1. Introduction

High-manganese steels are the most widely used material for the manufacture of parts and the application of coatings for such operational conditions when there are impact-abrasive wear and destruction under the influence of dynamic and cyclical loads. The main structural component of such steels is manganese austenite, capable of intensive deformation...
strengthening. Important properties that determine the performance of manganese austenite is its crystalline and electron structure. Its features determine the level of the structure stability, the value of elastic constants, as well as the energy condition of the crystal, and, consequently, the stacking fault energy (SFE) [1]. A value of SFE is a defining characteristic when establishing the fundamental possibility for the deformation strengthening of manganese steels by the twinning-induced plasticity (TWIP) [2] or the transformation induced plasticity (TRIP) [3] techniques. In this case, it is especially important to assess the influence of dissolved alloying elements in manganese austenite on its stability and the physical-mechanical characteristics. In addition, when applying coatings from manganese steels, it is important to assess the effect of alloying on the thermophysical characteristics, which emerge in different procedures for calculating temperature fields [4, 5]. Among these elements, it is important to assess the effects of silicon, which is part of almost all high manganese steels and coatings based on them. Chromium is introduced into such coatings to improve corrosion resistance [6], which is evaluated by electrochemical methods [7, 8].

One common procedure in materials science, which makes it possible to model the crystalline structure and determine its properties in a given time, is the quantum-mechanical calculations within the framework of the density-functional theory (DFT) using a method of nonlocal pseudopotential [9]. Such calculations typically employ software tools such as Vienna Ab initio Simulation Package (VASP) (Austria) [10], Cambridge Serial Total Energy Package (CASTEP) (UK) [11], and others. The main limitations of this procedure are the complexity of modeling unordered solid solutions, which include austenite. This necessitates modeling them in the form of supercells of orderly solid solutions that simulate individual cases of unordered structure. In addition, such calculations are carried out for defect-free structures at 0 K, so only the theoretical values of mechanical characteristics are reflected. In this case, an analysis of real solid solutions produces, basically, a qualitative pattern of the influence of dissolved components.

Studying such materials implies the design of new wear-resistant coatings with composite structures, which is relevant for materials science and tribology.

