Muonium in Stishovite: Implications for the Possible Existence of Neutral Atomic Hydrogen in the Earth’s Deep Mantle

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Hydrogen in the Earth’s deep interior has been thought to exist as a hydroxyl group in high-pressure minerals. We present Muon Spin Rotation experiments on SiO₂ stishovite, which is an archetypal high-pressure mineral. Positive muon (which can be considered as a light isotope of proton) implanted in stishovite was found to capture electron to form muonium (corresponding to neutral hydrogen). The hyperfine-coupling parameter and the relaxation rate of spin polarization of muonium in stishovite were measured to be very large, suggesting that muonium is squeezed in small and anisotropic interstitial voids without binding to silicon or oxygen. These results imply that hydrogen may also exist in the form of neutral atomic hydrogen in the deep mantle.

Hydrogen is the most abundant element in the solar system. It binds to oxygen and the resultant water makes the Earth a habitable blue planet. Ocean covers 70% of the Earth’s surface. Moreover, a significant amount of water may be hidden in the Earth’s interior. The water in anhydrous silicates is considered to be related to lattice defects and incorporated into the system, for example, by simultaneous substitutions such as Mg²⁺ → H⁺ and O²⁻ → OH⁻ (Ref. 1). If a structural change is induced by significant substitutions, the system should be classified in hydrous silicates. In the field of earth science, the water in anhydrous silicates has been studied mainly by the methods to probe the oscillation of hydroxyl group, such as infrared absorption and Raman scattering.⁶⁻⁸ It has been reported that the mantle-transition zone can accommodate a significant amount of H₂O component up to several times as much as the total mass of ocean¹ and, although it is not a consensus⁴, the lower mantle could also accommodate a non-negligible amount⁵.

At the same time, one can speculate that hydrogen might exist in silicates without substitutions. In the field of applied physics, hydrogen in perfect crystals of low-pressure phases of SiO₂ has been studied by the methods to probe H more directly⁶⁻¹⁰, such as Muon Spin Rotation (μSR) and Electron Paramagnetic Resonance (EPR), to obtain a better understanding of Metal-Oxide-Semiconductor (MOS) devices. In those studies, it has been reported that hydrogen can exist in interstitial voids of structure. Moreover, it has recently been revealed that a significant amount of small molecules, such as H₂ and He, dissolves into interstitial voids of low-pressure phases of SiO₂ (Refs. 11–15). Therefore, it seems possible that the hydrogen which is not directly related to lattice defects of silicates and has not so far been recognized sufficiently in earth science may exist in the mantle. However, because interstitial voids of low-pressure phases of SiO₂ are large, the above findings could be exceptional. The structures of two important SiO₂ minerals, stishovite and quartz, are compared in Figure 1. Stishovite is a rutile-type high-pressure phase and quartz is an ambient-pressure phase. The two structures show marked contrast in interstitial voids, in addition to the well-known difference in coordination number.
Here, we present the results of μSR spectroscopy to show the possible existence of neutral atomic hydrogen in small and anisotropic interstitial voids of stishovite. Although μSR spectroscopy is not very popular in earth science, it is a well-established technique to simulate the behavior of hydrogen in semiconductors and insulators. Positive muon can be considered as a light isotope of proton with a mass of about 1/9 that of proton. In μSR, spin-polarized positive muons are implanted into matter and their subsequent behaviors (changes in spin polarization with time) are measured under various conditions of magnetic field. μSR is a powerful tool to probe the position, electronic state, and other information of the hydrogen that exists irregularly in matter, because it tracks the behavior of each implanted particle (in contrast to diffraction). In this study, μSR was conducted on powder and pellet samples to clarify intrinsic properties of stishovite (see Methods).

Results
An example of the time evolution of muon-spin polarization in stishovite powder is shown in Figure 2a. These spectra were measured at a transverse field of 350 G (nominal) at TRIUMF (Canada). The upper and lower limits of the vertical axis of the graph correspond to the amplitude for fully spin-polarized muons. This figure clearly indicates that the fraction of muon staying in the initial diamagnetic state (corresponding to OH forming hydrogen) is small. The muon fraction in stishovite increased slightly with decreasing temperature; 14% at 300 K, 17% at 250 K, 16% at 200 K, 17% at 100 K, 17% at 50 K, and 19% at 2.5 K in stishovite powder.

