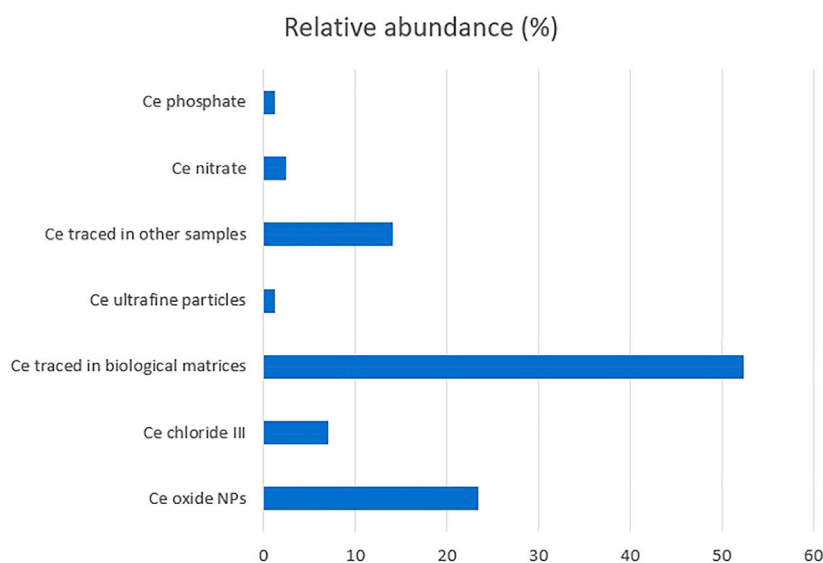
those in the soil near a mining area, and the same applied for urine and hair of residents near these two areas (Meryem et al., 2016). Tian et al. (2019) found an enrichment of Ce in dust from mining areas and the potential of causing health problems through inhalation. According to the Life Cycle Assessment of Wang C. et al. (2020), the most significant impact of  $\text{Sc}_2\text{O}_3$  production from Bayan Obo mine is the non-cancerous human



toxicity. All these studies indicated that the REE mining process influenced the environment in mining areas and the people living nearby.

However, REEs environmental exposure can occur, not only due to REE mining but also to other factors. Ugandan children presented significantly higher REE concentrations in their primary teeth than the children from the United Kingdom (Brown et al., 2004) due to an environmental exposure. The levels of REEs were significantly higher in the blood of

Moroccans than in Canarians, possibly related to the improper management of e-waste (Henríquez-Hernández et al., 2018). Lanthanum and Ce levels were higher in resident serum from an e-waste area and were associated with higher thyroid-stimulating hormone (TSH) (Guo et al., 2020). Hollriegl et al. (2010) associated the increased Ce levels in the serum of women in Madrid with the high traffic volume in this town. The accumulated concentration of REEs in the rib bone tissue was found to be age dependent by Zaichick et al. (2011). Street dust in



**FIGURE 4**

Comparison of the relative abundance (%) of each type of Ce studied about its toxicity to human health.

industrial cities is enriched with REEs mainly due to heavy local soil and less due to anthropogenic pollution, however, the existing REE levels were not found as a threat to cancer occurrence (Sun et al., 2017). Analysis of soil samples from a tropical area and a subsequent health risk assessment from Da Silva Ferreira et al. (2022) showed that REEs represent less than 20% of the metal content and the risk for cancer due to these metals is very low.

## 4 Rare earth elements-associated health disorders

Some studies have investigated the association between REEs and specific diseases, with different results. Brown et al. (2004) concluded that the higher Ce concentration in primary teeth of Ugandan children, compared to children from the United Kingdom, failed to be associated with endomyocardial fibrosis. Huo et al. (2017) suggested that although LREEs are higher in the hair of women with neural tube defects (NTDs) affected pregnancies, they are not associated with the risk of NTDs, even if Wei et al. (2020) found that the REEs levels were higher in the maternal serum of pregnant women affected by NTDs and that the risk for NTDs increases with REE concentrations, especially La. Henríquez-Hernández et al. (2017) have pointed out the possibility of REEs playing a role in the development of anemia since they were found to be higher in the blood of sub-Saharan immigrants with anemia and independent of Fe levels. De Vathaire et al. (1998) identified an excess of mortality due to lung and bladder cancer in a French

