

First Measurements of Deuterium–Tritium and Deuterium–Deuterium Fusion-Reaction Yields in Ignition-Scalable Direct-Drive Implosions

In direct-drive inertial confinement fusion (ICF) ignition designs, a cryogenic deuterium–tritium (DT) shell surrounding a vapor and encased in a thin ablator ($<10 \mu\text{m}$) is symmetrically heated with nominally identical laser beams. In most designs, laser ablation launches single or multiple shocks through the converging shell and into the vapor region. The shock-transit stage of the implosion is followed by a deceleration phase, where the kinetic energy of the converging shell is converted to the internal energy of the hot spot. Thermonuclear fusion reactions are initiated in both the shock phase and the compression phase once sufficiently high temperatures and densities are reached. To achieve conditions relevant for ignition implosion designs, the hot-spot size must exceed the mean free path of fusing ions and the mean free path of the alpha particles.

Previous experiments on OMEGA have reported anomalous $Y_{\text{DT}}/Y_{\text{DD}}$ values (different by as much as a factor of 4) with the measured pre-shot fuel composition and experimentally inferred ion temperatures in room-temperature implosions.¹ Several studies suggest that species separation of the hydrogen isotope resulting from multifluid effects^{2,3} is likely responsible for the observed discrepancies in the yield ratios. These classes of implosions—for example, exploding pushers that use thin glass ($\sim 3\text{-}\mu\text{m}$ SiO_2) or thin CH ($<16\text{-}\mu\text{m}$) shells—are, however, characterized by fusion reactions that occur predominantly during the shock phase at very high temperatures (≥ 10 keV)

and relatively low densities ($\leq 10 \text{ mg/cm}^3$). The mean free path for 90° deflection is given by $\lambda_{\text{ii}} \sim T_{\text{i}}^2 / Z_{\text{i}}^2 Z^2 \rho$ (Ref. 4) for ions of charge Z_{i} , average ion temperature T_{i} , ion charge Z , and density ρ . Conditions during the shock phase result in large mean-free-path lengths of the ions relative to the size of the fusing-plasma region (see Table 149.II). These conditions are also typical of ignition-relevant direct-drive cryogenic implosions⁵ during the shock phase; however, cryogenic targets differ from exploding-pusher targets in two respects: First, most of the neutron yield in a cryogenic implosion occurs later in the implosion, during the compression phase, when the kinetic energy is converted to the internal energy of the hot spot. Simulations using the spherically symmetric hydrodynamics code *LILAC*⁶ indicate that nearly 99% of the yield occurs in this compression phase. Second, compression yields occur at significantly higher densities ($\geq 20 \text{ g/cm}^3$) and lower temperatures (~ 3 keV), leading to mean free paths of thermal ions that are much shorter than the hot-spot size. Nonlocal transport of energetic ions is therefore not expected to significantly influence yields during compression. Evidence of fuel species separation that persists into the compression phase would suggest a reduction in the number of alpha particles produced from the dominant D–T fusion reactions. In ignition-scalable cryogenic implosions described in this article, however, measurements give the first evidence that species separation does not persist from the shock phase and has an insignificant influence on

Table 149.II: Calculated implosion parameters for various plasma conditions ranging from a highly kinetic exploding pusher (in the shock phase in the vapor) to a strongly hydrodynamic-like plasma regime (cold-fuel layer in the shock or compression phase).

Implosion Type	ρ (g/cm^3)	T_{i} (keV)	λ_{ii} (μm)	R_{shell} (μm)
Exploding pusher:				
Shock phase	0.03	10	400	100
Cryogenic implosions:				
Shock phase: vapor	0.1	8	80	100
Shock phase: cold-fuel layer	6.0	0.02	0.0002	$\Delta R_{\text{shell}} \sim 10$
Compression phase	20.0	3	0.08	25

the yield ratio into the compression phase in direct-drive D–T cryogenic implosions consisting of a near-equimolar mixture of deuterium and tritium.⁷