2. Literature review and problem statement

The construction of new, as well as the continuous improvement of existing algorithms of quantum-mechanical calculations of the crystalline structure and properties allow them to be used for modeling the most important phases in steels, in particular austenite of different compositions [12, 13]. In order to study the impact of Si and Al additives on the elastic characteristics, structure, and energy state of nickel-chromium austenite, the authors of work [14] modeled its crystalline structure in accordance with the Fe₂Cr₃Ni₃ formula. To simulate an unordered solution, the atoms of dissolved components were symmetrically arranged in the FCC (face-centered cubic) nodes of the supercell: 2×2×1, based on γ-Fe, which contained 16 atoms. According to the calculation results, it was found that the dissolution of Si in the austenite of the specified composition, additionally alloyed with Al, leads to a decrease in the elastic constants and theoretical hardness and an increase in the plasticity and Poisson coefficient. At the same time, the thermodynamic and electrochemical stability (calculated by the number of electrons at the Fermi level) increases significantly. However, the cited work disregarded the issue of modeling the introduction solutions involving elements such as carbon and nitrogen. Determining the effect of dissolved nitrogen on the crystalline structure and elasticity characteristics of manganese-chromium austenite of the Fe₈Cr₃Mn₈ formula by modeling with the use of DFT is reported in paper [15]. Austenite with different carbon content was modeled in study [16]; it was represented in the form of 5×5×5 supercells based on an FCC cell. They contained 125 Fe atoms in the nodes of the crystalline lattice and 1, 7, and 9 C atoms located in the octahedral pores. The results of the reported modeling showed a nonlinear growth of elasticity and shear modules while increasing the amount of dissolved C, which clearly manifests itself with an increase in the amount of dissolved carbon over 5 %. The structure was modeled in the form of a γ-Fe-based superlattice, which contained 32 metal atoms in the nodes of the lattice and 1 nitrogen atom in the octahedral pore located in the center of the lattice. According to the calculation results, the parameter of the lattice of the Fe₈Cr₃Mn₈ solid solution at nitrogen dissolution increases from 3,507 to 3,522 angstrom (Å), and its formation energy increases by 2–2.5 times. At the same time, the values of modules B and G increase while the Poisson coefficient decreases. However, all the structures studied with different mutual arrangements of Fe and Cr atoms, according to the B/G criterion [17], refer to plastic materials: B/G=2.23–3.25 (higher than 1.75). Determination of the yield strength of high-manganese steels alloyed with Al on the basis of a model that included the DFT calculations of the contribution of short-range ordering energy was carried out in [18]. In contrast to works [15, 16], instead of superlattices, the simulation was performed on the basis of calculating the energy of clusters with an FCC lattice and the octahedral pore filled with carbon in the center. The number and configuration of the atom arrangement resulted from searching for crystalline structures with the lowest ordering energy. As a result, six stable structures of the following formulas were identified: Fe₃Al₈C, Fe₅MnAl₈C, Fe₆Mn₂Al₈C, Fe₇Al₃C, Mn₆Al₈C, and Fe₃Mn₃Al₈C. Extrapolating the data about these structures on the composition of standard austenite steels has made it possible to determine the strength limit with a margin of error of ~10 %. The Ni and Co influence on the electron structure and the elastic characteristics of austenite of the Fe₃C formula was studied in work [19]. The simulations were carried out using a DFT procedure employing the CASTEP software for structures with an FCC lattice, whose Fe atoms were replaced with Co and Ni atoms in equal proportions in an amount of ~20 at. %. The resulting equilibrium structures were characterized by the presence of a tetragonal FCC lattice with modified parameters (a=3.7367–3.7451 and c=3.7843–3.8061) compared to austenite of the Fe₃C composition (a=3.7478 Å). All model structures met the criteria for mechanical stability and were plastic materials according to the B/G criterion. At the same time, in a general case, there is a simultaneous dissolution of Co and Ni in austenite of the Fe₃C formula. An analysis of values of the calculated elastic constants for structures with a different arrangement of atoms reveals that in a general case the simultaneous dissolution of Ni and Co leads to the higher values of G and a decrease in the B value and plasticity. The comprehensive approach to determining the elastic characteristics of high-manganese austenite was
reported in work [20], whose authors, in parallel with DFT calculations, performed experimental research by a nanoindentation method [21]. The samples for their experimental study were the coatings applied by magnetron spraying and alloys from pure components containing C in the amount of 0.5–3.0 at. %, and Mn, from 28–30 at. %. The comparison of the calculated and experimental values of the parameters of austenite lattices of different compositions shows that the values of the calculated magnitudes (3.51–3.54 Å) are lower compared to the calculated ones (3.59–3.61). However, the tendency to increase the lattice parameter when increasing the contents of Mn and C is reflected in both calculations and experimental results. The experimental values of the Young modulus are described by theoretical calculations with a ~15% accuracy and are within 185–220 GPa. At the same time, increasing the carbon content within 0–3 at. % does not lead to a noticeable increase in the elasticity module.

Our generalization of the above data from the scientific literature shows that for the alloyed austenite with the total content of alloying elements dissolved by replacing ≤50 at. % and in the amount of C≤3 at. %, the lattice parameters are within ~3.45–3.6 Å. The calculated values of the E, B, and G elasticity modules are within 185–346, 150–280, and 70–150 GPa, respectively. The data that most accurately describe the results from experiments were acquired by using the generalized gradient approximation (GGA) into the Perdew-Burke-Ernzerhof (PBE) images using the CASTEP and VASP programming codes. This allows us to argue that it is expedient to simulate alloyed manganese austenite in the form of an FCC superlattice by their central pore filled with a carbon atom.

3. The aim and objectives of the study

The aim of this study was to establish the impact of Si and Cr additives on the crystalline structure and properties of manganese austenite by experimentally determining the structure parameters and by ab initio simulation based on DFT methods.

To achieve the set aim, the following tasks have been solved:

– to determine the phase composition of high-manganese steel and to analyze its Mossbauer spectrum;
– to assess the impact of Si and Cr on the elastic constants, mechanical characteristics, and stability of manganese austenite;
– to establish patterns in the electron structure of manganese austenite with the dissolved atoms of Si and Cr and to establish its relation to the physical-mechanical characteristics.

