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Carbon-depleted outer core revealed by sound velocity measurements of liquid iron–carbon alloy

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The relative abundance of light elements in the Earth's core has long been controversial. Recently, the presence of carbon in the core has been emphasized, because the density and sound velocities of the inner core may be consistent with solid Fe_7C_3 . Here we report the longitudinal wave velocity of liquid $\text{Fe}_{84}\text{C}_{16}$ up to 70 GPa based on inelastic X-ray scattering measurements. We find the velocity to be substantially slower than that of solid iron and Fe_3C and to be faster than that of liquid iron. The thermodynamic equation of state for liquid $\text{Fe}_{84}\text{C}_{16}$ is also obtained from the velocity data combined with previous density measurements at 1 bar. The longitudinal velocity of the outer core, about 4% faster than that of liquid iron, is consistent with the presence of 4–5 at.% carbon. However, that amount of carbon is too small to account for the outer core density deficit, suggesting that carbon cannot be a predominant light element in the core.

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Sound velocity and density are important observational constraints on the chemical composition of the Earth's core. While properties of solid iron alloys have been extensively examined by laboratory studies to core pressures (> 136 GPa)^{1–3}, little is known for liquid alloys because of experimental difficulties. The core is predominantly molten, and the longitudinal wave (P-wave) velocity of liquid iron alloy is the key to constraining its composition. However, previous static high-pressure and -temperature (P - T) measurements of liquid iron alloys were performed only below 10 GPa using large-volume presses^{4–6}. Shock wave experiments have been carried out at much higher pressures but only along a specific Hugoniot P - T path^{7,8}.

Carbon is one of the possible light alloying components in the core because of its high cosmic abundance and strong chemical affinity with liquid iron⁹. Its high metal/silicate partition coefficients indicate that thousands of parts per million to several weight percent of carbon could have been incorporated into the core during its formation^{9–11}. In addition, recent experimental and theoretical studies^{12,13} have suggested that solid Fe_7C_3 may explain the properties of the inner core, in particular its high Poisson's ratio^{14,15}, supporting the presence of carbon in the core.

In this study, we determine the P-wave velocity (V_P) (equivalent to bulk sound velocity, V_Φ , in a liquid) of liquid $\text{Fe}_{84}\text{C}_{16}$ at high P - T based on inelastic X-ray scattering (IXS) measurements. Combined with its density data at 1 bar (ref. 16) both velocity and density (ρ) profiles of liquid $\text{Fe}_{84}\text{C}_{16}$ along adiabatic compression are obtained. They are compared with seismological observations, indicating that both V_P and ρ in the Earth's outer core are not explained simultaneously by liquid Fe-C.

Results

Longitudinal wave velocity measurements. We collected the high-resolution IXS spectra from liquid $\text{Fe}_{84}\text{C}_{16}$ (4.0 ± 0.3 wt.% carbon) at static high P - T using both resistance- and laser-heated diamond-anvil cells (Methods; Fig. 1). The starting material was synthesized beforehand as a mixture of fine-grained Fe and Fe_3C at 5 GPa and 1,623 K in a multi-anvil apparatus. Experimental P - T conditions were well above the eutectic temperature in the Fe- Fe_3C binary system (Supplementary Fig. 1). The carbon concentration in the eutectic liquid is known to be 3.8–4.3 wt.% at 1 bar to 20 GPa (ref. 17), almost identical to the composition of our sample. Above 20 GPa, we heated the sample to temperatures comparable or higher than the melting temperature of Fe_3C , a liquidus phase in the pressure range explored, assuring a fully molten sample. The molten state of the specimen was carefully confirmed, before and after the IXS measurements, by the absence of diffraction peaks from the sample (Fig. 2). We sometimes, depending on a sample volume, were also able to observe the diffuse diffraction signal typical of a liquid.

The V_P of liquid $\text{Fe}_{84}\text{C}_{16}$ was determined between 7.6 and 70 GPa (Fig. 3 and Supplementary Table 1) from dispersion curves for a range of momentum transfer (Fig. 4). It was found to be 15–30% smaller than that of solid Fe (refs 3,18–20) and Fe_3C (refs 21–23; note that a starting material in the present experiments was a mixture of these solid phases) (Fig. 5), confirming that we measured a liquid sample. The velocities of a fictive solid $\text{Fe}_{84}\text{C}_{16}$ alloy are also estimated assuming a linear velocity change between Fe (ref. 24) and Fe_3C (ref. 23) indicating that V_P drops by 13% upon melting at 2,300 K, a eutectic temperature at 45 GPa (ref. 17). Such a velocity change is comparable to that expected for pure Fe. The difference in V_Φ between solid and liquid $\text{Fe}_{84}\text{C}_{16}$ is very small (1.8%). On the