Direct-drive ICF targets consisting of a deuterated plastic (ablator) shell with a 460- μm outer radius are imploded at an ignition-scalable, on-target laser intensity with a laser energy of ~ 25 kJ (Ref. 8). The implosion velocity (V_{imp} , defined as the velocity of the compressing shell when the kinetic energy of the shell is at a maximum) ranged from 3.5×10^7 cm/s to 4×10^7 cm/s and the adiabat (α , defined as the ratio of the pressure to the Thomas–Fermi pressure at maximum shell density) ranged from 2.4 to 5. The average ion temperature T_i in this class of implosions is varied by adjusting the implosion velocity, $T_i \sim V_{\text{imp}}^{1.1}$, which, in turn, is governed by the thickness of the cryogenic DT layers or the CH (CD) ablator. The capsule is filled by a permeation technique at a temperature of 300 K, where increasing pressure is applied to the outside of the shell, allowing the gas to diffuse inside. Fill rates for a typical cryogenic target are carefully controlled by holding the pressure ramp rate at ~ 1 atm/min to ensure the integrity of the shell is not compromised.⁹ At the final fill pressure (between 400 and 800 atm) depending on the desired ice thickness, the capsule is cooled to a few mK below the triple point (~ 19.8 K), producing a DT ice layer ranging from 40 to 90 μm in thickness. The primary nuclear-fusion reactions examined in this study are given by



The neutron yields are measured using the time-of-flight (nTOF) diagnostics positioned around the OMEGA target chamber. The fusion yield is given by

$$Y_n^{\text{DD/DT}} = \int f_{\text{D}} f_{\text{D/T}} [\rho(\vec{r}, t)]^2 \langle \sigma v \rangle_{\text{DT/DD}} \times dr^3 dt / (1 + \delta_{\text{DD}}) \bar{m}^2,$$

where f_{T} and f_{D} are the atomic fractions of the reactants, ρ is the fuel-mass density, $\langle \sigma v \rangle$ is the Maxwellian-averaged reactivity for the D–T or D–D fusion reaction (which scales as $\sim T_i^{3.7}$ for the D–T reaction and $\sim T_i^{3.3}$ for the D–D reaction for the typical temperatures in OMEGA implosions), T_i is the average ion temperature, \bar{m} is the average reactant mass, and $\delta_{\text{DD}} = 1$ for DD and 0 for DT to account for double counting of the identical D–D reaction.

The primary D–T yields observed in cryogenic experiments are always lower relative to radiation–hydrodynamics codes that assume spherical symmetry and include the deposition of the laser energy through collisional absorption and account for laser–plasma interactions such as cross-beam energy transfer (CBET).¹⁰ These codes include nonlocal heat conduction¹⁰ and multigroup diffusive radiative transport.¹¹ Several multi-dimensional effects that reduce the overall yield relative to these state-of-the-art spherically symmetric fluid codes have been proposed, including nonuniformity growth caused by beam-to-beam energy imbalance,¹² on-target beam misalignment,¹³ single-laser-beam nonuniformity,¹³ and isolated defects on the target¹⁴ that potentially reduce T_i and/or fuel density. All these mechanisms include only hydrodynamic effects and do not exhibit yield ratio anomalies. More recently, an extension to fluid codes has been proposed. Calculations that include plasma barotropic diffusion,^{15,16} where hydrogen isotope species separation occurs during the shock phase into the hot spot because gradients in pressure and temperature, have been shown to influence the D–T and D–D fusion yields differently. Two phases of an ICF implosion have been analyzed using this model: the shock phase (when the shock is moving through the vapor toward the center of the capsule) followed by the rebound phase (outward-going shock). It was reported that during the shock phase, up to 5% of the deuterium can leave the fuel volume for an equimolar mixture of deuterium and tritium. During the subsequent shock-rebound phase, the barotropic diffusion rate decreases to zero and the ability for fuel to leave the volume is significantly reduced if not eliminated. Since the D–D fusion and D–T fusion reactivity are well-known¹⁷ and the composition of the fuel is measured prior to the implosion, the ratio of the neutron yields ($Y_{\text{DT}}/Y_{\text{DD}}$) from these reactions should follow a calculable trend with the measured ion temperature with the exclusion of diffusive effects. Table 149.II summarizes the mass-fuel density (ρ) and the key implosion parameters to calculate the ion–ion mean free path (λ_{ii}) for the plasma conditions across the class of implosions discussed earlier in this article. The radius of the shell (R_{shell}) is calculated from simulations for the different phases of the implosion.