4. Materials and methodology to study the influence of Si and Cr on the characteristics of manganese austenite

We theoretically determined the parameters of the crystalline structure and elastic constants of manganese austenite depending on its composition within the framework of DFT and pseudopotential methods using the basis of flat waves. The calculations were performed using the CASTEP programming code by the generalized gradient approximation (GGA) method in the representation by Perdew-Burke-Ernzerhof (PBE) [22]. The Brillouin’s zone was split into a grid of 4×4×4 k-points according to the Monkhorst-Pack scheme [23]. We described the potentials of ions by using the ultra-low potentials by Vanderbilt [24] for the electron configurations of ions: C–2s22p2, Si–3s23p2, Mn–3d54s2, Fe–3d64s2. The optimization of the geometry (relaxation of the structure) was considered complete at a volumetric tension of <0.05 GPa and the forces acting on an atom of <0.03 eV/Å. The 2×2×2 supercells based on an FCC lattice with 32 iron atoms and alloying elements per one cell were used to simulate the structure of austenite (Fig. 1). A carbon atom was placed in the central octahedral pore.

In the model superlattice, the following groups of symmetrically arranged atoms can be identified: along the edges (I) and faces (II), as well as at the cube vertices (III), and the octahedron vertices (IV) around the central octahedral pore. Thus, the composition of austenite corresponded to the formula A6B4C2D2F – C, where A, B, D, F are the alloying elements that form the substitutional solutions with iron in positions I, II, III, IV.

Samples for our experimental study were manufactured by the electric arc fusion of ferrosilicon-manganese and low-carbon steel. Based on the results from an X-ray fluorescence analysis, the composition of the alloys corresponded to high-manganese steel containing, % by weight: Mn ~19, Si ~4, C ~1.5, the rest - Fe.

We experimentally studied the parameters of the crystalline structure by a Mossbauer spectroscopy method using the nuclear gamma-resonance spectrometer YaGRS-4M. The Fe57 electron Mossbauer spectra were acquired by using Co57 in the chromium matrix as a source of γ-quantums and by calibrating the isometric offsets relative to sodium nitroprusside.

Thermodynamic calculations were carried out using the CALPHAD (CALculation of PHAse Diagrams) procedure employing the OpenCalphad programming code [25] and the thermodynamic functions given in works [26–28].

The phase composition of the samples was determined by X-ray phase analysis (XRD) using the DRON-3M diffractometer in the filtered CuKα radiation with the subsequent diffractionogram processing based on the Rietveld method.

5. The results of studying the crystalline structure and certain mechanical characteristics of the manganese austenite alloyed with Si and Cr

5.1. The results from the X-ray phase analysis, microstructure study, and Mossbauer spectroscopy of high-manganese steel

The results from the X-ray phase analysis of experimental alloys (Fig 2, a) showed that their base phase is Fe with a face-centered lattice (spatial group Fm-3m). The Fe lattice
parameter was $a=3.576\,\text{Å}$, which is close to the parameters of the austenite lattice in Hadfield's steel after casting [29] and in welded joints of $3.60–3.62\,\text{Å}$ [30]. The results of studying the microstructure of the experimental alloys obtained during the alloying with Si (Fig. 2, $b$) and Cr (Fig. 2, $c$) show that in both cases the structure of the alloys consists of uneven grains of austenite and eutectics with different morphology. In this case, alloying with Cr leads to the formation of a rougher structure of the austenite-carbide eutectics compared to alloying with Si. This is probably due to the different nature of the carbide phases that make up the eutectics. For alloys, alloyed with Cr, it is thermodynamically beneficial to form carbides of the type (Fe, Mn, Cr)\(_7\)C\(_3\), while when alloying with Si, there is only the formation of cementite-type carbides (Fe, Mn)\(_3\)C.

The experimental Mossbauer spectrum of Si, Cr-alloyed manganese austenite is formed by a combination of double and singlet components (Fig. 3). No manifestations of the Seeman nuclear effect are observed, indicating that there are no \(^{57}\text{Fe}\) nuclei in a magnetically-ordered state. It could be argued that the \(\alpha\)-Fe phase is absent and all Fe atoms are part of the austenite structure. Using the UNIVEM-MS software, subject to the Lorentz form of resonance lines, the experimental spectrum was represented as a superposition of two double and one singlet component. They correspond to the resonant absorption of \(^{57}\text{Fe}\) by the nuclei, which are in the crystalline- non-equivalent positions of the lattice. The estimation values of the characteristic parameters of the spectrum components (isomeric shift, quadrupole cleavage) of Lorentzian line shape agree well with the data reported in work [31] when analyzing the spectra of\(^{57}\text{Fe}\). In this case, doublet 1 corresponds to the crystallographic position of \(^{57}\text{Fe}\) number I, doublet 2 – positions number II and IV, singlet 1 – position number III (Fig. 1). Taking into consideration the value of the relative proportion \(p\) of Fe atoms in a particular environment (Table 1), the most likely is the localization of Si, Mn, and Cr alloying elements in the austenite superlattice in positions number II, III, and IV.