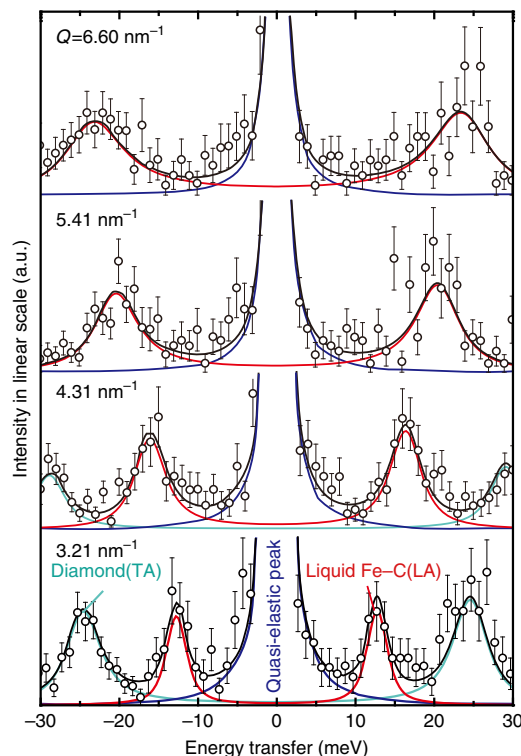


Figure 1 | Typical inelastic X-ray scattering spectra. These data were collected at 26 GPa and 2,530 K at momentum transfers Q , as indicated. The spectra include three components: a quasi-elastic peak near zero energy transfer (blue), longitudinal acoustic (LA) phonon mode of liquid $\text{Fe}_{84}\text{C}_{16}$ (red), and transverse acoustic (TA) phonon mode of diamond (turquoise).

other hand, the V_P of our liquid $\text{Fe}_{84}\text{C}_{16}$ sample is 3–14% faster at 8–70 GPa than that of liquid Fe determined by shock-wave study⁸ (Fig. 3).

Earlier ultrasonic measurements performed below 10 GPa reported a change in V_P by < 2 –3% per 1,000 K for liquid Fe-S alloys^{4,5}. Theoretical calculations^{25–27} and shock compression data⁸ on liquid Fe and Fe-S alloy demonstrated even smaller effects above 100 GPa ($< 0.5\%$ by 1,000 K). It is therefore very likely that the V_P of liquid $\text{Fe}_{84}\text{C}_{16}$ is also not sensitive to temperature with the temperature effect much smaller than the uncertainty in the present velocity determinations ($\pm 3\%$).

Thermodynamical equation of state. V_P of a liquid can be described using the Murnaghan equation of state⁴ (Methods) as;

$$V_P = \sqrt{\frac{K_{S0}}{\rho_0} \left(1 + \frac{K'_S}{K_{S0}} P \right)^{\frac{1}{2} - \frac{1}{2K'_S}}}, \quad (1)$$

where K_S and K'_S are adiabatic bulk modulus and its pressure derivative, respectively (zero subscripts denote values at 1 bar and $T = T_0$). Here, consistent with the discussion above, we neglect the temperature dependence of our V_P data, while ρ_0 is taken to be temperature dependent¹⁶ (Methods). We fit equation (1) to our P - V_P data for liquid $\text{Fe}_{84}\text{C}_{16}$ and find $K_{S0} = 110 \pm 9$ GPa and $K'_S = 5.14 \pm 0.30$ when $T_0 = 2,500$ K (Supplementary Table 2 and Supplementary Fig. 2). The choice of T_0 and, accordingly, the variation in ρ_0 practically changed K_{S0} and K'_S as $(\partial K_{S0}/\partial T) = -9.4 \times 10^{-3}$ GPa K⁻¹ and $(\partial K'_S/\partial T) = -2.7 \times 10^{-4}$ K⁻¹. Our value for K_{S0} is similar to that for liquid iron⁸ but for K'_S is higher than that for pure iron, $K'_S = 4.7$.