As shown in Table 149.II, the mean free path during the shock phase for the ions at the relevant average ion temperature approaches the radius of the shell. At this time, however, the vapor region is surrounded by a relatively cold (~ 20 -eV) and highly dense DT-fuel layer. The energetic and thermal ions that escape the vapor phase do not leave the target and instead are stopped in the cold dense DT shell. At peak neutron production, the mean free path is several orders of magnitude smaller ($\sim 10^{-2}$) than the boundary of the cold-fuel shell.

Cryogenic implosions are additionally different from shock-driven implosions that have been studied previously since the shell material is also made of DT fuel. When the shell decelerates in the compression stage of any ICF implosion, the cold fuel ablates into the hot spot. Simulations using the code *LILAC* indicate that, in the case of cryogenic layered DT implosions, nearly $5\times$ the mass of the original vapor⁶ is injected into the hot spot through the ablation process, which is the primary source of the fusion neutrons during compression. Therefore, it would be expected that the ions that are stopped in the cold-fuel shell would be restored into the hot core during the compression phase, compensating for any loss of particles that may have occurred earlier in the implosion.

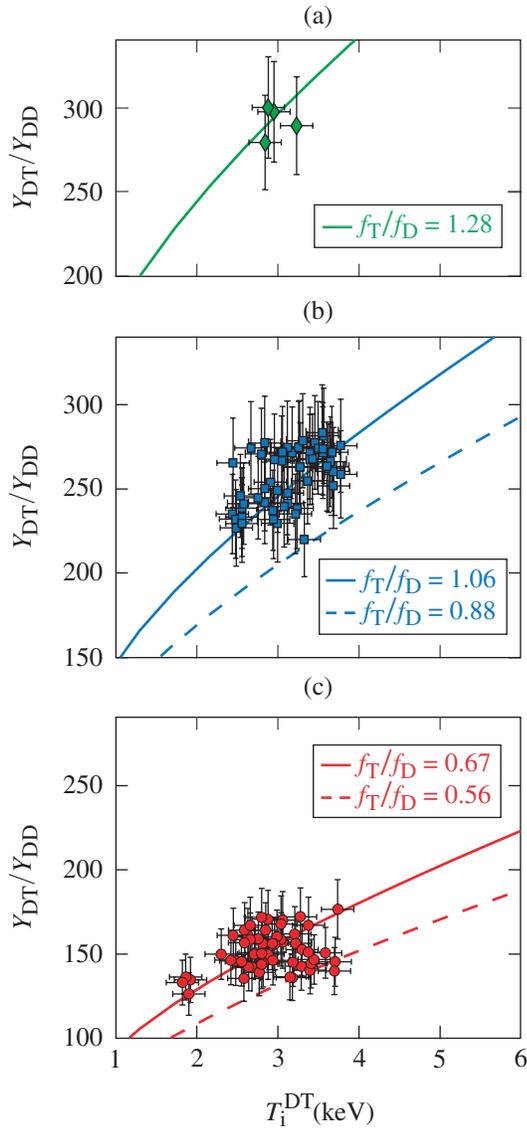
For this analysis, the yields (Y_{DT} and Y_{DD}) for the different reactions are measured along the same diagnostic line of sight using the 13.4-m high-resolution nTOF spectrometer.¹⁸ This diagnostic uses several microchannel-plate–based phototubes to increase the dynamic range required to measure the primary DT and DD signal in a single line of sight. The yield is inferred by fitting the recorded signal with a forward-fit approach using a relativistic model of the neutron distribution.¹⁹ Cross-calibration of the neutron diagnostics with standard measurements on OMEGA give an uncertainty in the D–T and D–D yields of 5% and 9%, respectively.^{20,21} In ignition-scalable implosions, the neutron yield is attenuated by the compressed fuel at peak neutron production (see Table 149.II). To recover the fusion birth yield, a correction to the measured yields must be included as a function of the areal density from the compressed fuel. The elastic scattering is proportional to the areal density of the implosion, which is inferred from separate measurements.²² The transmission factors (η_{DT} and η_{DD}) for the neutrons from the two fusion reactions are calculated using the well-known total scattering cross sections and the measured areal density. Typical values of these transmission factors for an areal density of 220 mg/cm^2 are 4% and 10% for the DT and DD neutrons, respectively. With the areal densities achieved on OMEGA, multiple scattering can be neglected, thereby providing an ideal platform to study the effects of fuel-species separation in ignition-scalable implosions. By adding the uncertainty of the D–T and D–D yields, the attenuation of the yield from the compressed fuel and the reaction rate for both of the primary reactions in quadrature, an error of 10% for the Y_{DT}/Y_{DD} ratio can be inferred.