population living near a site where Ce was extracted. Similarly, Kutty et al. (1996) found that deposits of monazite elements, like Ce, increase the rate of occurrence of endomyocardial fibrosis (EMF). In utero exposure to REEs, mainly La and Nd, increase the risk of orofacial clefts to infants (Liu L. et al., 2021). Exposure of pregnant women to Ce and Yb has decreased TSH levels in infants (Liu et al., 2019). REEs may accumulate in the cerebral cortex by long-term environmental intake of a small amount and cause subclinical damage (Zhu et al., 1997). REEs, especially La, Ce, and Gd, are significantly higher in brain-tumor tissues of patients with astrocytoma compared to normal brain tissues (Zhuang et al., 1996). People living near high-REE background areas have illnesses like indigestion, diarrhea, abdominal distension, anorexia, weakness, and fatigue. They have been shown to have lower levels of total protein, globulin, albumin, and serum-glutamic pyruvic transaminase (SGPT) and higher levels of IgM in their blood serum (Zhu et al., 2005). A high amount of REEs in hair has been associated with low levels of Ca in hair and a high risk of hypertension among housewives (Wang B. et al., 2017). Lanthanum levels in the serum of women undergoing *in vitro* fertilization-embryo transfer have been associated with a 30% increase in clinical pregnancy failure, a 230% increase in preclinical spontaneous abortion, and a negative correlation with the number of good-quality oocytes (Li et al., 2021). Cerium levels in toenails have been associated with an increased risk of acute myocardial infarction (Gómez-Aracena et al., 2006). In the study of Medina-Estévez et al. (2020), Ce levels were higher in control people group than in the subjects, and so, inversely associated with the risk of stroke. Increased REEs in the hair of housewives

have been correlated with indoor air pollution and the risk of hypertension by Wang et al. (2018).

## 5 Rare earth elements exposure as related to lifestyle

Everyday habits can also affect the concentration of metals, including REEs, in the human body. Marzec-Wroblewska et al. (2015) found that the concentrations of La, Ce, Eu, and Gd were significantly lower in the non-drinkers' and short-term smokers' semen than in the semen from drinkers and long-term smokers. Moreover, Zhang et al. (2020) evidenced that smoking was positively associated with Dy, Er, and Yb levels, whereas drinking habits showed no significant effect on REE levels, while La, Ce, Pr, and Nd were higher in hair and nails of smokers (Rodushkin and Axelsson, 2000). Poniedzialek et al. (2017) found that Nd concentration was significantly higher in the colostrum milk of smoking women compared to those who had never smoked. Zhang et al. (2020) confirmed also that REE levels are higher in biological matrices of smokers, but not in drinkers. Housewives exposed to REEs due to passive smoking presented elevated levels of REEs in their hair according to Na et al. (2022).

The rolling paper, especially a flavored paper, is the part of the cigarette that contains the highest amount of hazardous metals, including REEs, while black tobacco contains more REEs compared to the other tobacco types, and so, smokers are exposed to REEs mainly due to the rolling paper, as well as to the tobacco. (Zumbado et al., 2019). LREEs are contained in the flint of lighters, and even short-term contact with the flint during the light-up could result in exposure to LREEs (Rodushkin and Axelsson, 2000; Kastury et al., 2020).

Tobacco smoking has been rated as one of the most significant sources of REEs in indoor inhalable particles. Cerium and La levels significantly increased in the households of smokers and hospitality pubs like restaurants, pubs, dance clubs, offices, and cafeterias due to tobacco smoke, and imply a threat to the human respiratory system (Slezakova et al., 2009; Böhlandt et al., 2012). Increased indoor Ce and La have also been associated with indoor smoking and respiratory symptoms in children (Drago et al., 2018). Although REEs in PM<sub>10</sub> are higher in smokers' households, their dissolution in PM<sub>10</sub> and inhalation bioaccessibility are higher in non-smokers' households (Kastury et al., 2020).

