Research Article

Radiation Resistance of a Structural Material Based on Modified Titanium Hydride


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This work investigates the radiation resistance of a structural material based on modified titanium hydride and a Portland cement in a flux of neutron and γ-radiation. An assessment of the geometric and physicomechanical properties is given, along with the surface structure of irradiated cement composites, and the phase composition of the main hydrosilicates of the hydrated cement matrix during its γ-irradiation. It is shown that the use of a shot of titanium hydride increases the radiation resistance of radiation shielding based on a cement matrix, in comparison with the unmodified shot. A composite based on a modified shot of titanium hydride retains its basic properties after γ-irradiation, at an absorbed dose of up to 10 MGy. At an absorbed dose of 2 MGy in the Portland cement matrix of a composite based on a modified shot of titanium hydride, the formation of suolunite hydrosilicates occurs. It was established using X-ray fluorescence that, in the titanium hydride, a redistribution of the electron density occurs at an absorbed dose of γ radiation of 5 MGy, caused by structural phase changes due to the ongoing dehydrogenation processes.

1. Introduction

When designing a radiation protection structure, it is necessary to use materials that retain their operational properties under the influence of reactor irradiation (physicomechanical, physicochemical, structural, thermophysical, etc.). Radiation exposure, to a large extent, affects the mechanical properties of materials, their strength, and ductility. This influence must be taken into account when building critical structures for radiation protection (nuclear and thermonuclear reactors, spacecraft, etc.) from ionizing radiation. Radiation-induced changes in mechanical properties is the result of a complex of various processes occurring in the structure of the material, including the formation of point defects and their accumulations (dislocation loops and pores); the formation of gaseous impurities (helium, hydrogen), stimulating gas swelling and embrittlement; decomposition of solid solutions; formation and dissolution of second phases; formation of radiation-stimulated diffusion and segregation of components; and radiation creep [1–4].

The ability of materials to retain their basic properties after irradiation (within the specified range) is determined by the concept of radiation resistance. These properties, first of all, include the strength characteristics of the material, changes in its geometric parameters, as well as radiation-protective properties. The quantitatively characteristic is the maximum value of the absorbed radiation dose at which the material changes its properties to a subcritical level. The concept of the subcritical level depends on the specific operating conditions of the radiation-protective material. For example, for materials inside the elements of metal structures, radiation resistance is understood as the absorbed dose of gamma radiation causing the formation of macrocracks in the material. For materials operating in a stressed state, radiation resistance is determined by a decrease in strength characteristics, the specific value of which depends on the function performed by the material.
Many well-known studies have been devoted to the subject of radiation resistance of materials, in particular, structural steels, various metals, and their alloys, as well as the mechanisms of radiation defect formation occurring during irradiation. The problem of radiation-induced degradation of the mechanical properties of materials is of particular importance when considering the almost twofold extension of the service life of nuclear reactors. For this reason, increasing the radiation resistance of structural and functional materials is one of the most important in modern radiation materials science [5–11].

Composite materials based on cement binders (concretes) containing various functional fillers have found the most use as radiation protection materials. This is due to the possibility of obtaining materials of various geometries and filling hollow radiation-shielding metal structures with mobile cement mortar. At the same time, the concrete has high radiation resistance (it does not change the geometric dimensions in the integral neutron fluxes up to $10^{21}$ neutrons/cm$^2$) and has only negative radiation deformations (reduces the volume) [12–15].

When exposed to gamma radiation, the mechanism of cracking in Portland cement concrete is determined by the dislocation of atoms in the cementing phases [16–21], dehydration of cement hydrates [17, 18, 21], and amorphization of aggregates [22–24]. Hydrogen ions formed as a result of water radiolysis are capable of being attracted by hydrated ions of clinker minerals, forming new phases, solid solutions, and intrusions in the structure of cement paste [25]. The impact of gamma rays on chemically bound water may directly impart sufficient energy in order to dehydrate phases such as C–S–H or Ca(OH)$_2$ [26]. Neutron radiation can lead to an increase in the volume of the filler. The change in the properties of concrete shielding during irradiation mainly depends on changes in the properties of aggregates and in the interphase interaction between the aggregate and the cement matrix. Concrete is least susceptible to cracking processes, in which the radiation deformations of the aggregate are close to or less than the radiation deformations of the concrete itself [27].