As indicated earlier, it is important to know the ion temperature in the implosion and the fuel composition. The energy spread of the primary neutron distribution provides a good measure of the ion temperature characteristics of peak neutron

production. If mass flow within the reaction region is present, this effect can lead to a broadening of peak distribution and an incorrect interpretation of ion temperature.²³ On OMEGA, several nTOF detectors measure the width of the DT neutron spectrum temperature from various lines of sight around the target chamber.²⁴ The ion temperature inferred from the width of the neutron spectrum in ignition-scalable implosions can vary up to ~ 1 keV across the three different detectors. Simulations indicate that this variation in the temperature is caused by bulk fluid motion of the fusing plasma.²⁵ The uncertainty in the inferred ion temperature, excluding effects caused by bulk fluid motion, is ± 0.2 keV for implosions between 2 keV and 5 keV. To minimize the effect of bulk motion, the minimum ion temperature will be used in this analysis as an approximation of the thermal temperature. It should be noted that the implosions that can vary up to 1 keV are only 3% of the data points. The remaining 2σ that vary up to 0.8 keV account for 90% of the implosion analyzed. Using this variation in the ion temperature, the calculated fuel fraction has an uncertainty of less than 7%.

The observed reaction yield ratio is plotted as a function of the minimum ion temperature in Fig. 149.24 for each cryogenic shot on OMEGA (35 experimental campaigns with 120 implosions taken over a period of three years). The composition of the DT inventory in the assay volume is periodically measured on OMEGA to within an accuracy of 1.5%. In this case, the gas used to fill the targets was taken at various stages during the pressurization of the fuel so that the deuterium-to-tritium (D:T) concentration could be calculated. Over time, the tritium supply in the system gradually changes as a result of beta decay of the hydrogen isotope. Figure 149.24 also shows the calculated ratios using the measured fuel fraction and the minimum ion temperature. The measured ratios show good agreement with the calculated ratios expected from the DT inventory and experimentally inferred ion temperatures. It should be noted that while the accuracy of the fuel composition in the both the assay volume and the pressurized system are well understood, an extrapolation of the fuel fraction is required of the gas composition during the fill process in the permeation cell that is used to fill cryogenic capsules. A project is underway to better characterize the fuel composition of the gas as it is sent into the permeation cell used to fill the capsules. Presently, this effect is known to change the composition between 3% and 5%. The calculated reaction yield ratios follow the form $Y_{DT}/Y_{DD} \sim 2T_i^{0.4} (f_T/f_D)$ using the assumption that hydrodynamic models of an ICF implosion predict that the reactant density ratio (f_T/f_D) is spatially and temporally constant during all phases. This indicates that additional effects that change this ratio or the volume over

which each of the D–T and D–D reactions are produced do not significantly influence yields from the hot-spot stagnation. Pre-shot fuel fractions are measured during each fill process for every campaign. Variations in the yield ratio measurements resulting from the fuel composition are reflected in Fig. 149.24 with the solid and dashed lines representing the initial and final

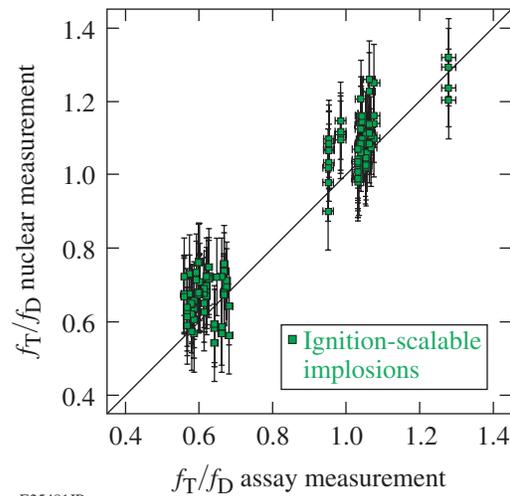


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Figure 149.24 With each cryogenic implosion, the Y_{DT}/Y_{DD} yield ratio is plotted with the minimum DT neutron averaged ion temperature. The T:D fuel fraction used to fill the cryogenic targets varied between (a) $f_T/f_D \sim 0.58$ and (b) $f_T/f_D \sim 1$ for the majority of the cryogenic targets. (c) A single campaign had a fill fraction of $f_T/f_D \sim 1.28$. The solid lines represent the initial measurement of fuel inventory and the dashed lines show how much the fuel has changed over time resulting from tritium decay. The hydrogen concentration does not contribute to the fusion yield and is not included in this analysis.

measurement, respectively, before the inventory underwent a scheduled refinement.