The spectrum at 300 K in Figure 2a is enlarged in terms of the horizontal axis and is compared with quartz in Figure 2b,c. High-frequency oscillation of muonium-spin polarization is seen. The fraction of muonium (= muon-electron bound paramagnetic state, corresponding to neutral atomic hydrogen) can be determined from the amplitude of oscillation at time = 0 after subtracting the contribution of muon-spin polarization. Only the triplet state, in which muon and electron have spins of the same direction, appears in these spectra and the singlet state, in which they have spins of the opposite direction, has the same population as the triplet and does not appear; twice the amplitude of the triplet state corresponds to the fraction of muonium. It was determined to be 56% in stishovite at 300 K. The sum of muon and muonium fractions is smaller than 100% (Table 1). The remaining 30% was missing due to fast relaxation in spin polarization. These spectra also show that the relaxation rate of muonium-spin polarization in stishovite is large.

The beat of oscillation due to the splitting of the triplet state under transverse field is also seen in Figure 2b,c. The frequencies determined from the spectrum for stishovite are 452(10) and 562(32) MHz. The hyperfine-coupling parameter of muonium A/h, where h is the Plank constant, can be determined from these frequencies. A/h’s for powder and pellet samples agree well with each other (Table 1). A/h for muonium in stishovite at 300 K is determined to be 4.67(3) GHz with all the spectra for the two samples at transverse fields from 200 to 1000 G (nominal). A/h at 2.5 K is 5.17(14) GHz, estimated only from the two frequencies at 350 G (nominal). These values are significantly larger than 4.49(2) GHz in quartz (Table 1) and 4.463 GHz in vacuum. Data for stishovite plotted on the Breit-Rabi diagram are shown in Figure 3.

Experimental results at TRIUMF are summarized in Table 1. In both powder and pellet samples of stishovite, the muon fraction was small and decreased with increasing temperature. Also, A/h’s are the same for the two samples as mentioned above. On the other hand, some differences are found; the muon fraction and the relaxation rate of muonium-spin polarization were smaller in powder than in pellet. In this study, experiments were also conducted at RIKEN/RAL (UK), PSI (Switzerland), and MLF/J-PARC (Japan). Preliminary results at these facilities, such as the muon fraction and A/h in stishovite powder (roughly estimated based on decoupling measurements under...
concentration. Interstitial voids of stishovite, which consists of SiO$_6$ octahedra, are much smaller than those of quartz, which consists of SiO$_4$ tetrahedra (Fig. 1). Therefore, the present results suggest that the formation of muonium is not controlled by the size of interstitial voids. In stishovite, not only the muonium fraction was large but also the muon fraction decreased with increasing temperature, suggesting that implanted muons become more stable by capturing an electron to form muonium than by staying in diamagnetic state. Hyperfine-coupling parameter is a measure of 1s electronic orbital size; the cube of Bohr radius is inversely proportional to this parameter$^{16,17}$. Therefore, a very large hyperfine-coupling parameter of muonium in stishovite, which is even larger than that in quartz, suggests that muonium is squeezed in small interstitial voids without binding to silicon or oxygen. A very large relaxation rate of muonium-spin polarization cannot be explained by the effect of nuclear magnetic moments of silicon and oxygen because they are small. It may be due to anisotropy in hyperfine interactions. This explanation seems plausible because the relaxation rate becomes larger at low temperatures where the mobility of muoniums becomes smaller. In fact, interstitial voids in stishovite are largely anisotropic (i.e., having channels along the c-axis; Fig. 1).

The difference in muonium fraction and relaxation rate of muonium-spin polarization between powder and pellet suggests that they are affected by the grain size and/or surface condition (= defects, in a broad sense) of samples. The pellet sample is a sintered body of nanocrystals and therefore has a much smaller grain size than the powder sample. A negative correlation between grain size and muon fraction may be explained by the inhibition of delayed-muonium formation, in which positive muons capture radiolysis electrons and become muoniums after time $t = 0$ (Ref. 10). In quartz single crystal, it has been reported that muon and muonium fractions were 20% and 80%, respectively, and about the half of muonium was formed promptly after time $t = 0$ (as delayed muoniums)$^{10}$. On the other hand, in quartz powder, the muon fraction was 28% (Table 1). Therefore, a negative correlation between grain size and muon fraction is also seen in quartz. The delayed formation of muonium substantially (but very shortly) after time $t = 0$ causes the de-phasing of spin polarization. This may explain the missing fraction in powder and pellet samples; i.e., the sum of muon and muonium fractions is smaller than 100% (Table 1). At low temperatures, the muon fraction becomes larger probably because the mobility of radiolysis electrons becomes smaller and therefore the formation of delayed muonium is inhibited. In any case, muonium seems to be more easily formed at higher temperatures and/or in more perfect crystals of stishovite.

