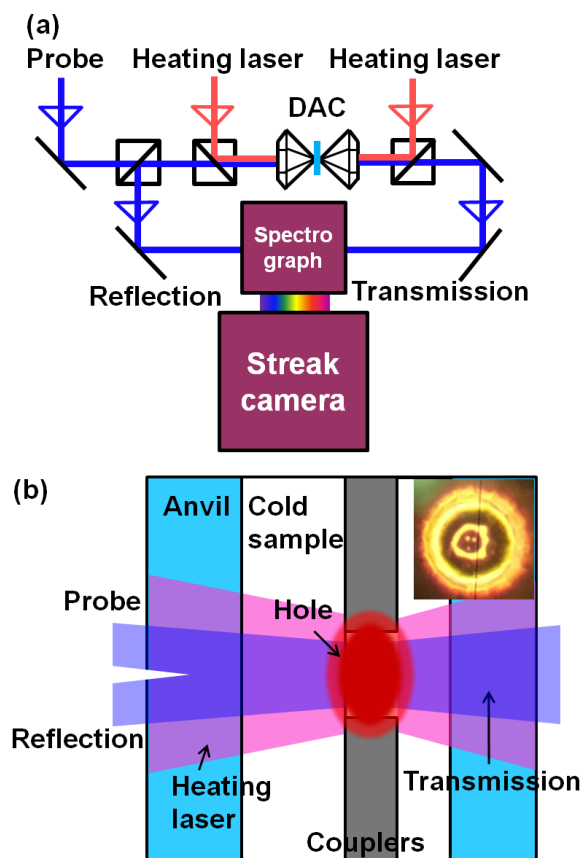
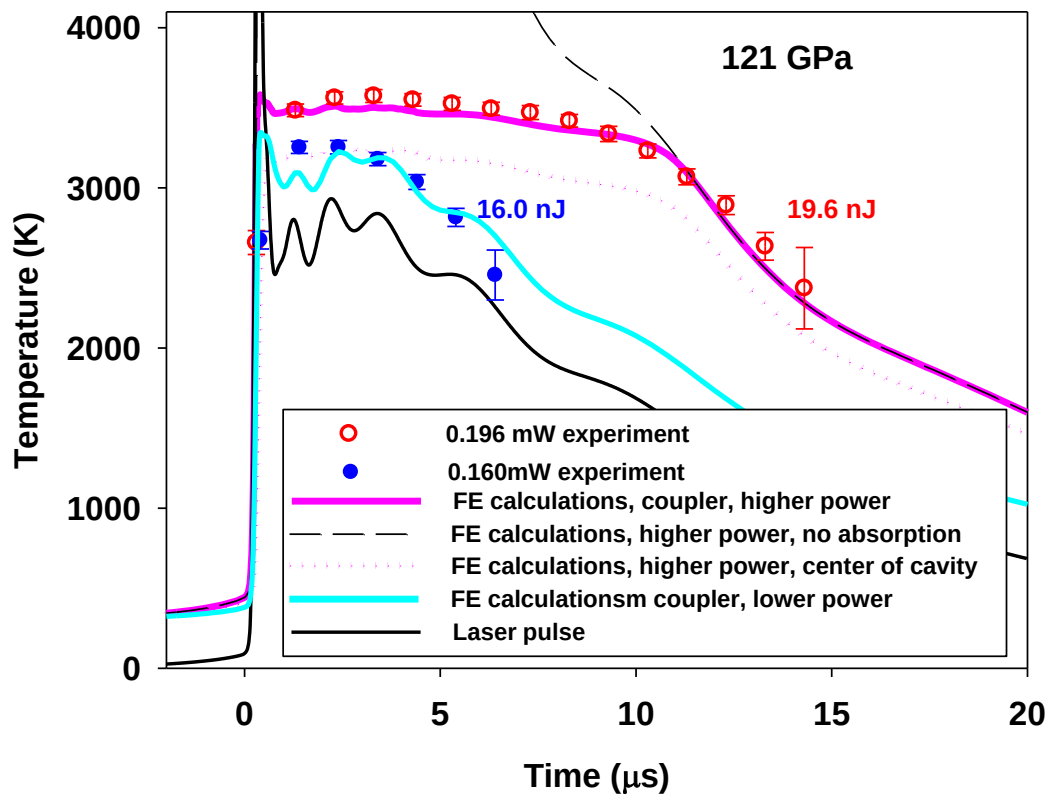