E-cigarettes can be considered also as a source of an increased REE intake. Badea et al. (2018) found that REE levels are higher in serum of e-cigarette users than cigarette users, and the longer someone uses e-cigarettes, the higher the levels of Ce and Er in their blood serum. Ytterbium levels were found higher in teeth of people wearing porcelain bridges, and so the latter can be considered as well as a another potential source of REEs intake (Rodushkin and Axelsson, 2000).

## 6 Rare earth elements exposure through the food chain

REEs can be transferred in the human body also via the food chain. In the study by Zhang et al. (2020), centenarians who consumed smoked and pickled food, egg, milk, and high amounts of salt had higher REE levels in their hair. REE oxides (LREOs) in Oolong tea consider a negligible threat for human health as its consumption does not exceed the accepted daily intake (ADI) of REE levels (Guo et al., 2015). Cereals, especially wheat, and vegetables, especially leaf vegetables, bioaccumulate significantly high amounts of LREEs when growing near mining areas; however, these concentrations are much lower than the ADI (Zhuang et al., 2017a; Zhuang et al., 2017b). Guo et al. (2012) investigated the REE levels in the most consumed foods in China and concluded that REEs are at a shallow concentration and do not exert a hazard for human health. Freshwater and marine fish in Shandong Province, the largest center for fishery production and processing in China, contain a much higher amount of LREEs versus HREEs; however, the estimated daily intake (EDI) of REEs through fish food were significantly lower than the ADI; therefore, its consumption presented little risk to human health (Yang et al., 2016). The health risk assessment of Li et al. (2013) concluded that vegetable consumption would not exceed the EDI values for REEs. The same applies for the health risk assessment of Shi et al. (2022) concerning fruit and vegetable consumption. REE levels are higher in the aquatic than the terrestrial food, get lower at higher trophic levels in the environment and their human intake at present is acceptable (Dai et al., 2022). To the best of our knowledge, the studies, up to now, about the REEs level in food showed that they are present in low concentrations and can hardly exert problems in human health. Squadrone et al. (2019) analyzed REEs level in several terrestrial and marine matrices and organisms. Although the highest REE levels were found at low trophic levels in both environments, they suggested a biomonitoring of these elements for possible cumulative effects to human health, since they can be transferred to human body through the trophic chain.

Zhang et al. (1999) indicated that exposure to REEs through the food chain could result in low total serum protein (TSP), albumin,  $\beta$ -globulin, glutamic pyruvic transaminase, serum triglycerides, and immunoglobulin, but high cholesterol. Cerium concentration was found higher in endomyocardial samples of people dying from endomyocardial fibrosis and has been associated with the Ce levels in leafy vegetables and root tubers like tapioca and yam (Valiathan et al., 1986).

## 7 Gd contrast agents are causing health problems

Gadolinium-based contrast agents (GBCAs) have been used for years in magnetic resonance imaging (MRI) and magnetic

resonance angiography (MRA). In the first research about the safety of these agents to human health, no side effects were caused to patients. Carr et al. (1984) found that a small intravenous administration of Gd-diethylenetriamine pentaacetic acid (Gd-DTPA) could enhance the contrast of tumors during an MRI operation, causing no side effects to patients. However, subsequent studies reported that these agents are not completely safe for human health. GBCAs can cause acute renal failure in patients with underlying chronic renal insufficiency (Sam et al., 2003). Chien et al. (2011) found out that potential acute kidney injury is after administration of a GBCA, under sepsis condition, at the dose given for MRI and MRA examinations in patients with renal impairment. Ergun et al. (2006) reported that an acute renal failure (ARF) can occur after a GBCA administration in patients with moderate to severe chronic renal failure. Risk factors for ARF after Gd toxicity include diabetic nephropathy and low glomerular filtration rate (GFR). Similarly, Thomsen (2004) reported that GBCAs can induce nephropathy even at doses below 0.2 mmol/kg body weight (a usual dose of GBCA for MRI) in patients with multiple risk factors. Gd-DTPA could play a role in developing nephrogenic fibrosing dermopathy (NFD) in renal disease patients undergoing MRA (Grobner, 2006). There is a high risk of nephrogenic systemic fibrosis in patients with chronic kidney disease at stage 5 due to exposure to gadodiamide during MRI (Rydahl et al., 2008). Gadolinium remained in human bone tissue after administration of a standard clinical dose (0.1 mmol/kg) of Omniscan or Prohance a few days before surgery to patients undergoing hip replacement (Gibby et al., 2004; White et al., 2006). GBCAs can induce central torso and peripheral arm and leg distribution pain in patients (Semelka et al., 2016). Gadovist, a GBCA was reported to cause fibromyalgia in one patient following repeated administrations (Lattanzio and Imbesi, 2020).