One of the most effective fillers used for reactor protection is titanium hydride, in the form of spherical granules (shot). Its high hydrogen content (up to 3.6 wt. %) and density (up to 3.8 g/cm$^3$) make it possible to effectively use titanium hydride to simultaneously attenuate both neutron and $\gamma$ radiation [28–32]. Previous reports [33–38] have shown that modification of the surface of titanium hydride shot with borosilicate coatings or ion-plasma spraying of titanium metal significantly increases its thermal stability. The creation of a modifier shell on the surface of a shot of titanium hydride prevents thermal diffusion of hydrogen into the environment and increases the initial temperature of dehydrogenation to 500°C. This expands the possibility of using titanium hydride as a radiation protection material.

However, studies of the radiation resistance of a cement composite based on shot of titanium hydride have not been carried out. To this end, this paper presents the results of studies of the effect of high-energy $\gamma$ and neutron radiation on the structure and properties of a radiation protection material, based on a modified shot of titanium hydride and a Portland cement binder. For comparison, the results of studies for cement stone and a composite based on an unmodified shot of titanium hydride are also presented.

2. Materials and Methods

2.1. Materials. The following materials were used as objects of research:

1. A composite material based on a mixture of a modified shot of titanium hydride, Portland cement, water, and plasticizing additives was examined. The ratio of Portland cement and a shot of titanium hydride in the mixture was 1:3; the water–cement ratio was 0.3. Titanium hydride was modified by depositing a borosilicate shell on its surface [35]. The material was obtained by vibration compaction on the KV-56 stand, with a vibration frequency of 3000 counts/min. After compaction, the material was hardened for 28 d and then heat-treated at 300°C for 500 h.

2. Composite material based on an unmodified shot of titanium hydride, Portland cement, water, and plasticizing additives. The ratio of Portland cement and a shot of titanium hydride in the mixture was 1:3; the water–cement ratio was 0.3. The material was obtained by vibration compaction on the KV-56 stand, with a vibration frequency of 3000 counts/min. After compaction, the material was hardened for 28 d and then heat-treated at 300°C for 500 h.

3. Cement paste based on a mixture of Portland cement and water. Water–cement ratio 0.3.

The material was obtained by vibration compaction on the KV-56 stand, with a vibration frequency of 3000 counts/min. After compaction, the material was hardened for 28 d and then heat-treated at 300°C for 500 h.

2.2. Methods. 1. Irradiation of the composite material in a neutron flux was carried out in the zone of the reflectors of the core of the IVV-2M research nuclear reactor, at a temperature gradient on the composite sample of no more than 1.5°C/mm. The temperature gradient on the composite sample was determined on the basis of the thermal calculation of the reactor irradiation device taking into account the required neutron fluence of $2 \times 10^{19}$ neutrons/cm$^2$. The calculated value of the heat transfer coefficient was taken equal to 8000 W/m$^2$·K. The specific energy release of the composite material was 0.0216 W/(g·MW).

The irradiation device consisted of an aluminum ampoule with ten composite samples, placed in a vessel sealed from the reactor water, which, in turn, was installed in the “wet” channel of the reactor (Figure 1). The geometry of the material samples was a cylinder 22 mm in diameter and 45 mm in height.
The samples were located symmetrically relative to the central plane of the reactor core. Inserts between the samples in the form of steel washers served to equalize the temperature of the samples located below and above the central plane of the core, where the gamma field is weaker. The body of the irradiation device was sealed from water circulating in the gap between the walls of the body and the “wet” channel.

The samples were located symmetrically, relative to the central plane of the reactor core. The arrangement of the samples ensured their irradiation in the temperature range 245–255°C, with an average flux density of fast neutrons ($E > 0.5$ MeV) for 15 MW power, the reactor power $5.55 \times 10^{12}$ neutrons/(cm$^2$·s), and a fluence of $2 \times 10^{19}$ neutrons/cm$^2$. The temperature in the material sample was determined by the calculation method based on the temperature of the coolant in the cooling zone of the body of the irradiation device and the heat transfer coefficient. The calculation of the temperature fields in the samples of the material of the assembly of the irradiation device was carried out using the calculation program "ANSYS" [39]. The calculation results are presented in Figures 2 and 3.