### Table 1 | Summary of μSR measurements for stishovite$^a$

| Material    | Form   | Temperature [K] | Statistics [events] | $f_m$ [%] | $\lambda_m$ [μs$^{-1}$] | $f_{\text{Mu}}$ [%] | $\lambda_{\text{Mu}}$ [μs$^{-1}$] | $A/h$ [GHz] |
|-------------|--------|-----------------|---------------------|-----------|------------------------|-----------------|--------------------------|-------------|
| Stishovite  | Powder | 300             | 100 M               | 14.4(3)   | 0.10(1)                | 56(5)           | 24(4)                    | 4.65(2)     |
|             |        | 2.5             | 100 M               | 19.2(4)   | 0.13(1)                | 4.68(5)         | 5.17(14)                 |             |
| Pellet$^b$  | Powder | 300             | 100 M               | 32.1(5)   | 0.05(1)                | 55(9)           | 55(9)                    | 4.68(5)     |
| Quartz      | Powder | 300             | 25 M                | 39.6(8)   | 0.06(1)                | 56(8)           | 7(5)                     | 4.49(2)     |

$^a$ $f_m$: fraction of muon, $\lambda_m$: relaxation rate of muon, $f_{\text{Mu}}$: fraction of muonium, $A/h$: hyperfine-coupling parameter of muonium, where $h$ is the Plank constant (see Methods for more details). Measurements shown in Figure 3 were used to determine $A/h$ for stishovite and measurements at 350 and 700 G (nominal) were used to determine $A/h$ for quartz. Measurements at 350 G (nominal) were used to determine the other parameters.

$^b$ Although the simultaneous determination of $f_{\text{Mu}}$ and $\lambda_{\text{Mu}}$ was not successful due to the very fast relaxation, the relation between the two parameters can be expressed well as $f_{\text{Mu}} = 0.83\lambda_{\text{Mu}} + 8.1$.

In this study, μSR experiments have suggested that the neutral atomic hydrogen (corresponding to muonium) can exist in stishovite without having a direct relation to lattice defects. So, the question is whether invisible hydrogen really exists in stishovite, i.e., without being detected by infrared absorption, Raman scattering, and other standard methods. Although proving the existence is difficult, a clue can be found in a report on MOS devices$^{29}$. Infrared absorption due to Si-OH and Si-H oscillations was detected in $\gamma$-ray irradiated SiO$_2$ films of MOS devices, while it was not detected in non-irradiated films. It has been interpreted as a sign that H and/or H$_2$ released from an electrode of the device (hydrogen source) exist in the film and react with dangling bonds of silicon and oxygen generated by the $\gamma$-ray irradiation. Therefore, we believe that invisible hydrogen exists similarly in stishovite if hydrogen sources are available. However, even in the presence of hydrogen sources, oxidizing environment would favor the incorporation of water into stishovite by a substitution of Si$^{4+}$ $\rightarrow$ 4H$^+$ (Ref. 5). The situation may be similar in other high-pressure silicates. In previous studies of earth science, oxidizing environment has often been assumed and many experiments have been conducted in various silicate-water systems. However, the redox state of the deep mantle (from past to present) is a challenging issue$^{20-23}$ and it may be much more reducing compared to the present shallow mantle$^{20-22}$.