The measured D–T and D–D yield ratios and the ion temperature are used to instead infer a fuel fraction (f_D and f_T) for each of these shots. The measured fuel fraction is compared against values inferred from nuclear measurements in Fig. 149.25. The average of the ratio of the inferred fuel fraction from the nuclear measurement over the composition obtained from the permeation cell is 1.07 with a standard deviation of 0.09. Although error on the mean is small with 1% for 120 implosions used for this study, given the 10% systematic error on the Y_{DT}/Y_{DD} ratio, both measurements of the fuel fractions are consistent within the experimental uncertainties.



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Figure 149.25 The measured fuel fraction determined at the fill station (assay measurement) is compared with the values inferred from the nuclear measurements over a three-year period. The changes in the fuel fraction used to fill the targets is clearly visible between $f_D/f_D \sim 0.58$ and $f_D/f_D \sim 1$. A single campaign with four implosions had a fill fraction of $f_T/f_D \sim 1.28$.

In summary, nuclear measurements of the D–T to D–D yield ratio from OMEGA cryogenic implosions scale predictably with the known composition of the fuel and experimentally inferred ion temperatures with a calculated 7% systematic offset. These observations indicate that multifluid effects that may take place during the shock phase of the implosion (and potentially influence species profiles in the compressing target) do not persist into the subsequent compression phase of the implosion. A plausible explanation for this rests on the composition of the target; the shell is also DT fuel. During the deceleration phase of cryogenic DT implosions, the fuel from the inner DT wall is ablated into the hot spot. Simula-

tions indicate that nearly $5\times$ the mass of the neutron-emitting region is from the ablation of the cold DT shell. Therefore, the energetic ions that may be lost because of their long mean free paths earlier in the implosion return to the hot spot during peak neutron production, leading to an unchanged fusion yield ratio. These observations indicate that multifluid effects have an insignificant influence on the yield ratio in ignition-scalable cryogenic implosions.

The largest contribution to the uncertainty in the yield ratio measurement is caused by the D–D yield. Upcoming experiments are designed to increase the accuracy of this measurement to 5%. These experiments will reduce the uncertainty in the Y_{DT}/Y_{DD} ratio to 7%, which, in turn, will also increase the accuracy of the inferred fuel fractions obtained from this measurement.

Presently, there is no measurement available of the true temperature of the plasma, which is very important for this measurement. Several projects are being considered that will provide a true thermal temperature that is not influenced by the bulk motion of the plasma.