A few studies have investigated the potential toxicity of GBCAs to human health by a number of cytotoxicological experiments. Four GBCAs, Gadovist, Magnevist, Multihance, and Omniscan, were not toxic to the embryonic fetal lung fibroblast cell line Hel-299, which rapidly recovered after the initial antiproliferative effects of the agents (Wiesinger et al., 2010). Omniscan increased the proliferation and the levels of the matrix metalloproteinase-1 (MMP-1) and the inhibitor of metalloproteinases-1 (TIMP-1) to monolayer culture of human dermal fibroblasts and to organ culture of human dermal skin, but they did not stimulate the proliferation of epidermal keratinocytes (Bhagavathula et al., 2009; Varani et al., 2009). Magnevist, Multihance, and Prohance (i.e., another GBCAs) increased the proliferation and levels of MMP-1 and TIMP-1 in human dermal fibroblasts like Omniscan did (Varani et al., 2009). Edward et al. (2010) examined the effect of several GBCAs on human skin fibroblasts. Some of them stimulated cell proliferation and hyaluronan production, while no GBCA could affect collagen synthesis. Omniscan increased

MMP-1 and TIMP-1 in a human skin organ culture, but reduced the collagenolytic activity (Perone et al., 2010). Gadolinium chloride had similar effects with the GBCAs, having no effect on epidermal keratinocytes, enhancing the proliferation and the levels of MMP-1 and TIMP-1, and having no effect on type I procollagen in fibroblasts via activation of the PDGF receptor signaling pathway (Bhagavathula et al., 2010).

## 8 Occupational exposure to rare earth elements

The content of REEs in the hair of miners, particularly light REEs (La, Ce, Pr, and Nd), was much higher than the values in the hair of non-miners from both mining area and control area (Wei et al., 2013). Liu et al. (2015) found higher REE hair levels in miners working at the Baiyun Obo mining area (China), affecting the levels of proteins in serum that participate in various biological processes. Increased La levels were also found in the lung tissue of deceased smelter workers (Gerhardsson and Nordberg, 1993). Elevated REE levels were found in the hair of miners, which were associated with Fe and Ca in hair and a lower bone mineral density at lumbar vertebrae, femoral neck, greater trochanter, and intertrochanter (Liu H. et al., 2021). These studies suggested that the miners can be exposed to increased REE levels in their occupational environment.

Rare earth pneumoconiosis is an uncommon occupational disease caused by the inhalation of REE-containing dust and has been associated with dendriform pulmonary ossification (Yoon et al., 2005). Some early studies showed that workers active in movie projection were exposed to REEs, produced in the fumes and dust of arc carbon lamps that contained a significant REEs amount. A movie projectionist who had approximately 25 years of occupational exposure to carbon arc lamp fumes was found to be exposed to La, Ce, and Nd, with evidence of their redistribution throughout the reticuloendothelial system. However, the exposed did not suffer from pneumoconiosis, as there were no respiratory symptoms or radiographic or histological pulmonary changes attributable to a progressive REEs accumulation (Waring and Watling 1990). Porru et al. (2000) investigated another movie projectionist exposed for 12 years to REE-containing dust from cored arc light carbon electrodes. The exposed presented interstitial lung disease, emphysema, and a severe obstructive impairment with a marked decrease of carbon monoxide diffusion capacity. The biopsy confirmed the diffuse interstitial lung fibrosis and significantly higher concentrations of Ce, La, Nd, Sm, Tb, and Yb compared to the unexposed control subjects. Three other cases of photoengravers, exposed for many years to the fumes and dust from carbon arc lamps, suffered from pneumoconiosis and the clinical analyses showed evident high REEs concentration in the pulmonary and lymph node biopsies, suggesting a relationship between the pneumoconiosis and the

occupational exposure to REE dusts, and the need to define air exposure threshold limit values of REEs to prevent workers occupational exposure (Sabbioni et al., 1982; Vocaturo et al., 1983).