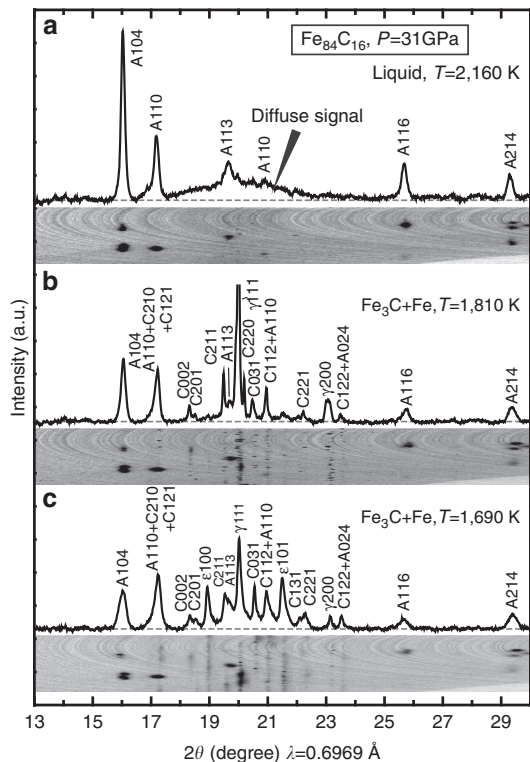


Figure 2 | X-ray diffraction spectra before and after melting. They were collected at 2,160 K (a), 1,810 K (b) and 1,690 K (c) during heating at 31 GPa. The starting material was composed of Fe (ϵ or γ) and Fe_3C (c), and the peaks of Al_2O_3 (a) were from a thermal insulator. The coexistence of ϵ - and γ -Fe phases at 1,610 K was due to a sluggish solid–solid phase transition⁴⁹ and the peaks from the ϵ -phase were lost at 1,810 K. All sample peaks disappeared between 1,810 and 2,160 K. In addition, the background was enhanced slightly, indicating a diffuse scattering signal from a liquid sample.

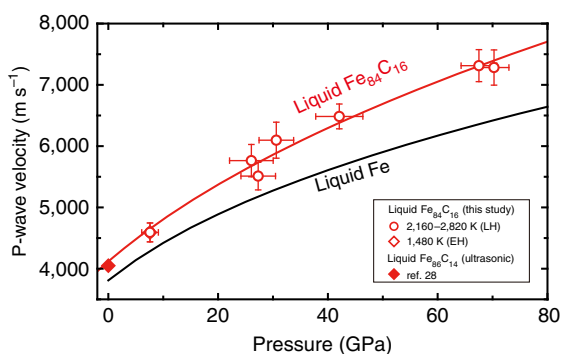


Figure 3 | Compressional wave velocity of liquid $\text{Fe}_{84}\text{C}_{16}$. Open circles, obtained by laser-heated DAC; open diamond, by external-resistance-heated DAC. The data at 1 bar is from ultrasonic measurements²⁸ (closed diamond). The red curve represents a thermodynamical fitting result for liquid $\text{Fe}_{84}\text{C}_{16}$, compared with the velocity of liquid Fe (black curve)⁸.

This suggests that liquid $\text{Fe}_{84}\text{C}_{16}$ becomes progressively stiffer than liquid Fe with increasing pressure. We also found $V_{P0} = 4,121 \pm 177 \text{ m s}^{-1}$ for liquid $\text{Fe}_{84}\text{C}_{16}$ from K_{S0} and ρ_0 , in good agreement with a previous study²⁸ of liquid $\text{Fe}_{86}\text{C}_{14}$ at 1 bar ($4,050 \text{ m s}^{-1}$) and faster than $V_{P0} = 3,860 \text{ m s}^{-1}$ for liquid Fe (ref. 8).

To compare the present results with earlier density measurements of liquid Fe–C alloys at high pressure^{29,30} the isothermal

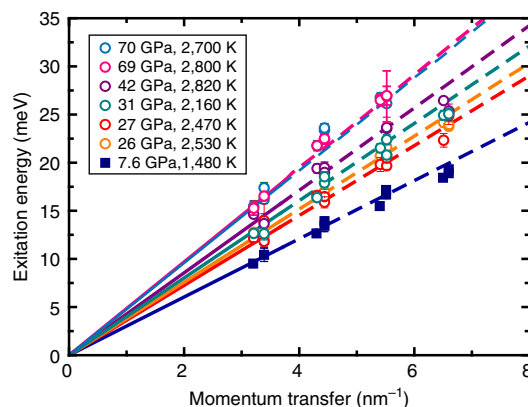


Figure 4 | Longitudinal acoustic phonon dispersion of liquid $\text{Fe}_{84}\text{C}_{16}$. The dispersion data were obtained at pressures from 7.6 to 70 GPa. Only data collected with the low momentum transfer ($<3.5 \text{ nm}^{-1}$) were used to determine the velocity to avoid possible anomalous dispersion for liquid (see Methods).