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REFERENCES

1. D. T. Casey, J. A. Frenje, M. Gatu Johnson, M. J.-E. Manuel, H. G. Rinderknecht, N. Sinenian, F. H. Séguin, C. K. Li, R. D. Petrasso, P. B. Radha, J. A. Delettrez, V. Yu. Glebov, D. D. Meyerhofer, T. C. Sangster, D. P. McNabb, P. A. Amendt, R. N. Boyd, J. R. Rygg, H. W. Herrmann, Y. H. Kim, and A. D. Bacher, *Phys. Rev. Lett.* **108**, 075002 (2012).
2. M. J. Rosenberg, A. B. Zylstra, F. H. Séguin, H. G. Rinderknecht, J. A. Frenje, M. Gatu Johnson, H. Sio, C. J. Waugh, N. Sinenian, C. K. Li, R. D. Petrasso, P. W. McKenty, M. Hohenberger, P. B. Radha, J. A. Delettrez, V. Yu. Glebov, R. Betti, V. N. Goncharov, J. P. Knauer, T. C. Sangster, S. LePape, A. J. Mackinnon, J. Pino, J. M. McNaney, J. R. Rygg, P. A. Amendt, C. Bellei, L. R. Benedetti, L. Berzak Hopkins, R. M. Bionta, D. T. Casey, L. Divol, M. J. Edwards, S. Glenn, S. H. Glenzer, D. G. Hicks, J. R. Kimbrough, O. L. Landen, J. D. Lindl, T. Ma, A. MacPhee, N. B. Meezan, J. D. Moody, M. J. Moran, H.-S. Park, B. A. Remington, H. Robey, M. D. Rosen, S. C. Wilks, R. A. Zacharias, H. W. Herrmann, N. M. Hoffman, G. A. Kyrala, R. J. Leeper, R. E. Olson, J. D. Kilkenny, and A. Nikroo, *Phys. Plasmas* **21**, 122712 (2014).
3. H. G. Rinderknecht, M. J. Rosenberg, C. K. Li, N. M. Hoffman, G. Kagan, A. B. Zylstra, H. Sio, J. A. Frenje, M. Gatu Johnson, F. H. Séguin, R. D. Petrasso, P. Amendt, C. Bellei, S. Wilks, J. Delettrez, V. Yu. Glebov, C. Stoeckl, T. C. Sangster, D. D. Meyerhofer, and A. Nikroo, *Phys. Rev. Lett.* **114**, 025001 (2014).
4. D. B. Henderson, *Phys. Rev. Lett.* **33**, 1142 (1974).
5. V. N. Goncharov, T. C. Sangster, R. Betti, T. R. Boehly, M. J. Bonino, T. J. B. Collins, R. S. Craxton, J. A. Delettrez, D. H. Edgell, R. Epstein, R. K. Follet, C. J. Forrest, D. H. Froula, V. Yu. Glebov, D. R. Harding, R. J. Henchen, S. X. Hu, I. V. Igumenshchev, R. Janezic, J. H. Kelly, T. J. Kessler, T. Z. Kosc, S. J. Loucks, J. A. Marozas, F. J. Marshall, A. V. Maximov, R. L. McCrory, P. W. McKenty, D. D. Meyerhofer, D. T. Michel, J. F. Myatt, R. Nora, P. B. Radha, S. P. Regan, W. Seka, W. T. Shmayda, R. W. Short, A. Shvydky, S. Skupsky, C. Stoeckl, B. Yaakobi, J. A. Frenje, M. Gatu-Johnson, R. D. Petrasso, and D. T. Casey, *Phys. Plasmas* **21**, 056315 (2014).
6. J. Delettrez, R. Epstein, M. C. Richardson, P. A. Jaanimagi, and B. L. Henke, *Phys. Rev. A* **36**, 3926 (1987).
7. S. P. Regan, V. N. Goncharov, I. V. Igumenshchev, T. C. Sangster, R. Betti, A. Bose, T. R. Boehly, M. J. Bonino, E. M. Campbell, D. Cao, T. J. B. Collins, R. S. Craxton, A. K. Davis, J. A. Delettrez, D. H. Edgell, R. Epstein, C. J. Forrest, J. A. Frenje, D. H. Froula, M. Gatu Johnson, V. Yu. Glebov, D. R. Harding, M. Hohenberger, S. X. Hu, D. Jacobs-Perkins, R. T. Janezic, M. Karasik, R. L. Keck, J. H. Kelly, T. J. Kessler, J. P. Knauer, T. Z. Kosc, S. J. Loucks, J. A. Marozas, F. J. Marshall, R. L. McCrory, P. W. McKenty, D. D. Meyerhofer, D. T. Michel, J. F. Myatt, S. P. Obenschain, R. D. Petrasso, R. B. Radha, B. Rice, M. Rosenberg, A. J. Schmitt, M. J. Schmitt, W. Seka, W. T. Shmayda, M. J. Shoup III, A. Shvydky, S. Skupsky, S. Solodov, C. Stoeckl, W. Theobald, J. Ulreich, M. D. Wittman, K. M. Woo, B. Yaakobi, and J. D. Zuegel, *Phys. Rev. Lett.* **117**, 025001 (2016); **117**, 059903(E) (2016).
8. C. D. Zhou and R. Betti, *Phys. Plasmas* **14**, 072703 (2007).
9. W. T. Shmayda, M. D. Wittman, R. F. Earley, J. L. Reid, and N. P. Redden, *Fusion Eng. Des.* **109–111**, 128 (2016).
10. I. V. Igumenshchev, W. Seka, D. H. Edgell, D. T. Michel, D. H. Froula, V. N. Goncharov, R. S. Craxton, L. Divol, R. Epstein, R. Follett, J. H. Kelly, T. Z. Kosc, A. V. Maximov, R. L. McCrory, D. D. Meyerhofer, P. Michel, J. F. Myatt, T. C. Sangster, A. Shvydky, S. Skupsky, and C. Stoeckl, *Phys. Plasmas* **19**, 056314 (2012).
11. V. N. Goncharov, O. V. Gotchev, E. Vianello, T. R. Boehly, J. P. Knauer, P. W. McKenty, P. B. Radha, S. P. Regan, T. C. Sangster, S. Skupsky, V. A. Smalyuk, R. Betti, R. L. McCrory, D. D. Meyerhofer, and C. Cherfils-Clérouin, *Phys. Plasmas* **13**, 012702 (2006).
12. I. V. Igumenshchev, V. N. Goncharov, F. J. Marshall, J. P. Knauer, E. M. Campbell, C. J. Forrest, D. H. Froula, V. Yu. Glebov, R. L. McCrory, S. P. Regan, T. C. Sangster, S. Skupsky, and C. Stoeckl, *Phys. Plasmas* **23**, 052702 (2016).
13. S. X. Hu, V. N. Goncharov, P. B. Radha, J. A. Marozas, S. Skupsky, T. R. Boehly, T. C. Sangster, D. D. Meyerhofer, and R. L. McCrory, *Phys. Plasmas* **17**, 102706 (2010).
14. I. V. Igumenshchev, V. N. Goncharov, W. T. Shmayda, D. R. Harding, T. C. Sangster, and D. D. Meyerhofer, *Phys. Plasmas* **20**, 082703 (2013).
15. P. Amendt *et al.*, *Phys. Rev. Lett.* **105**, 115005 (2010).
16. G. Kagan and X.-Z. Tang, *Phys. Lett. A* **378**, 1531 (2014).
17. H.-S. Bosch and G. M. Hale, *Nucl. Fusion* **32**, 611 (1992).