### Figure 3 | Energy diagram of muonium in stishovite as a function of applied transverse field, so called Breit-Rabi diagram$^{16,17}$. $x$ denotes the transverse field that is normalized by the field corresponding to $A/h$ (Eq. 8). The three solid lines, in order from top to bottom, represent theoretical $E_1/A$, $E_2/A$, and $E_3/A$ (Eqs. 12–14). The data of $E_1/A$ and $E_2/A$, from left to right, correspond to the measurements at 200, 350, 500, 700, and 1000 G (nominal). Because of a larger $A/h$ at 2.5 K than at 300 K, the data for 2.5 K are located at the left side of corresponding data at 300 K.
Now we assume reducing environment in the deep mantle. Then, the question is whether hydrogen sources are available in the Earth’s interior. A candidate may be the core. In some models of early earth evolution, a tremendous amount of hydrogen dissolved into the core\(^2\text{-}3\). Or, hydrogen may have dissolved directly into the mantle at an early stage and stay there quietly\(^20\text{-}22\); the chemoaffinity between silicates and liquid H\(_2\) under high pressure has been demonstrated recently\(^27\). Therefore, it is possible that the neutral atomic hydrogen which is not directly related to lattice defects of silicates may exist in the mantle. To further test the hypothesis, \(\mu\)SR experiments should be conducted on more realistic mantle minerals and rocks. In-situ experiments under high pressure and high temperature are also indispensable. Definitely, studies from multiple viewpoints are required to know how much hydrogen can dissolve into silicates in a thermodynamically stable state. For example, accurate calculation of the formation enthalpy should be made with the help of knowledge obtained by experimental work. Neutral atomic hydrogen in the mantle will be an important research target in future earth science, because the existence form of this important element has a fundamental importance in physicochemical properties of mantle minerals, thus controls the dynamics and evolution of our planet.

**Methods**

**Sample preparation.** \(\mu\)SR spectroscopy requires a relatively large amount of samples. So, nominally anhydrous stishovite samples were synthesized under high pressure and high temperature with a belt-type large-volume press\(^4\) at National Institute for Materials Science (Japan) and with a Kawai-type large-volume press\(^5\) at Geodynamics Research Center of Ehime University (Japan), respectively. To clarify intrinsic properties of stishovite, two kinds of samples were prepared: powder sample was synthesized with a standard technique at the former institute and pellet sample (sintered nanocrystals)\(^28\) was synthesized with a recently developed technique at the latter institute. Stishovite powder of about 0.5 g was synthesized from noncrystalline SiO\(_2\) powder (Kanto Chemical Co., Inc., purity 99.9%) at 10 GPa and 1373 K with the former press. For reference in \(\mu\)SR experiments, quartz powder was also synthesized from the same starting material at 2 GPa and 1373 K with the same type of press. Each synthetic powder was packed in a capsule made of aluminum foil (having a thickness of 12 \(\mu\)m). Stishovite pellet (sintered nanocrystals) of 5.5 mm in diameter and 1.0 mm in thickness was synthesized with the latter press. More detailed information on the stishovite pellet has been given elsewhere\(^6\).

**Experimental procedure.** \(\mu\)SR experiments were conducted for stishovite and quartz with surface-\(\mu\)on beams at RIKEN/RAL (UK), PSI (Switzerland), MLF/J-PARC (Japan), and TRIUMF (Canada). Spin-polarized positive muons were implanted into the capsulated powder or pellet and subsequent behaviors (changes in spin polarization with time) were measured under transverse magnetic field (perpendicular to the initial spin-polarization direction, up to 1000 G) or longitudinal magnetic field (parallel to the initial spin-polarization direction, up to 4000 G) at temperatures between 2.5 and 300 K (room temperature). Stishovite powder was measured at all the four facilities and stishovite pellet was measured only at TRIUMF. Quartz powder was measured at MLF/J-PARC and TRIUMF. Based on preliminary results at RIKEN/RAL, PSI, and MLF/J-PARC, experiments with a high timing resolution, which is suitable for the observation of muonium, were conducted by using a continuous beam and a spectrometer called HiTime at TRIUMF.

**Analytical procedure.** Standard procedure was followed to analyze \(\mu\)SR data\(^6\text{-}17\). It is briefly summarized below.

First, data obtained with HiTime under transverse magnetic field were analyzed in frequency domain. Angular frequencies of muon-spin polarization \(\omega_0\) and muonium-spin polarization \(\omega_{\text{Mu1}}\) and \(\omega_{\text{Mu2}}\) were determined by the Fourier transform. Because of the splitting of the triplet state, two frequencies appear for muonium. Thus determined \(\omega_0\) and gyromagnetic ratio of muon \(\gamma_0\) were used to determine the transverse magnetic field \(H\) actually applied to the sample with the equation

\[
\omega_0 = \frac{\gamma_0}{2\pi} H.
\]

Here, \(\gamma_0\) is the gyromagnetic ratio of electron.