Rare earth elements are biopersistent and can remain in the human body many years after occupational exposure (Pairon et al., 1994; Pairon et al., 1995). High Ce, La, and Nd levels were found in the lung tissue of a subject having worked in printing shops and exposed to carbon-arc lamp emissions (Dufresne et al., 1994). REE dust exposure in a lens grinder was reported by Yoon et al. (2005), who found pneumoconiosis associated with dendriform pulmonary ossification. The urines of workers producing ultrafine and nano-sized particles containing Ce and La oxide; especially those at separation and packaging locations presented greater levels of La and Ce (Li et al., 2017). In the study by Li et al. (2016), the REE levels in the urine of workers that manufacture cerium and lanthanum oxide ultrafine and nano-sized particles were more than five times higher versus controls. Their results, however, suggested that only the urinary levels of La, Ce, Nd, and Gd among the exposed workers were significantly higher than the levels measured in the control subjects. High REE levels in the blood have been associated with significant DNA oxidative damage and high blood concentration of Cr in chromate exposed workers (Bai et al., 2019). The urines and blood of e-waste recyclers contain more Eu, but for La, this applies only for urines (Takyi et al., 2021). An uptake of REEs has been found in triple negative breast cancer cells of a woman having worked in the ceramic industry (Roncati et al., 2018).

## 9 Cytotoxicity of rare earth elements

Nanoparticles (NPs) of Ce can significantly mitigate reactive oxygen species (ROS) production and DNA oxidative damage in the BEAS-2B cell line (Rubio et al., 2016). The co-exposure of CeO<sub>2</sub> NPs and diesel exhaust on a sophisticated *in vitro* 3D co-culture model of the human epithelial airway barrier indicates that a short-term co-exposure led to result in adverse effects, long-term co-exposure caused undesired effects to human respiratory health (Steiner et al., 2012). CeO<sub>2</sub> NPs induced mitochondrial damage and overexpression of apoptosis to human peripheral blood monocytes (CD14<sup>+</sup>) (Hussain et al., 2012) and human lung adenocarcinoma (A549) (Mittal and Pandey, 2014). CeO<sub>2</sub> NPs protected human colon cells (CRL 1541) from radiation by reducing ROS production and increasing the expression of superoxide dismutase 2 (SOD2) (Colon et al., 2010). CeO<sub>2</sub> NPs enhanced the antitumor effect of anthracycline doxorubicin, one of the most effective anti-cancer drugs, in human melanoma cells (A375), and even protected human dermal

fibroblasts from doxorubicin-induced cytotoxicity (Sack et al., 2014). Nanoparticles of CeO<sub>2</sub> did not cause any cytotoxicity to human hepatic carcinoma cell line (HepG2) and human lung carcinoma cell line (A549) during a short-term exposure (24 h) but only during a long-term period (10 days); moreover, they did not cause any cytotoxicity to cells from human colon carcinoma (CaCo<sub>2</sub>) at any of the investigated times (De Marzi et al., 2013).