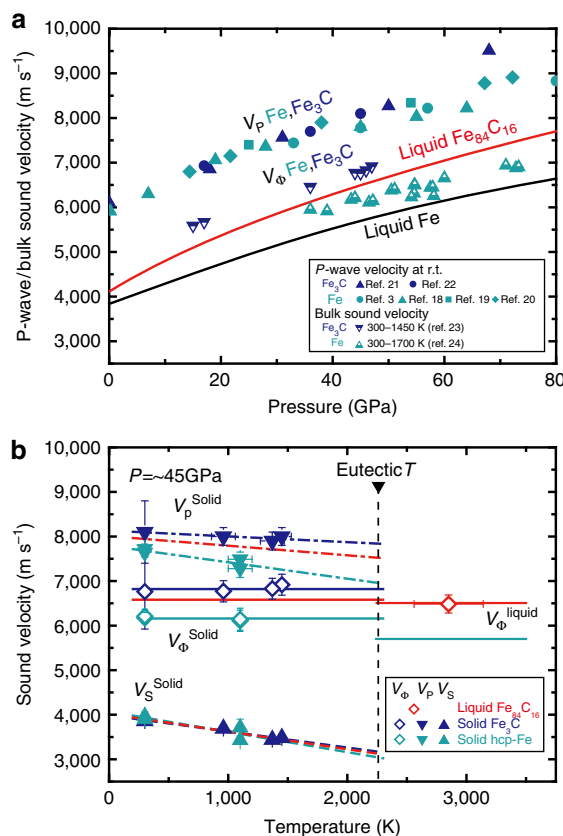


Figure 5 | Comparison of velocities between liquid and solid Fe–C alloys. (a), P-wave velocities (V_p) and bulk sound velocities (V_ϕ) of solid Fe (turquoise)^{3,18–20,24} and Fe_3C (blue)^{21–23} were determined by previous IXS and nuclear inelastic scattering (NIS) measurements. The V_p for liquid Fe is from shock-wave study⁸. (b), Temperature effects on the sound velocities of Fe–C alloys at $\sim 45 \text{ GPa}$. The $V_p (= V_\phi)$ of liquid $\text{Fe}_{84}\text{C}_{16}$ and liquid Fe is from the present work at 42 GPa and shock-wave data⁸, respectively. The V_p , shear velocity (V_s), and V_ϕ for solid Fe and Fe_3C were reported by NIS measurements^{23,24}. Red lines for fictional solid $\text{Fe}_{84}\text{C}_{16}$ are estimated from a linear relationship between Fe and Fe_3C .

bulk modulus for liquid $\text{Fe}_{84}\text{C}_{16}$ is estimated to be $K_{T0} = 100$ (82) GPa at 1,500 K (2,500 K) from our determination of K_S combined with Grüneisen parameter $\gamma_0 = 1.74$ (ref. 8) and

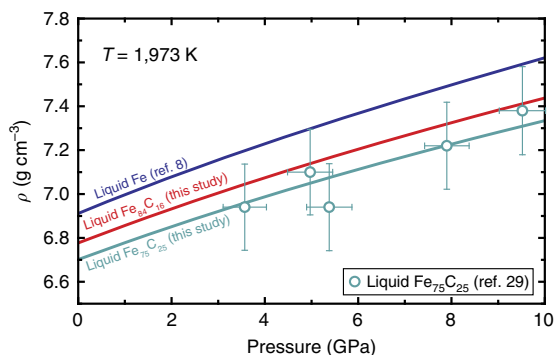


Figure 6 | Comparison with previous density measurements. Blue and red curves demonstrate calculated densities at 1,973 K for liquid Fe (ref. 8) and $\text{Fe}_{84}\text{C}_{16}$ (present study). The density of liquid $\text{Fe}_{75}\text{C}_{25}$ (turquoise curve) is estimated assuming linear compositional dependence between pure Fe and $\text{Fe}_{75}\text{C}_{25}$, which shows good agreement with the previous measurements at 1,973 K (ref. 29).

thermal expansion coefficient¹⁶ (Methods). When applying $C_p/C_V = 1.125$ at 1,820 K for liquid $\text{Fe}_{86}\text{C}_{14}$ derived from theoretical calculations³¹, $K_{T0} = 106\text{--}98$ GPa is obtained at the same temperature range. These K_{T0} values for liquid $\text{Fe}_{84}\text{C}_{16}$ are similar to $K_{T0} = 95\text{--}63$ GPa for liquid Fe at 1,500–2,500 K (ref. 8). On the other hand, they are significantly larger than $K_{T0} = 55.4$ GPa for liquid $\text{Fe}_{86}\text{C}_{14}$ at 1,500 K and $K_{T0} = 50$ GPa for liquid $\text{Fe}_{75}\text{C}_{25}$ at 1,973 K from previous density measurements^{29,30}. However, the calculated density for $\text{Fe}_{84}\text{C}_{16}$ using the present EoS are in reasonable agreement with the previous density measurements of $\text{Fe}_{75}\text{C}_{25}$ (ref. 29) (Fig. 6). The disagreement of elastic parameters with such earlier experiments may be attributed either to the limited pressure range of the previous density determinations, or to a different structure or magnetic (or electronic) change in the state of the liquid Fe–C at low pressure, as has been suggested from the change in compressional behaviour of liquid $\text{Fe}_{78}\text{C}_{22}$ around 5 GPa (ref. 6). Our data were collected above 7.6 GPa, so that the physical properties of liquid Fe–C obtained here should be more applicable to the Earth's core.