For research, 20 samples of composites were used, obtained under the same conditions from one batch of a composite mixture. The conditions of the reactor tests made it possible to irradiate one batch of samples (10 samples) simultaneously loaded into the irradiation device. Investigations of changes in the properties of the composite samples were carried out after the collection of the average fast neutron fluence ($E > 0.5$ MeV) over the samples, equal to $1.9 \times 10^{16}$ neutrons/cm$^2$. All work with irradiated samples was carried out remotely in hot chambers. The accuracy of determining the geometric dimensions and mass was the same as for nonirradiated samples.

2. Irradiation of the composite material in a flow of high-energy $\gamma$ radiation was carried out at various dose loads in the barrier geometry of the shield, on a $^{60}$Co ($E = 1.3$ MeV) research $\gamma$ installation with a dose rate of 21 kGy/h. For the experiment, we used 25 samples based on an unmodified fraction of titanium hydride (does not contain a borosilicate shell) and 25 samples based on a modified fraction of titanium hydride (contains a borosilicate shell). In each experiment, 5 samples of composite material $3 \times 3 \times 3$ cm in size, mounted parallel to the plane of a wide beam of gamma radiation, were simultaneously exposed to irradiation until they reached the appropriate dose of radiation. In each experiment, all five samples received the same radiation dose. Five samples from each batch were evaluated as controls. The geometry of the experiment is shown in Figure 4. The control of integral doses of irradiation of the samples was performed using cellulose triacetate dosimetric films. The temperature of the samples was measured using a thermocouple.

<table>
<thead>
<tr>
<th>Content [wt. %]</th>
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<tbody>
<tr>
<td>TiH$_{1.8}$</td>
</tr>
<tr>
<td>77.71</td>
</tr>
</tbody>
</table>

Table 1: Chemical composition of a composite material based on a modified shot of titanium hydride and Portland cement after heat treatment at 300°C for 500 h.
3. Results and Discussion

External examination of the composite samples after irradiation showed that their integrity, color, and surface structure were all retained, in comparison with their initial state (Figure 5).

The results of studies of changes in the linear dimensions, strength, and density of composite specimens based on modified titanium hydride and Portland cement, after fast neutron irradiation at a fluence of $1.9 \times 10^{19}$ neutrons/cm$^2$, are presented in Table 2.

It follows from Table 2 that the geometry of the samples after irradiation did not change or changed only slightly. The maximum change in the height of the samples ($\Delta h$) is 0.05% of the initial value. The maximum change in the average diameter ($d$) is 0.09% of the initial value. All samples under investigation had reduced volume, associated with the removal of water bound in the hydrosilicate cement matrix. The average density of the samples prior to irradiation was 3.27 g/cm$^3$ and 3.28 g/cm$^3$ after irradiation. Changes in density (upward) of most of these samples are within the volume determination error. Thus, this work did not reveal obvious changes in the geometric and physicomechanical properties of composites based on modified titanium hydride and Portland cement after neutron irradiation. There does not appear to be a statistical significance.

According to SEM data (Figures 6–9), at an irradiation dose of more than 2 MGy, changes in the surface structure of the unmodified titanium hydride shot in the bulk of the cement composite and stratification (an increase in the interphase boundary) between the shot and the hydrated cement matrix occur due to the ongoing processes of dehydrogenation of the shot during radiogenic heating.

The released hydrogen creates channel microcracks and leads to the destruction of the composite structure. In this case, in a composite based on a modified shot of titanium hydride, no noticeable interphase disturbances occur up to an absorbed dose of 10 MGy. There is a visual anchoring of the filler surface with the cement matrix (Figures 6(b)–9(b)), which occurs due to significant adhesion of the cement matrix to the borosilicate modification shell of the titanium hydride shot.

Figure 10 shows data regarding changes in the strength properties of materials based on unmodified (UTHS) and modified (MTHS) shot of titanium hydride, under $\gamma$-irradiation with different dose loads.

The most significant decrease in strength occurs in a composite based on an unmodified fraction of titanium hydride at an absorbed dose of gamma irradiation of more than 2 MGy (temperature 350°C), which is associated with both thermal diffusion of hydrogen and dehydration of calcium hydrosilicates as a result of concomitant radiation heating.

Figures 11–14 show diffraction patterns of cement paste without a modified shot of titanium hydride and cement paste in a composite based on a modified shot of titanium hydride after $\gamma$-irradiation with a dose load from 1.0 to 10.0 MGy.