Based on this conclusion, in the further calculations of the model crystalline structure, the Mn atoms were arranged in the crystallographic position of phase III, and the Si or Cr atoms – in position IV. The initial parameter of the superlattice was set to $2\times a$, where $a$ is the parameter of the austenite lattice, defined experimentally (3.576 Å). This arrangement of atoms ensured that all the necessary pairs of adjacent atoms were available: C-Mn, C-(Si,Cr), C-Fe, Mn-(Si,Cr), Mn-Fe, Fe-(Si,Cr).

| Phase            | Doublet 1 | Doublet 2 | Singlet 1 |
|------------------|-----------|-----------|-----------|
|                  | $Q_o$, mm/s | $G_o$, mm/s | $p$ | $Q_o$, mm/s | $G_o$, mm/s | $p$ | $Q_o$, mm/s | $p$ |
| $\gamma$-Fe(Mn,C) | 0.65      | 0.31      | 38.7     | 0.16  | 0.21      | 50.0   | 0.18      | 11.3  |
| $\gamma$-Fe$_2$C [31] | 0.65      | 0.30      | 57.0     | 0.10  | 0.24      | 34.0   | 0.23      | 9.0   |

![Fig. 2. The analysis of phase composition and microstructure of experimental materials: $a$ – diffractogram of high manganese experimental steel; $b$ – microstructure of high-manganese steel alloyed with silicon; $c$ – microstructure of high-manganese steel alloyed with chrome](image_url)

![Fig. 3. Mossbauer spectrum of alloyed high manganese steel](image_url)
The comparative analysis of the parameters of the Mössbauer spectra of carbon and manganese austenite reveals the proximity of the parameters of quadrupole uncoupling and line width, which indicates the stability of the austenite phase in experimental alloys.

5. 2. The results of determining the stability and mechanical characteristics of manganese austenite when dissolving Si and Cr

One of the attributes of stability of a crystalline structure is, in particular, its mechanical stability, which, depending on the type of crystal lattice, is determined by the Bourne-Huang criterion. For a tetragonal lattice, which was the basis for model structures, the stability, depending on the elastic constants \((C_{ij})\), is determined by meeting a series of criteria: 
- \(C_{11} > 0, C_{33} > 0, C_{44} > 0, C_{66} > 0, C_{11}C_{12} > 0, C_{11} + C_{33} > 0\), and
- \(2(C_{11} + C_{12}) + C_{33} + 4C_{44} > 0\). The elastic constants were calculated after optimizing the geometry (structure relaxation) in accordance with the relationship between deformation and stresses under the generalized Hook law:

\[
\sigma_{ij} = C_{ijkl} \varepsilon_{kl}.
\]

where \(\sigma_{ij}\) is the stretching strain; \(\varepsilon_{ij}\) is the longitudinal deformation.

Comparing the results of our calculations of values for the elastic constants, the lattice parameter \((a)\), and the full energy of the crystalline lattice \((E_f)\) for pure \(\gamma\)-Fe, according to the selected methodology, to the data reported in [32] shows that they are in good agreement (Table 2).

| Characteristics of the crystalline structure of \(\gamma\)-Fe |
|---------------------------------|
| Elastic constants               | \(C_{11}\) | \(C_{12}\) | \(C_{44}\) | \(a, \text{Å}\) | \(E_f, \text{eV/atom}\) |
| Calculation                     | 445.591   | 294.884   | 254.627   | 3.45342       | −27,685.6               |
| Data from [32]                  | 484       | 234       | 287       | 3.45          | −34,634.5               |

The elastic constants and the parameters of the model structures’ lattices are given in Table 3. An analysis shows that the value of the austenite \((\text{Fe-C})\) lattice parameter \((3.5743 \text{ Å})\), calculated by extrapolating the empirical equation proposed in work [33] to \(T=0\) K, and that of the model structure are within the typical error for calculations based on a DFT method (10%).