Supplementary Information for Metallization and molecular dissociation of dense fluid nitrogen

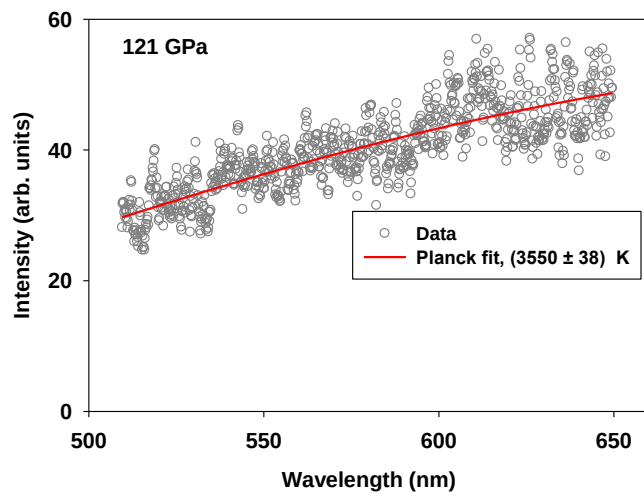
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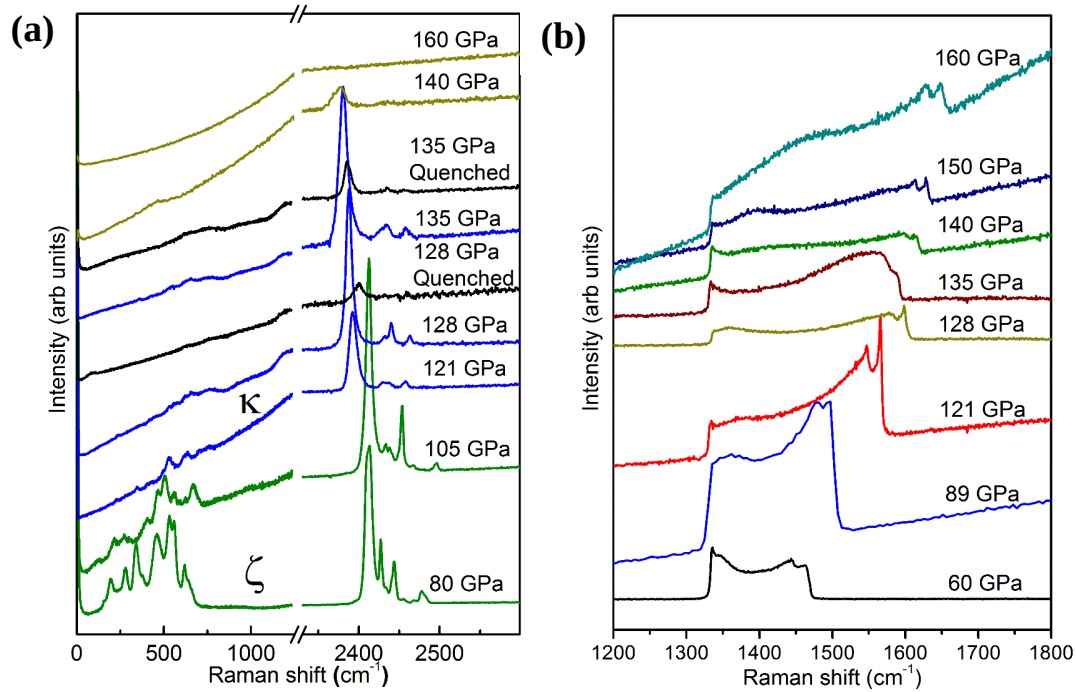
Supplementary Figure 1. Configuration of a single shot laser heating experiment of nitrogen to probe transient transmission and reflection in the diamond anvil cell. (a) Optical schematic: time resolved reflection or transmission can be measured one at a time superimposed with the thermal emission- see our previous publications ^{1,2} for the timing of acquisition of the optical probes and thermal emission using a streak camera ; (b) Microscopic view of the diamond cell cavity, which contained the nitrogen sample and a metal Ir foil (coupler) which had a small hole(s) (5~10 μm in diameter); coupler is heated by an IR laser from both sides and it converts laser radiation to heat, which is transferred to the adjacent sample inside the hole (if the laser heats the rim of the hole) or from both sides of the coupler (if a flat regular surface is exposed). The transmission probe is passing through the nitrogen sample inside the hole in a coupler, and the reflection probe is detected from the same side as the incident probe beam.