ρ of liquid $\text{Fe}_{84}\text{C}_{16}$ is then given, using the elastic parameters determined above, by;

$$\rho = \rho_0 \left(1 + \frac{K'_S}{K_{S0}} P \right)^{\frac{1}{K'_S}}. \quad (2)$$

Equations (1) and (2) give the V_P and ρ profiles for adiabatic compression (Methods), assuming $\gamma_0 = 1.74$, the same as that of liquid Fe (ref. 8) (Fig. 7). We find $V_P = 9,200 \text{ m s}^{-1}$ and $\rho = 9.82\text{--}9.61 \text{ g cm}^{-3}$ at the core-mantle boundary (CMB) for $T_{\text{CMB}} = 3,600\text{--}4,300$ K (refs 32,33). This indicates that V_P of liquid $\text{Fe}_{84}\text{C}_{16}$ is 19.6% faster than that of liquid Fe at the CMB⁸, implying that the addition of 1 at.% carbon increases the V_P of liquid Fe by 1.2%. The extrapolation of the present experimental data using the Murnaghan equation of state may overestimate the V_P by 2–4% at the CMB (Supplementary Note 1 and Supplementary Fig. 3), but, even if this is the case, 1 at.% carbon enhances the V_P of liquid Fe by as large as 0.8%. Indeed, the effect of carbon is much larger than a recent theoretical prediction of only 0.2% increase in velocity per 1 at.% carbon at 136 GPa (ref. 34). On the other hand, our data show that the incorporation of 1 at.% carbon reduces the density of liquid Fe by 0.6–0.7%, while theory suggested only 0.3% density reduction by 1 at.% carbon³⁴.

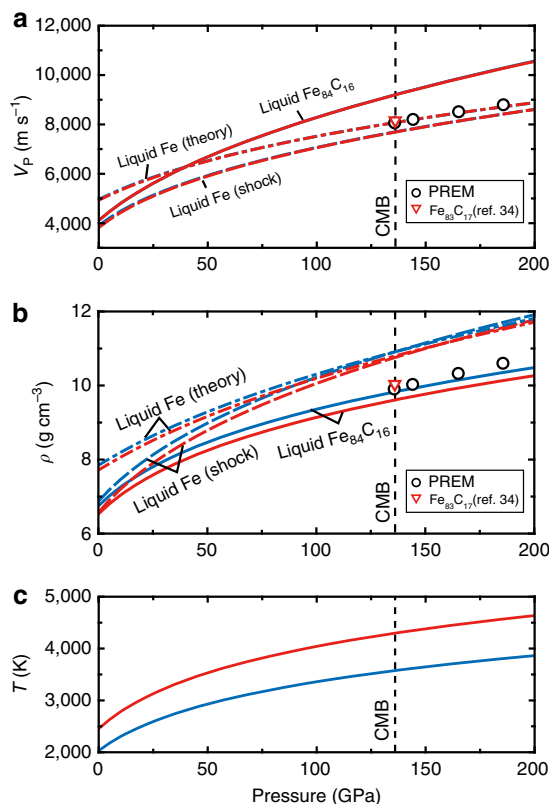


Figure 7 | Velocity and density of liquid $\text{Fe}_{84}\text{C}_{16}$ extrapolated to core pressures. The P-wave velocity (a) and density (b) profiles of liquid $\text{Fe}_{84}\text{C}_{16}$ are calculated along two adiabatic temperature curves (c) of 3,600 K (blue) and 4,300 K (red) at the CMB. Those for liquid Fe are from shock-compression experiments⁸ and theoretical calculations²⁶. The velocity and density for liquid $\text{Fe}_{83}\text{C}_{17}$ at the CMB (4,300 K) are by theory³⁴. PREM denotes seismologically deduced Preliminary Reference Earth model³⁵.