Hyperfine-coupling parameter of muonium A/H relates to \(\omega_{\text{Mu1}}\) and \(\omega_{\text{Mu2}}\) with the following equations;

\[
A = \frac{1}{2\pi} \frac{\omega_0^2 - (\omega_{\text{Mu1}} - \omega_{\text{Mu2}})^2}{\omega_{\text{Mu1}} - \omega_{\text{Mu2}}},
\]

\[
A = \frac{1}{2\pi} \frac{\omega_0^2 - (\omega_{\text{Mu1}} - \omega_{\text{Mu2}})^2}{\omega_{\text{Mu1}} - \omega_{\text{Mu2}}}.
\]

Next, data at 350 G (nominal) were analyzed in time domain with angular frequencies obtained by the Fourier transform. The equation

\[
A(t) = A_0 \cos(\omega_{\text{Mu1}} t) \exp(-\lambda_{\text{Mu1}} t) + A_{\text{Mu2}} \cos(\omega_{\text{Mu2}} t + \phi) \exp(-\lambda_{\text{Mu2}} t),
\]

where

\[
A_{\text{Mu1}} = \frac{1 + x + x^2}{1 - x + x^2},
\]

\[
x = \frac{\gamma_0 + \gamma_H}{2\pi} \frac{h}{A},
\]

Here, phase \(\phi\) was introduced to correct the error in time \(t = 0\). The difference in relaxation rates of two frequencies was assumed to be negligible. Then, muon fraction \(f_\mu\) and muonium fraction \(f_{\text{Mu}}\) were determined with the equations

\[
f_\mu = \frac{A_{\text{Mu1}}}{A_{\text{Mu1}} + A_{\text{Mu2}}}.
\]

\[
f_{\text{Mu}} = \frac{2 A_{\text{Mu1}} + A_{\text{Mu2}}}{A_{\text{Mu1}} + A_{\text{Mu2}}},
\]

where

\[
e_1 = \frac{1}{4} A + \frac{1}{4} \left( 4 \times \frac{1}{2} h \omega_{\text{Mu1}} \right)^2
\]

\[
e_2 = \frac{1}{4} A + \frac{1}{4} \left( 4 \times \frac{1}{2} h \omega_{\text{Mu1}} \right)^2.
\]

1. Smyth, J. R. \(\beta\)-\(\text{Mg}_2\text{SiO}_4\): A potential host for water in the mantle? *Am. Mineral.* 72, 1051–1055 (1987).
2. Williams, Q. & Hemley, R. J. Hydrogen in the deep Earth. *Annu. Rev. Earth Planet. Sci.* 29, 365–418 (2001).
3. Murakami, M., Hirose, K., Yurimoto, H., Nakashima, S. & Takafuji, N. Water in Earth’s lower mantle. *Science* 295, 1885–1887 (2002).
4. Bolfan-Casanova, N. Water in the Earth’s mantle. *Mineral. Mag.* 69, 229–257 (2005).
5. Spektor, K. et al. Ultrahydrostatic stishovite from high-pressure hydrothermal treatment of SiO\(_2\). *Proc. Natl. Acad. Sci. USA* 108, 20918–20922 (2011).
6. Myasishcheva, G. G., Obukhov, Yu. V., Roganov, V. S. & Firsov, V. G. A search for atomic muonium in chemically inert substances. *Sozhy. Phys. JETP* 26, 298–301 (1968).
7. Gurievich, I. I. et al. Muonium in magnetic field. *Phys. Lett.* 29B, 387–390 (1969).
8. Brown, J. A. et al. A precise determination of the hyperfine splitting of muonium in quartz. *Solid State Commun.* 33, 613–614 (1980).
9. Isoya, J., Weil, J. A. & Davis, P. H. EPR of atomic hydrogen \(1\)H and \(2\)H in z-quartz. *J. Phys. Chem. Solids* 44, 335–343 (1983).
10. Brewer, J. H. et al. Delayed muonium formation in quartz. *Physica B* 289–290, 425–427 (2000).
11. Sato, T., Funamori, N. & Yagi, T. Helium penetrates into silica glass and reduces its compressibility. Nature Commun. 2, 345 (2011).