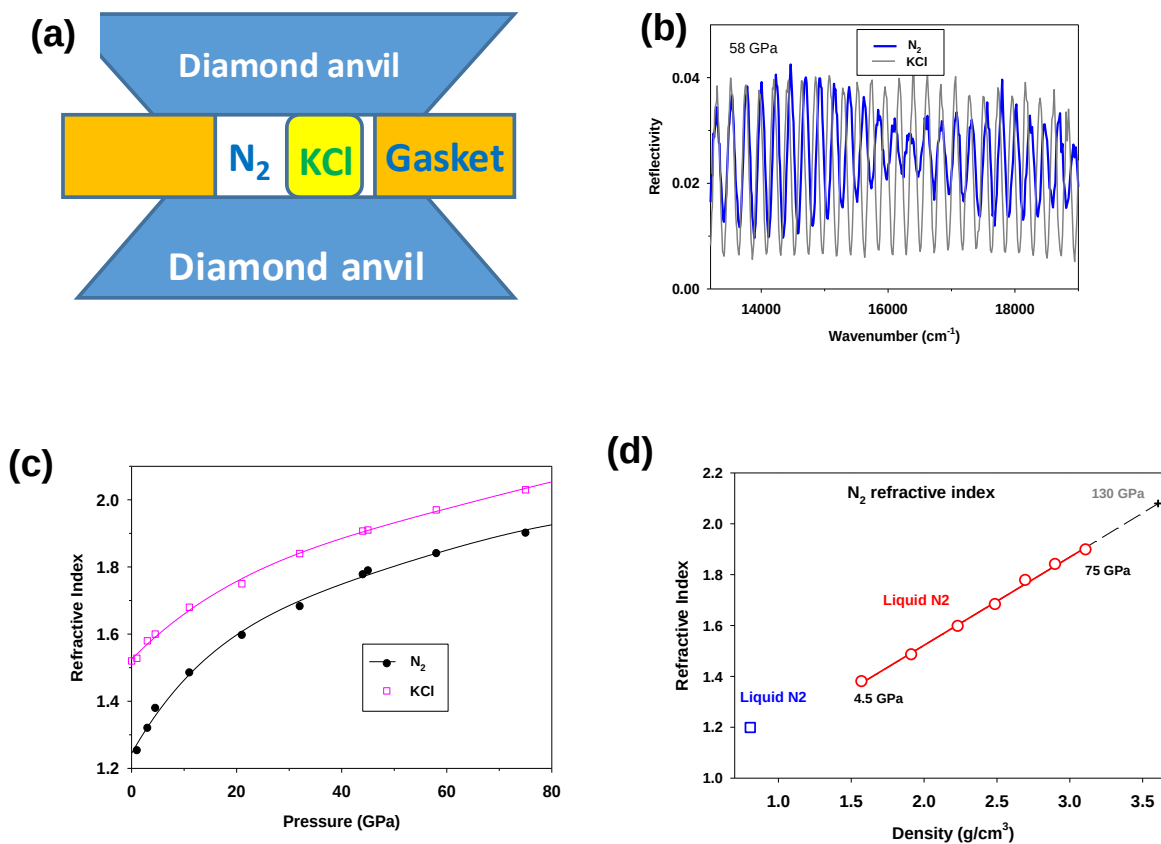
Supplementary Figure 2. Finite element (FE) calculations of the temperature histories in nitrogen at 121 GPa in comparison with the experimentally determined dependencies. Experiments show a plateau, which becomes longer with the increase in heating laser pulse energy (tabulated and measured through the pulsed laser energy). FE calculations picture well this observations. If no absorption is introduced, FE calculations show much higher temperatures instead of plateau approximately corresponding to the onset of the sample absorption. The laser pulse intensity is shown in arbitrary units.