Discussion

We now compare the sound velocity and density of liquid $\text{Fe}_{84}\text{C}_{16}$ and liquid Fe with the seismologically based PREM model³⁵ for the outer core (Fig. 8). The V_P and ρ of liquid Fe are 4.6% slower and 10.1–8.6% denser, respectively, than the PREM at the CMB (3,600–4,300 K). To match the PREM values, considering the uncertainty of data extrapolation to higher pressures (Supplementary Note 1), only 5.2–4.0 at.% (1.2–0.9 wt.%) carbon is required to match the velocity, whereas 15.4–12.0 at.% (3.8–2.9 wt.%) carbon is necessary to account for the density. Therefore, carbon cannot be a predominant light element in the outer core.

These results suggest there is <5.2 at.% (1.2 wt.%) carbon in the outer core, consistent with the previous cosmochemical and geochemical arguments. In particular, the silicate portion of the Earth exhibits much higher $^{13}\text{C}/^{12}\text{C}$ isotopic ratio than that of Mars, Vesta and chondrite meteorites, as may be attributed to a strong enrichment of ^{12}C in core-forming metals⁹. The carbon isotopic fractionation that occurred during continuous core-formation process proposed previously^{36,37} will give a reasonable $^{13}\text{C}/^{12}\text{C}$ ratio in the silicate Earth, and yields 1 wt.% carbon in the core⁹. In addition, Wood *et al.*⁹ demonstrated that carbon strongly affects the chemical activity of Mo and W in liquid metal, so that their abundance in the mantle can be explained by partitioning between silicate melt and core-forming metal with ~ 0.6 wt.% carbon. It has been repeatedly suggested that the inner core may be composed of Fe_7C_3 , which accounts

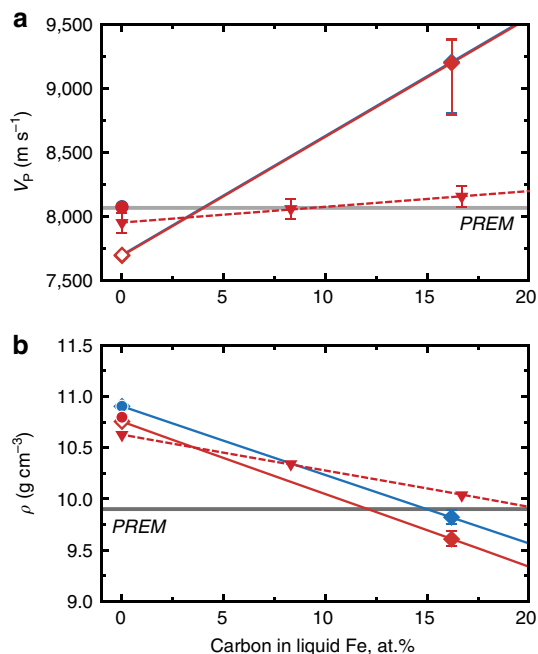


Figure 8 | Effect of carbon on the velocity and density of liquid Fe at 136 GPa. (a) Velocity and (b) density for liquid $\text{Fe}_{84}\text{C}_{16}$ at $T_{\text{CMB}} = 4,300$ K (red) and 3,600 K (blue). Present results (closed diamonds, solid curves) are compared with theoretical calculations³⁴ (triangles, broken curve). The data for pure Fe are from shock compression study⁸ (open diamonds) and theoretical calculations²⁶ (closed circles). PREM denotes seismological observations³⁵ at the CMB.

for high Poisson's ratio observed^{14,15}. The crystallization of solid Fe_7C_3 from a liquid outer core with < 1.2 wt.% carbon may still be possible if sulfur is also included in the core³⁸.

Methods

High P - T generation. Molten Fe-C alloy was obtained at high P - T in an external-resistance-heated (EH) or laser-heated (LH) diamond-anvil cell (DAC; Supplementary Table 1) using facilities installed at SPring-8. A disc of pre-synthesized $\text{Fe}_{84}\text{C}_{16}$ sample, 20–25 μm thick and 100–120 μm in diameter, was loaded into a hole of a rhenium gasket, together with two 12–17 μm thick single-crystal Al_2O_3 sapphire discs that served as both thermal and chemical insulators. The sample was compressed with 300 μm culet diamond anvils to a pressure of interest before heating.