Diesel particulate matter containing CeO<sub>2</sub> and Fe(C<sub>5</sub>H<sub>5</sub>)<sub>2</sub> NPs decrease the viability of human-type II cell alveolar epithelial cells (A549) (Zhang and Balasubramanian, 2017). Nanoparticles of CeO<sub>2</sub> enhanced the proliferation of human keratinocytes and human microvascular endothelial cells (Chigurupati et al., 2013). Nanoparticles of CeO<sub>2</sub> were not found to be genotoxic to eye lens epithelial cells (ATCC-LGC CRL-11421), suggesting a potential safe use in drug delivery for treating cataract (Pierscionek et al., 2010). They were not toxic to the cell lines T98G (from a human glioblastoma multiform tumor) but caused cell death to BEAS-2B cell line (Park et al., 2008) and human bronchoalveolar carcinoma derived cell line (A549) (Lin et al., 2006) by an apoptotic process, mainly by increasing the levels of ROS and decreasing the levels of GSH and  $\alpha$ -tocopherol. In the study of Alpaslan et al. (2015), the dextran-coated nanoceria killed much more effectively the osteosarcoma cell line MG-63 (ATCC CRL-1427) than the osteoblast noncancerous cell line (PromoCell, C-12720), suggesting that nanoceria can promisingly be used for treating bone cancer without the appearance of adverse effects to healthy bone cells. CeO<sub>2</sub> NPs showed a rapid uptake from human keratinocyte cells (HaCaT) and co-localized with mitochondria, lysosomes, endoplasmic reticulum, cytoplasm, and nucleus, suggesting that they act as cellular antioxidants in multiple compartments (Singh et al., 2010). CeO<sub>2</sub> NPs are taken up into caveolin-1 endosomal compartments by BEAS-2B cells without causing inflammation or cytotoxicity (Xia et al., 2008). Tarnuzzer et al. (2005) showed that CeO<sub>2</sub> NPs could protect almost 100% of the normal breast epithelial cell line CRL8798 and simultaneously did not protect the breast carcinoma cell line MCF-7. According to the study by Celardo et al. (2011), in two tumor cell lines, the human tumor monocytes U937 and the human tumor T lymphocytes Jurkat, the intracellular antioxidant effect of CeO<sub>2</sub> NPs is due to their anti-apoptotic and pro-survival activity. They also found that doping of CeO<sub>2</sub> NPs with increasing concentrations of Sm<sup>3+</sup> blunted these effects by decreasing Ce<sup>3+</sup> and not affecting oxygen vacancies, demonstrating that Ce<sup>3+</sup>/Ce<sup>4+</sup> redox reactions might be responsible for the biological properties of nanoceria.

In Yongxing et al. (2000), La and Gd were indicated as possible mutagenic factors for human primary peripheral lymphocytes. The compounds LaCl<sub>3</sub> and CeCl<sub>3</sub> inhibited the growth of the leukemic cell lines HL-60 and NB4, respectively, and simultaneously LaCl<sub>3</sub> had no inhibitory effect on normal bone marrow hematopoietic progenitor cells (Dai et al., 2002). Lanthanum and Ce nitrates decrease the proliferation of the



human osteoblast MG63 cell line when combined with radiation but not alone (Iwahara et al., 2019). Lanthanum oxyiodide nanosheets with low dose of doxorubicin, an anticancer drug, can kill more effectively the A375 cells compared to the doxorubicin alone (Xu et al., 2020). Nd<sub>2</sub>O<sub>3</sub> and La<sub>2</sub>O<sub>3</sub> NPs are cytotoxic, while Gd<sub>2</sub>O<sub>3</sub> and CeO<sub>2</sub>-Gd NPs enhance ROS production to the U-87 MG tumor cell line (Lu et al., 2019). Lanthanum is cytotoxic through the necrosis pathway and genotoxic by increasing ROS production in Jurkat cells and human peripheral lymphocytes (Paiva et al., 2009). The ionic forms La<sup>3+</sup> and Tb<sup>3+</sup> caused disruptions on potassium channels on the plasma membrane of HEK293 cells (Wang L. et al., 2017). Lanthanides can decrease the ATP production and the cell viability of the HepG2 cells, and also can decrease the mitochondrial membrane potential most probably by interfering with the calcium regulation due to the similar ionic radius (Kajumba et al., 2021). In the report by Feyerabend et al. (2010), lanthanides are cytotoxic against the human osteosarcoma cell line MG63, but not to the human umbilical cord perivascular (HUCPV) cells, with some of them increasing the levels of the inflammatory factors TNF- $\alpha$  and IL-1 $\alpha$ . Lanthanides did not stimulate the growth of HaCaT cells (keratinocytes), while in human dermal fibroblasts they enhanced the proliferation and the production of matrix metalloproteinase-1 (MMP-1) and did not influence the production of type I procollagen (Jenkins et al., 2011). The “Raman fingerprints” of human sperm DNA exposed to CeCl<sub>3</sub> demonstrated that Ce can cause DNA damage through oxidative damage and destruction of the antioxidant defense systems of sperm and even lead to damage or death (Chen et al., 2015). This implies that excess REEs exposure may be a risk factor for human infertility. A cytotoxicity study of CeO<sub>2</sub> NPs in six cancers and two normal cell lines was elaborated by Pešić et al. (2015). Only two cancer cell lines were sensitive to CeO<sub>2</sub> NPs, melanoma 518A2 and colorectal adenocarcinoma HT-29, showing a median inhibitory concentration (IC<sub>50</sub>) of 125  $\mu$ M and 183  $\mu$ M, respectively. In this study, human cells also resulted in lower ROS production and higher antioxidant capacity, implying that normal cells are more resistant to CeO<sub>2</sub> NPs versus cancer cells. Ultrafine particles of Y<sub>2</sub>O<sub>3</sub> were not cytotoxic against human aortic endothelial cells and resulted in an inflammatory response by increasing the concentration of IL-8, ICAM-1, and MCP-1 (Gojova et al., 2007), but CeO<sub>2</sub> ultrafine particles failed to cause any inflammation on the same cell lines (Gojova et al., 2009). Lanthanum and Ce, alone or in combination, inhibited cell proliferation and altered genes involved in oxidative stress pathways in HepG2 and HT-29 cell lines (Benedetto et al., 2018). Particles enriched with REEs, collected from an REE mining area, showed no cytotoxicity to lung epithelial A549 cells; however, they increased ROS production (Tian et al., 2020). Erbium-laser photoablation was found to cure oligoasthenozoospermia and enhance fertilization (Antinori et al., 1994). Europium selenide nanoparticles synthesized *in*