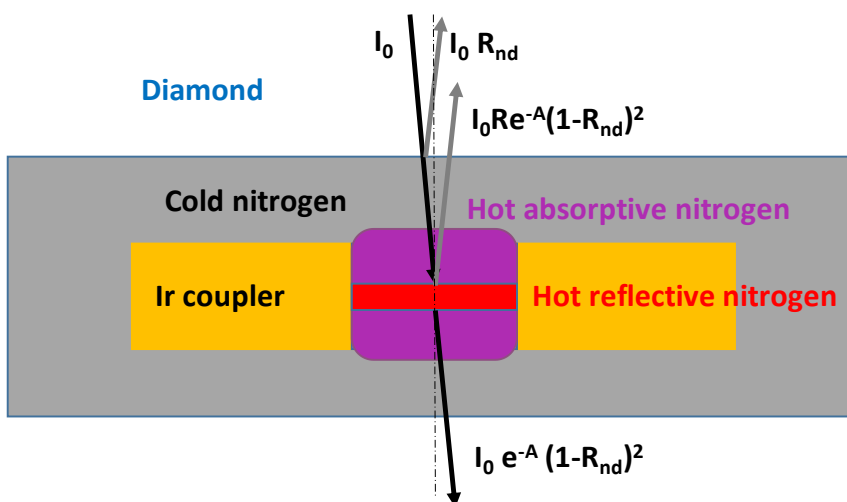
Supplementary Figure 3. The thermal radiation spectrum near the maximum (at the $2 \mu\text{s}$, Fig. 1 (a), the bottom panel) used to determine the sample temperature.



Supplementary Figure 4. The Raman spectra of nitrogen at various pressures before and after laser heating experiments (a) and stressed diamond anvils (b). The pressure was determined using the spectral positions of the N-N stretching mode of N₂ (vibron)³ (a) and stressed diamond⁴ (b) measured by Raman spectroscopy. At above 121 GPa N₂ is in the κ molecular phase⁵. The black lines show the Raman spectra after laser heating at 128 and 135 GPa, which demonstrate a gradual transformation into amorphous η state⁶ but no sign of cg or LP nonmolecular nitrogen^{7,8}.



Supplementary Figure 5. Refractive index of nitrogen to 75 GPa (a) Experimental arrangement: a thin slab of KCl is squeezed between diamond anvils and N₂ serves as a pressure medium, reflectivity spectra are measured from the KCl and N₂ regions yielding a series of interference fringes due to interference from the two sample-diamond interfaces, the refractive index of N₂ is determined at each pressure point by referencing to the refractive index of KCl, which is determined by a linear extrapolation as a function of density of measurements of Ref. ⁹ using the equation of state of Ref. ¹⁰; (b) Examples of optical interference at high pressures; (c) Results up to 75 GPa as a function of pressure; (d) results extrapolated to 130 GPa using a linear dependence on the density; the equation of state of Ref. ⁵ is used.



Supplementary Figure 6. A schematic of absorption and reflectivity measurements of liquid conducting nitrogen in the DAC. Hot conducting nitrogen is created in a hole of a metallic Ir coupler. The transmission spectra determine the absorbance A . Once nitrogen transforms to a strongly reflective state in the hottest area of the cell, a phase boundary appears between the metallic and semiconducting (absorptive) states. To extract the reflectance spectra we normalize for the optical absorption in semiconducting nitrogen.

Supplementary references

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