In LH-DAC experiments, the sample was heated at high pressure from both sides by using two 100 W single-mode Yb fibre lasers (YLR-100-AC, IPG Photonics Corp.). The Gaussian-type energy distribution of the laser beam was converted into flat-top one with a refractive beam shaper (GBS-NIR-H3, Newport Corp.). A typical laser spot was 50–70 μm in diameter on the sample, much larger than X-ray beam size (~ 17 μm). We determined temperature by a spectroradiometric method, and its variations within the area irradiated by X-rays and fluctuations during IXS measurements were $< \pm 10\%$. The pressure was obtained from the equation of state for Fe_3C (ref. 39) from the lattice constant observed before melting at 1,800–2,500 K. Its error was derived from uncertainties in both temperature and the volume of Fe_3C . A typical image of a sample recovered after the laser heating experiment at 70 GPa and 2,700 K is given in Supplementary Fig. 4.

Only run #FeC08 was conducted in an EH-DAC. The whole sample was homogeneously heated by a platinum-resistance heater placed around the diamonds. The temperature was obtained with a Pt-Rh (type-R) thermocouple whose junction was in contact with the diamond near a sample chamber. The temperature uncertainty was < 20 K. We determined the pressure based on the Raman shift of a diamond anvil⁴⁰ before heating at 300 K, whose uncertainty may be as much as $\pm 20\%$.

IXS measurements. The sound velocity of liquid Fe-C alloy was determined in the DAC by high-resolution IXS spectroscopy at the beamline BL35XU, SPring-8 (ref. 41). Both LH- and EH-DACs were placed into vacuum chambers to minimize background scattering by air. The measurements were carried out with ~ 2.8 meV energy resolution using Si (999) backscattering geometry at 17.79 keV. The

experimental energy resolutions were determined using scattering from Polymethyl-methacrylate. The incident X-ray beam was focused to about 17 μm size (full width at half maximum) in both horizontal and vertical directions by using Kirkpatrick-Baez mirrors⁴². The X-ray beam size was much smaller than heated area (50–70 μm for LH-DAC). Scattered photons were collected by an array of 12 spherical Si analyzers leading to 12 independent spectra at momentum transfers (Q) between 3.2 and 6.6 nm^{-1} with a resolution $\Delta Q \sim 0.45$ nm^{-1} (full width) that was set by slits in front of the analyzer array. The energy transfer range of ± 30 (or -10 to ± 30) meV was scanned for 1–3 h. Before and after IXS data collections, sample melting was confirmed by X-ray diffraction data (Fig. 2) that was collected, *in situ*, by switching a detector to a flat panel area detector (C9732DK, Hamamatsu Photonics K.K.)⁴³.

The IXS spectra included three (sometimes five) peaks (Fig. 1) of Stokes and anti-Stokes components of the longitudinal acoustic (LA) phonon mode from the sample (sometimes also from a diamond), and a quasi-elastic contribution near zero energy transfer. These spectra were fitted with the damped harmonic oscillator (DHO) mode⁴⁴ for acoustic phonon modes and with Lorentzian function for quasi-elastic peaks convolved by experimental resolution function. The DHO model function can be described as;

$$S^{\text{DHO}}(Q, \omega) = \left[\frac{1}{1 - e^{-\hbar\omega/k_B T}} \right] \frac{A_Q}{\pi} \frac{4\omega\omega_Q \Gamma_Q}{(\omega^2 - \Omega_Q^2)^2 + 4\Gamma_Q^2 \omega^2}, \quad (3)$$

where A_Q , Γ_Q , Ω_Q , k_B and \hbar are the amplitude, width, and energy of inelastic modes, Boltzmann constant and Planck constant, respectively. In the fitting, temperature T was fixed at a sample temperature obtained by a spectroradiometric method or a thermocouple. The excitation energy modes appearing at both Stokes and anti-Stokes sides correspond to the phonon creation and annihilation, respectively. With increasing temperature, as given by the Bose function in equation (3), the intensities of such Stokes and anti-Stokes peaks become similar to each other. A symmetric shape of the present IXS spectra therefore assures that the IXS signals originated from a high-temperature area.

The peak at a finite energy transfer gives the frequency of each mode (Fig. 1). The excitation energies for the LA phonon mode of liquid $\text{Fe}_{84}\text{C}_{16}$ obtained in a pressure range of 7.6–70 GPa are plotted as a function of momentum transfer (Q) in Fig. 4. The compressional sound wave or P-wave velocity (V_P) corresponds to the long-wavelength LA velocity at $Q \rightarrow 0$ limits;

$$V_P = \left(\frac{dE}{dQ} \right)_{Q \rightarrow 0}. \quad (4)$$

We made a linear fit to the data obtained at low Q below 3.5 nm^{-1} to determine the P-wave velocity (Supplementary Table 1), because positive dispersion can appear at higher $Q \gg 3$ nm^{-1} (ref. 45). For comparison, the results based on a sine-curve fit to all Q -range data, as is usually applied for polycrystalline samples in similar high-pressure IXS measurements⁴⁶, are also given in Supplementary Table 1. In general, the error bars of the two determinations of V_P overlap, though the sine fit to large Q does give slightly larger V_P , as would qualitatively be expected from previous measurements on liquid iron⁴⁷.