*vivo* by recombinant *Escherichia coli* cells were cytotoxic to the cancer cell lines 293T, HeLa and SKOV-3, and thus could be promising drug carrying agents against cancers (Kim et al., 2016). An Eu complex bearing 8-hydroxyquinoline-N-oxide and 1,10-phenanthroline ligands exhibits low toxicity to normal HL-7702 cells, but high antiproliferative activity against cisplatin-resistant A549/DDP cells by upregulating the expression of LC3 and Beclin1 and downregulating p62 to induce apoptosis which is related to its cell autophagy-inducing properties (Yang et al., 2021). Neodymium-iron-boron magnets are cytotoxic against human oral mucosal fibroblasts (Donohue et al., 1995). Although not causing cell death, the powder of Nd<sub>2</sub>Fe<sub>14</sub>B magnet can cause a dose-dependent ROS production to A549 cells (Rumbo et al., 2021). Gadolinium oxide nanoparticles were found cytotoxic to human umbilical vein endothelial cells, inducing lipid peroxidation, ROS production, mitochondrial dysfunction and autophagic modulation through apoptosis and necrosis (Akhtar et al., 2020). Nanoparticles of Y<sub>2</sub>O<sub>3</sub> induced apoptosis and necrosis to human embryonic kidney (HEK293) cells, by elevating the cellular ROS levels, increasing the mitochondrial membrane permeability, and causing DNA damage (Selvaraj et al., 2014). Yttrium aluminum borates nanoparticles are cytotoxic to human bronchial epithelial cells (Cieslik et al., 2019). The study by Perry et al. (2020) was the first to provide *in vitro* data on the efficacy of radiation therapy with Tm nanoparticles for the management of metastatic cutaneous squamous cell carcinoma (cSCC). REE salts are not skin sensitizers and do not exhibit endocrine disruption to epidermal keratinocytes (Rucki et al., 2021). The chlorides of Ce, La, Eu and Yb are cytotoxic against the embryonal kidney HEK-293 cells (Heller et al., 2019). La citrate induces anoikis (a type of programmed cell death) in HeLa cells by reorganizing actin cytoskeleton and increasing the co-localization of F-actin with mitochondria (Su et al., 2009). La citrate is also cytotoxic against SiHa cells causing a mitochondrial dysfunction that induces apoptosis (Shen et al., 2010). EuCl<sub>2</sub> and EuCl<sub>3</sub> reduced the viability of the human monocyte (U937) and primary human mononuclear cells (PBMNCs) (Bladen et al., 2013). In the study of Wang L. et al. (2020) a REE-doped up conversion nanoparticle (UCNP), sodium yttrium fluoride NP, doped with Yb and Er (NaYF<sub>4</sub>: Yb<sup>3+</sup>, Er<sup>3+</sup>), induced cytotoxicity to HepG2 cells after their internalization by promoting ROS generation, and the induced apoptosis was related to the mitochondria mediated pathway after seeing an increase of cleaved caspase-9, cleaved caspase-3 and Bax and a decrease in the anti-apoptotic protein, Bcl-2. They also compared to the cytotoxicity of the UCNP with that of Y<sup>3+</sup> ions and concluded that the properties of NPs did not play any significant role in the cytotoxicity of the UCNP. In contrast to the above mentioned, GdCl<sub>3</sub> increased the survival and the proliferation of HeLa cells by promoting S phase and activating FAK and JPK, but also attenuated the serum deprivation-induced cell loss (Zhang et al., 2009).