18. C. J. Forrest, V. Yu. Glebov, V. N. Goncharov, J. P. Knauer, P. B. Radha, S. P. Regan, M. H. Romanofsky, T. C. Sangster, M. J. Shoup, and C. Stoeckl, *Rev. Sci. Instrum.* **87**, 11D814 (2016).
19. L. Ballabio, J. Källne, and G. Gorini, *Nucl. Fusion* **38**, 1723 (1998).
20. O. Landoas, V. Yu. Glebov, B. Rossé, M. Briat, L. Disdier, T. C. Sangster, T. Duffy, J. G. Marmouget, C. Varignon, X. Ledoux, T. Caillaud, I. Thfoin, and J.-L. Bourgade, *Rev. Sci. Instrum.* **82**, 073501 (2011).
21. C. Waugh, “An Improved Method for Measuring the Absolute DD Neutron Yield and Calibrating Neutron Time-of-Flight Detectors in Inertial Confinement Fusion Experiments,” M.S. thesis, Massachusetts Institute of Technology, 2014.
22. C. J. Forrest, P. B. Radha, V. Yu. Glebov, V. N. Goncharov, J. P. Knauer, A. Pruyne, M. Romanofsky, T. C. Sangster, M. J. Shoup III, C. Stoeckl, D. T. Casey, M. Gatu-Johnson, and S. Gardner, *Rev. Sci. Instrum.* **83**, 10D919 (2012).
23. H. Brysk, *Plasma Phys.* **15**, 611 (1973).
24. V. Yu. Glebov, T. C. Sangster, C. Stoeckl, J. P. Knauer, W. Theobald, K. L. Marshall, M. J. Shoup III, T. Buczek, M. Cruz, T. Duffy, M. Romanofsky, M. Fox, A. Pruyne, M. J. Moran, R. A. Lerche, J. McNaney, J. D. Kilkenny, M. J. Eckart, D. Schneider, D. Munro, W. Stoeffl, R. Zacharias, J. J. Haslam, T. Clancy, M. Yeoman, D. Warwas, C. J. Horsfield, J.-L. Bourgade, O. Landoas, L. Disdier, G. A. Chandler, and R. J. Leeper, *Rev. Sci. Instrum.* **81**, 10D325 (2010).
25. T. J. Murphy, *Phys. Plasmas* **21**, 072701 (2014).