3. The microstructure of the surface of the composite material was studied by scanning electron microscopy (SEM) in the diffraction mode of reflected (backscattered) electrons. In this mode, the contrast largely depends on the crystallographic orientation of the grains relative to the sample surface and on the chemical composition of the phases. In this case, the microstructure of the material surface can be studied without preliminary etching of the section. The studies were carried out on a Quanta 600FEG SEM (FEI Company, USA), at an accelerating voltage of 20 kV and an electron beam current of 36 nA.

4. The study of the phase composition of the main hydrosilicates of the hydrated cement matrix during its irradiation and the effect of shot of titanium hydride on the change in the structure of the cement paste in the composite was carried out using X-ray phase analysis (XPA). X-ray diffraction (XRD) was performed using CuKa radiation with a Ni filter, on an ARL XTRA X-ray diffractometer (Switzerland) using an ASTM file (USA). The spectra were recorded with an MSTR-4 ionization counter at angles from 14 to 56°. The diffraction patterns were measured with a RK-3A comparator. The XRD patterns of the sample being studied were processed separately for each shooting interval, using the PDWin program. Determination of the phase composition was carried out by analyzing the interplanar distance $d$ and the intensity of spectral reference lines.

5. X-ray fluorescence (XRF) analysis was performed on an ARL9900 Work Station XRF spectrometer (Switzerland) with a built-in diffraction system, a high-performance tube with an Rh anode, and a 50 µm (75 µm) Be-window. The analysis software used was GuantAS, UniGuant, and OptiGuant. The XRF method is based on the excitation of the characteristic radiation of different elements in a sample using an external source (X-ray or isotope photon source), and registration of said characteristic radiation in the form of a line spectrum [40]. The position of the peaks in the $K$, $L$, and $M$–series of characteristic radiation on the energy scale is responsible for the identification of chemical elements in the sample, and the intensity of the peaks broadly represents the concentrations of elements.
Analysis of the data showed that modified titanium hydride introduces some changes in the structure of the hydrosilicate phases of the cement paste, at absorbed γ-radiation doses of 1.0–2.0 MGy.

At a dose of 2 MGy, the presence of a modified shot of titanium hydride contributes to the formation of low-basicticy calcium hydrosilicates in the system, in the form of bar-like crystalline rods Ca₂Si₂O₆(OH)₂•H₂O (Figure 15). According to known data [41], this can lead to compaction of cement paste and a decrease in its water and gas permeability.

Analysis of diffraction patterns of irradiated composites at a dose of 5–10 MGy showed the formation of silicate phases of the same chemical composition and crystallographic parameters in the cement paste and cement paste in the composition of the composite based on the modified shot of titanium hydride.

The electronic structure of titanium atoms in the composition of the composite was estimated using XRF, for various dose loads of γ radiation. Titanium is characterized by electronic transitions of the M, L — K type, and by the appearance of $K_α$ and $K_β$ lines in the XRF spectrum (Figure 16). The parameters of XRF spectra of titanium atoms in a composite subjected to γ-irradiation, with an absorbed dose of 0.5–10.0 MGy, are presented in Table 3.
With an absorbed dose of γ radiation from 0.5 to 2.0 MGy, the intensity of the $K_{\alpha}$ and $K_{\beta}$ lines of the XRF spectrum decreases by an average of 5.2% (Figure 17).

A significant increase in fluorescence in the titanium atom at 5.0 MGy (by 31% for $K_{\alpha}$ lines and 51% for $K_{\beta}$ lines) indicates the intensity of the electron-photon interaction. The change in fluorescence intensity is also associated with the selective excitation of titanium atoms under γ- and X-ray-irradiation, which, in turn, may be due to structural and phase changes in titanium hydride, due to

**Figure 7:** Surface microstructure (SEM) of a fracture of a cement composite chip based on unmodified (a) and modified (b) shot of titanium hydride subjected to γ-irradiation, at a dose $D = 2$ MGy.

**Figure 8:** Surface microstructure (SEM) of a fracture of a cement composite chip based on unmodified (a) and modified (b) shot of titanium hydride subjected to γ-irradiation, at a dose $D = 5$ MGy.
the diffusion of hydrogen from the bulk of the shot into the surface layers. In this case, a shift of the $K_\alpha$ and $K_\beta$ lines to the high-energy region of the spectrum ($E \approx 4.49 \text{ eV}$) is observed.