## 10 Conclusion

More than one third of the literature about REE effects on human health regards cytotoxicity. Hair, blood and blood serum are the most biomonitoring matrices. In the near future, research should focus on other potential REE targeted tissues and organs. The effects on LREEs are much more studied than for MREEs or HREEs, thus requiring more attention. Cerium is the most investigated among all REEs.

The REEs mining showed to affect human health as they are accumulated in hair, urine or blood, not only of mining workers, who are exposed directly to these elements, but also of residents near the mining areas. Precautionary regulation should be evaluated for their protection—in the perspective of the “prevention is better than cure” paradigm.

Human daily exposure to REEs can occur also due to other factors such as improper management of e-wastes or high traffic volume since they are used for developing technological devices and as additives in diesel fuels. Although REE levels in the environment and food are currently present at low levels and do not possess a huge threat for human health, they should be considered as a potential future risk since REEs concentration is expected to increase along time.

REEs exposure has been associated with several diseases like NTD, anemia or endomyocardial fibrosis. The lifestyle (smoking and drinking) could contribute to increase the exposure to REEs.

A number of studies have shown that Gd used as a contrast media in MRA and MRI can be accumulated in kidney or muscles causing renal failure and fibromyalgia. Through cytotoxicity studies, GBCAs showed to increase the levels of MMP-1 and TIMP-1, possibly interfering in biological processes where collagen is involved.

A regulation should be implemented also in occupational REE exposures since many jobs may lead to adverse effects. To date, most studies have focused on REE miners and workers involved in metal production. More focus should be given also to other jobs, concerning occupational exposure to REEs, like workers in diesel or technological devices production.

The mechanism of action and the metabolic pathways that REEs activate are still unclear in most cases. GBCAs can activate the pathways related to MMP-1 and TIMP-1. Activation of the PDGF receptor pathway is referred above. REEs can activate the metabolic pathways resulting to ROS production, DNA damage and apoptosis to several cell lines. They seem to be more cytotoxic to cancer cells than normal cells, probably by promoting apoptosis, suggesting their potential protective action against cancer. In some studies, they acted as antioxidants activating pathways related to numerous proteins such as SOD2, GSA, TNF- $\alpha$ , IL-1 $\alpha$ , IL-8, ICAM-1, MCP-1, LC3, Beclin1, p62, caspase-9, caspase-3, BAX, Bcl-2, FAK and JPK. They can be uptaken by several cellular compartments like

mitochondria, lysosomes, cytoplasm and nucleus. Attention should be given to the case of mitochondria, since they have been mentioned to cause mitochondrial disruption. REEs have similar ionic radius to calcium and so they may interfere with its regulation. Future cytotoxic studies should focus on the mechanisms of action of REEs to elucidate the metabolic pathways they can activate.

As a general conclusion, REEs seem to constitute a potential risk for human health. Specific human biomonitoring programs should take place especially for occupational exposures (i.e., miners, e-waste workers), evidencing, particularly, how tobacco smoke can change the background exposure level in the population. Further investigations are necessary to highlight stronger cause-effect relationships between pathological status and REEs exposure, looking for their presence in the right target matrix.

## Author contributions

Conceptualization, AB, GP, GL, MT, MG; Literature gathering, AB, AG; Writing—original draft preparation, AB, GP, AG, MT, GL; Writing—review and editing, AB, GP, AG, MT, GL; Visualization, AB, GP, GL, MT, MG; Supervision, MT, GL, MG; Project administration, MT, GL; Funding acquisition, MT, MG, GL. All authors have read and agreed to the published version of the manuscript.

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## Conflict of interest

The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

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