Equation of state for liquid $\text{Fe}_{84}\text{C}_{16}$. We constructed an equation of state (EoS) for liquid $\text{Fe}_{84}\text{C}_{16}$ to extrapolate the present V_P data and to estimate its density at the core pressure range. V_P of liquid can be written as;

$$V_P = \sqrt{\frac{K'_S}{\rho}}. \quad (5)$$

The pressure dependence of K'_S is assumed to be

$$K'_S = K'_{S0} + K''_S P, \quad (6)$$

where K'_S is the pressure derivative of K_S and subscript zero indicates a value at 1 bar. The adiabatic Murnaghan EoS can be described as (for example, ref. 4);

$$\rho = \rho_0 \left(1 + \frac{K''_S P}{K'_{S0}} \right)^{\frac{1}{K'_S}}. \quad (7)$$

Equation (5) is thus rewritten as;

$$V_P = \sqrt{\frac{K'_{S0}}{\rho_0}} \left(1 + \frac{K''_S P}{K'_{S0}} \right)^{\frac{1}{2} - \frac{1}{2K'_S}}. \quad (8)$$

The temperature effect on ρ_0 can be expressed by;

$$\rho_0(T) = \rho_0(T_0) / \exp\left(\int_{T_0}^T \alpha dT\right). \quad (9)$$

The thermal expansion coefficient α is also dependent on temperature as;

$$\alpha(T) = a + bT, \quad (10)$$

where a and b are constants. Previous density measurements¹⁶ of liquid Fe-C alloys at 1 bar give $a = 6.424 \times 10^{-5} \text{K}^{-1}$ and $b = 0.606 \times 10^{-8} \text{K}^{-2}$ for liquid

$\text{Fe}_{84}\text{C}_{16}$ using $\rho_0 = 6.505 \text{ g cm}^{-3}$ at $T_0 = 2,500 \text{ K}$ as a reference. The result of fitting equation (8) to the present $P - V_p$ data is given in Fig. 3.

Isothermal bulk modulus. We estimate isothermal bulk modulus K_T from isentropic bulk modulus K_S in two ways. The relationship between these two is described as follows;

$$\frac{K_S}{K_T} = \frac{C_P}{C_V} = 1 + \alpha\gamma, \quad (11)$$

where C_P and C_V are heat capacities at constant pressure and volume, respectively. Although γ for liquid Fe-C alloys is not known, $\gamma_0 = 1.74$ has been reported for liquid Fe at 1 bar and 1,811 K (ref. 8) It is close to 1.58 for liquid $\text{Fe}_{90}\text{O}_8\text{S}_2$ estimated from the shock compression data set⁴⁸.

Extrapolation of present data to core pressures. With the EoSs determined above (equations (7) and (8)), we extrapolate the P-wave velocity and density of liquid $\text{Fe}_{84}\text{C}_{16}$ to the core pressure range along adiabatic compression, in which temperature is given by;

$$T = T_0 \exp \left[\int_{\rho_0}^{\rho} (\gamma/\rho) d\rho \right]. \quad (12)$$

Assuming $\gamma = \gamma_0 \times (\rho_0/\rho)$, temperature is simply represented as;

$$T = T_0 \exp \left[\gamma_0 \left(1 - \frac{\rho_0}{\rho} \right) \right]. \quad (13)$$

γ_0 is fixed at 1.74 previously obtained for liquid Fe (ref. 8). Using the temperature dependence of K_{S0} and ρ_0 shown above, we calculate density, velocity and temperature profiles along adiabatic compression with various reference temperatures at the CMB. The adiabatic compression profiles of liquid $\text{Fe}_{84}\text{C}_{16}$ for the low ($T_0 = 2,045 \text{ K}$ and $T_{\text{CMB}} = 3,600 \text{ K}$)³² and high ($T_0 = 2,457 \text{ K}$ and $T_{\text{CMB}} = 4,300 \text{ K}$)³³ temperature cases are calculated in Fig. 7.

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Author contributions

Y.N. synthesized a starting material and performed experiments and data analysis. Y.N., S.I., K.H., T.K., S. Tateno, S. Tsutsui, Y.K. and A.B. were involved in IXS measurements. H.O., S. Tateno, S.I. and Y.N. were involved in developing the laser heating system at the beamline. Y.N., K.H. and A.B. wrote the paper. All authors discussed the results and commented on the manuscript.

Additional information

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