A continuous change in the intensity ratio of the $K_\alpha/K_\beta$ excitation lines over the entire range of dose loads indicates a redistribution of the electron density in titanium hydride, under the action of $\gamma$ learning.

Figure 9: Surface microstructure (SEM) of a fracture of a cement composite chip based on unmodified (a) and modified (b) shot of titanium hydride subjected to $\gamma$-irradiation, at a dose $D = 10 \text{ MGy}$.

Figure 10: Strength of titanium hydride composites subjected to $\gamma$-irradiation.
Figure 11: Diffraction patterns of cement paste (A) and cement paste in a composite based on a modified shot of titanium hydride (B), subjected to γ-irradiation dose $D = 1$ MGy: $1$-2CaO·SiO$_2$·H$_2$O; $2$-Ca(OH)$_2$.

Figure 12: Diffraction patterns of hardened Portland cement paste (A) and cement paste in a composite based on a modified shot of titanium hydride (B), subjected to γ-irradiation dose $D = 2$ MGy: $1$-2Ca$_3$Si$_2$O$_7$·H$_2$O; $2$-Ca$_5$SiO$_4$; $3$-Ca(OH)$_2$; $4$-Ca$_3$Si$_2$O$_5$(OH)$_2$·H$_2$O. 
Figure 13: Diffraction patterns of cement paste (A) and cement paste in a composite based on a modified shot of titanium hydride (B), subjected to γ-irradiation dose \( D = 5 \) MGy: 1-Ca_2SiO_4·0.3O_2; 2-Ca_2SiO_4; 3-Ca(OH)_2.

Figure 14: Diffraction patterns of cement paste (A) and cement paste in a composite based on a modified shot of titanium hydride (B), subjected to γ-irradiation dose \( D = 10 \) MGy: 1-Ca_2SiO_4; 2-Ca(OH)_2.
Figure 15: Microstructure of the cleavage surface of a composite based on a modified shot of titanium hydride, subjected to γ-irradiation at a dose of $D = 2$ MGy (SEM): (a) 2500x magnification; (b) 25000x magnification.

Figure 16: XRF spectrum of titanium atoms, in a composite heat-treated at 300°C.

Table 3: Parameters of XRF spectra of titanium atoms, in a composite subjected to γ-irradiation.

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<tr>
<th>Dose $D$ [MGy]</th>
<th>$\lambda$ (mÅ)</th>
<th>$I$ (imp./c)</th>
<th>$\Delta_{1/2}$ (mÅ)</th>
<th>$I_{\alpha}/I_{\beta}$</th>
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4. Conclusions

A composite material based on a modified shot of titanium hydride and Portland cement, hydrated under hydrothermal conditions, has a high radiation resistance when irradiated with fast neutrons. In a flow of fast neutrons with energy >0.5 MeV, at a fluence of $1.9 \times 10^{19}$ neutrons/cm$^2$, the material retains its integrity, color, and surface structure in comparison with the initial state. Evaluation of the geometric and physicomechanical properties of the irradiated composite samples did not reveal any obvious changes after irradiation with fast neutrons.

The use of a modified shot of titanium hydride increases the radiation resistance of the cement composite in a high-energy $\gamma$-ray stream, compared to an unmodified shot. The homogeneity of the composite structure is maintained, up to an absorbed dose of $\gamma$ radiation of 10 MGy. The presence of a modified shot of titanium hydride in the cement composite under $\gamma$-irradiation dose loads of 1–2 MGy leads to the formation of low-basicity hydrosilicates, which does not significantly affect the geometric nor strength properties of the cement paste.

Evaluation of the electronic structure of titanium atoms in the cement composite by the XRF method showed that, under the action of $\gamma$-irradiation, a redistribution of electron density occurs in the hydride phase of the titanium hydride. This is caused by structural and phase changes in the modified titanium hydride, due to the ongoing dehydrogenation processes at increased radiation loads.

Data Availability

The data are available on request to R. N. Yastrebinsky (astrebinskiy.rn@bstu.ru).

Conflicts of Interest

The authors declare that there are no conflicts of interest regarding the publication of this paper.

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