12. Shen, G. et al. Effect of helium on structure and compression behavior of SiO2 glass. Proc. Natl. Acad. Sci. USA 108, 6004–6007 (2011).

13. Efimchenko, V. S., Fedotov, V. K., Kuzovnikov, M. A., Zhuravlev, A. S. & Bulychev, B. M. Hydrogen solubility in amorphous silica at pressures up to 75 kbar. J. Phys. Chem. B 117, 422–425 (2013).

14. Sato, T. et al. Anomalous behavior of cristobalite in helium under high pressure. Phys. Chem. Miner. 40, 3–10 (2013).

15. Matsuı̂, M., Sato, T. & Funamori, N. Crystal structures and stabilities of cristobalite-helium phases at high pressures. Am. Mineral. 99, 184–189 (2014).

16. Schenck, A. Muon Spin Rotation Spectroscopy: Principles and Applications in Solid State Physics (Adam Hilger Ltd., Bristol and Boston 1985).

17. Yaouanc, A. & de Réotier, P. D. Muon Spin Rotation, Relaxation, and Resonance: Applications to Condensed Matter (Oxford University Press, Oxford 2011).

18. Liu, W. et al. High precision measurements of the ground state hyperfine structure interval of muonium and of the muon magnetic moment. Phys. Rev. Lett. 82, 711–714 (1999).

19. Nagasawa, Y. et al. The study of the thermal oxide films on silicon wafers by Fourier transform infrared attenuated total reflection spectroscopy. J. Appl. Phys. 68, 1429–1434 (1990).

20. Ballhaus, C. Is the upper mantle metal-saturated? Earth Planet. Sci. Lett. 132, 75–86 (1995).

21. Rohrbach, A. et al. Metal saturation in the upper mantle. Nature 449, 456–458 (2007).

22. Frost, D. J. & McCammon, C. A. The redox state of Earth’s mantle. Annu. Rev. Earth Planet. Sci. 36, 389–420 (2008).

23. Siebert, J., Badro, J., Antonangeli, D. & Ryerson, F. J. Terrestrial accretion under oxidizing conditions. Science 339, 1194–1197 (2013).

24. Fukai, Y. The iron-water reaction and the evolution of the Earth. Nature 308, 174–175 (1984).

25. Shinozaki, A. et al. Influence of H2 fluid on the stability and dissolution of Mg2SiO4 forsterite under high pressure and high temperature. Am. Mineral. 98, 1604–1609 (2013).

26. Kanke, Y., Akaishi, M., Yamaoka, S. & Taniguchi, T. Heater cell for materials synthesis and crystal growth in the large volume high pressure apparatus at 10 GPa. Rev. Sci. Instrum. 73, 3268–3270 (2002).

27. Irifune, T., Isobe, F. & Shinmei, T. A novel large-volume Kawai-type apparatus and its application to the synthesis of sintered bodies of nano-polycrystalline diamond. Phys. Earth Planet. Inter. 228, 255–261 (2014).

28. Nishiyama, N. et al. Synthesis of nanocrystalline bulk SiO2 stishovite with very high toughness. Scr. Mater. 67, 955–958 (2012).

Acknowledgments
μSR experiments were conducted at RIKEN/RAL, PSI, MLF/J-PARC, and TRIUMF. The authors are grateful to beam line staffs of these facilities for experimental support and K. Nishiyama, K. Shimomura, and S. Tsuneyuki for invaluable discussion on the muonium state in matter. This work was in part supported by a Grant-in-Aid for Scientific Research (Japan) and Special Coordination Funds for Promoting Science and Technology (Japan).

Author contributions
N.F., K.K.M., D.W., T.S., A.K. and R.K. designed the study. T.T., N.N. and T.I. prepared the samples. N.F., K.K.M., D.W., T.S., T.M., M.M., M.H., A.K. and R.K. conducted the μSR experiments and analyzed the data. N.F., K.K.M., D.W., T.S. and R.K. wrote the manuscript and others gave comments to improve the manuscript.

Additional information
Competing financial interests: The authors declare no competing financial interests.

How to cite this article: Funamori, N. et al. Muonium in Stishovite: Implications for the Possible Existence of Neutral Atomic Hydrogen in the Earth’s Deep Mantle. Sci. Rep. 5, 8437; DOI:10.1038/srep08437 (2015).

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