The results of our calculations of values for the elastic constants are the initial ones to calculate some mechanical characteristics of a polycrystalline material [34]. According to the average Voigt-Reuss-Hill (VRH) scheme [35], the values of the volumetric elasticity module \((B)\) and the shear module \((G)\) are determined from the following equations:

\[
B = \frac{1}{9} \left( C_{11} + C_{22} + C_{33} \right) + \frac{2}{9} \left( C_{12} + C_{13} + C_{23} \right),
\]

\[
G = \frac{1}{15} \left( C_{11} + C_{22} + C_{33} - C_{12} - C_{13} - C_{23} \right) + \frac{4}{5} \left( C_{44} + C_{55} + C_{66} \right).
\]

Given the known values of \(B\) and \(G\) for polycrystalline structures, the Young module \((E)\) and the Poisson coefficient \((\nu)\) can be calculated from the following equations:

\[
E = \frac{9GB}{G + 3B},
\]

\[
\nu = \frac{3B - 2G}{2(G + 3B)}.
\]

The results of our calculations (Fig. 4) show that the dissolution of \(C\) and the studied alloying elements (\(\text{Mn, Si, Cr}\)) leads to a slight decrease in the values of modules \(E\) and \(G\) and the growth of module \(B\). In this case, there is an increase in plasticity, hardness, thermodynamic stability. Dissolution of Si and Cr in manganese austenite leads to its increased thermodynamic stability, hardness, and plasticity, while the values of elasticity modules \((B, G, E)\) decrease.
The effect of silicon on the change in these characteristics manifests itself more clearly. However, dissolution of silicon leads to a decrease in the anisotropy of the austenite phase ($A_z$) while, when dissolving Cr, anisotropy increases. Structures (Fig. 5), calculated according to the CALPHAD methodology, shows the same tendencies to stabilize the austenite phase as the DFT calculations. In particular, the dissolution of C leads to a slight decrease in $G_m$ in the temperature stability region $\gamma$-Fe (~1,184–1,665 K), according to the equilibrium Fe state diagram. Further dissolution of Mn leads to a significant decrease in the $G_m$ of the austenite phase, which is observed throughout the entire temperature range (800–1,800 K). With the simultaneous dissolution of Mn and Si, the intensity of reducing the free energy of austenite increases by almost two times, compared to Fe-C alloys, where only Mn dissolved. Dissolution of Cr in manganese austenite also leads to a decrease in its Gibbs energy but, compared to Si, its effect is much weaker.

The DFT calculation data at 0 K agree well with the CALPHAD thermodynamic calculations at 800–1,600 K. This indicates that the identified trends in the impact of Si and Cr on elastic constants would be maintained under normal conditions and elevated temperatures.

5.3. The results of analyzing the electron structure of manganese austenite when dissolving Si and Cr

The calculated density of electron states (DOS) of austenite structures (Fig. 6) indicates the similarity of their electron structure, and the lack of energy cracks near the Fermi level (zero energy point) is a sign of the bond of metallic type. According to [14], the number of electrons at the Fermi level determines the level of electrochemical stability of a structure, namely, when it increases, the level of stability decreases. According to this, the model austenite structures in the order of growth of their electrochemical stability can be placed in a series: $\text{Fe}_2\text{C} \rightarrow \text{Fe}_3\text{Mn}_2\text{C}(\text{Fe}_{18}\text{Mn}_{8}\text{Si}_6\text{C}) \rightarrow \text{Fe}_{18}\text{Mn}_8\text{Cr}_6\text{C}$. Thus, dissolving Mn in austenite increases its electrochemical stability while the further dissolution of Si in it does not significantly affect its value. In this case, the dissolution of Cr in manganese austenite leads to a significant increase in electrochemical stability, which can manifest in the growth of corrosion resistance.

The analysis of electron distribution density maps for model structures in the plane (002) (Fig. 7) shows that interatomic bonds with high concentrations of Fe-C and Cr-C electrons are formed in $\text{Fe}_2\text{Mn}_2\text{C}$ and $\text{Fe}_{18}\text{Mn}_8\text{Si}_6\text{C}$ structures, respectively. For the $\text{Fe}_{18}\text{Mn}_8\text{Si}_6\text{C}$ structure, the density of electrons in Si–C pairs is much lower but the relatively high density of electrons is found in Fe-Si pairs. Bonds
in pairs of metal-carbon are characterized by a covalent type, which, in a general case, is reflected on the mechanical properties such as a decrease in plasticity. Therefore, the nature of the distribution of electron density indicates that, for Fe$_{24}$Mn$_8$C, the plasticity would be lower compared to Fe$_{24}$Mn$_8$SiC. For Fe$_{24}$Mn$_8$SiC, the low concentration of electrons around the C atom implies the increased plasticity, on the one hand, while the presence of the developed covalent metal bonds metal–silicon increases hardness, on the other hand.

![Fig. 6. Complete density of electron states of austenite of different formulation](image)

Thus, the dissolution of Si and Cr in both cases causes significant enough changes in the distribution of the density of electrons, which leads to a simultaneous increase in the hardness and plasticity indicators of the austenite phase.

6. Discussion of results of analyzing the impact of alloying with Si and Cr on the structure of manganese austenite

The generalization of results from our experimental research and the results of modeling shows that the dissolution of Si and Cr leads to the increased stability of the austenite phase (Fig. 4, d). Increasing the stability of austenite (Fig. 5) is a prerequisite for preventing the process of cement phase evolution, which negatively affects the capability of manganese austenite to be used for surfacing. On the other hand, the dissolution of Si and Cr leads to changes in mechanical characteristics, in particular to a decrease in the elastic constants (Fig. 4, a) and an increase in plasticity (Fig. 4, b). The theoretical hardness of manganese austenite, calculated from equation (5) (Fig. 4, d), correlates with the level of its stability and the nature of the formation of interatomic bonds (Fig. 7). When dissolving Si in manganese austenite, its stability is higher than when dissolving Cr; hence, its theoretical hardness is also higher although the level of electrochemical stability (Fig. 6) is higher when dissolving Cr.

Our results are useful for the development of new systems to alloy materials for wear-resistant surfacing based on high-manganese steel, under operating conditions involving dynamic and cyclical loads. The scope of application of such materials includes the working surfaces of milling units, agricultural machinery, as well as equipment for woodworking and processing of timber biomass.

The main limitation in the use of our data in relation to actual systems is the need to extrapolate them to the ratio of components that cannot be set for supercells (Fig. 1) due to a significant labor intensity of calculations. In addition, due to the presence of defects in actual structures, such as vacancies, dislocations, grain boundaries, etc., the calculated hardness values could mainly be used for a comparative analysis only.

The presence of Si dissolved in the austenite is a prerequisite for the fundamental possibility of applying coatings by electric arc methods, that is, Si is primarily a technological additive. In contrast to Si, introducing Cr into austenite aims to enhance performance, namely corrosion resistance and hardness. Based on this, further studies should investigate the total impact of Si and Cr, as well as other carbide-forming elements that may dissolve in small quantities in manganese austenite.

7. Conclusions

1. It was established that the main phase in high-manganese steel, containing, % by weight: Mn ~19 %, Si ~4 %, C ~1.5, made by electric arc fusion, is γ-Fe with a lattice parameter of $a=3.576\,\text{Å}$. Analyzing it by a Mossbauer spectroscopy method shows that the Si and Mn dissolved in γ-Fe mainly occupy those positions in the lattice relative to which the carbon atom is in the first or second coordinating sphere.

2. The ab initio simulation and the thermodynamic calculations using a CALPHAD method helped establish that dissolving Si and Cr in manganese austenite leads to an increase in its stability. In this case, plasticity and theoretical hardness increase while the value of elastic constants decreases. Si dissolution has a more significant impact compared to Cr, both to increase the stability of austenite and to change its mechanical characteristics.

3. Based on the results of calculating the density of electron states, it is found that the dissolution of Si and Cr does not significantly affect the electron structure of manganese austenite. This, in particular, indicates that the stacking fault energy (SFE) value would not change significantly, and, consequently, the mechanism of the deformation strengthening of manganese austenite would not change. The lowest density of electron states at the Fermi level was found for the manganese austenite alloyed with chromium, indicating the highest level of its electrochemical stability among the structures examined.
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