



# Multi-millijoule hollow-core fiber compression of short-wave infrared pulses to a single cycle

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**Abstract:** Approaches for efficient pulse compression can enable dramatic increases in the available peak power, as well as enable the generation of isolated attosecond X-ray pulses. Achieving high compression ratios for longer wavelength drivers has, however, been challenging. We present the compression of few-cycle 2.1  $\mu\text{m}$  central wavelength short-wave infrared laser pulses to 6.9 fs with 2.35 mJ pulse energy at a 10 kHz repetition rate. Electric field resolved measurements reveal a single cycle light field oscillation. With a carrier-envelope-phase stability of 131 mrad and average power fluctuations below 1 %, the system constitutes an excellent light source for strong-field experiments and attosecond physics.

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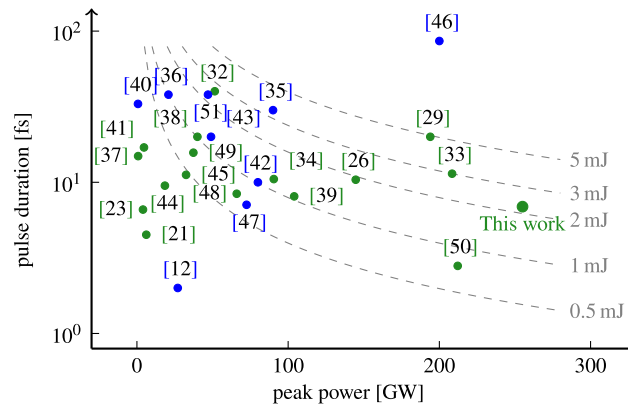
## 1. Introduction

Advances in laser development propel modern science and technology [1–3]. In particular the combination of a high peak power at infrared wavelengths and increasingly shorter laser pulse envelopes paves the way for strong-field experiments even in the relativistic region [4], or for attosecond science reaching the X-ray range [5]. In pursuit of the generation of such short laser pulses, frequency broadening in hollow-core fibers (HCFs) has emerged as the gold standard [6–8]. Whereas the generation of additional frequency components in the laser spectrum was initially decoupled from subsequent compression (e.g. via chirped mirrors [9,10]), novel approaches tune the fiber parameters for self-compression [11,12], soliton propagation [13], or the generation of dispersive waves in the ultraviolet [14–16] or mid-infrared [17]. In particular, the interplay between the dispersion introduced by the gas, the non-linear interaction and the dispersion of the waveguide allows self-compressing schemes, where those contributions counteract each other [11,14]. This development is profiting from the concept of stretched hollow-core fibers that can be extended to several meters in length [18]. Besides that, hollow-core photonic crystal fibers have successfully been employed for the generation of super broadband continua [7,15,19,20]. The maximum achievable peak power is limited by the damage threshold, but peak powers up to the gigawatt range have been demonstrated [21–23].

For the waveguide dispersion to become significant, smaller fiber core radii  $r$  are preferable due to the  $\propto r^{-2}$  scaling of the waveguide dispersion [14], even though they are prone to losses [14]. Typically, additional limitations are set by self-focusing and ionization [24–26] that prevent the usage of arbitrarily small fibers for high power lasers. For the achievement of self-compression in fibers, infrared laser sources are especially appealing due the increasing waveguide dispersion with longer wavelength [12,14,27,28]. A combination of state of the art short-wave infrared

(SWIR) laser systems with self-compressing spectral broadening systems thus promises the generation of very high-peak powers due to the aforementioned scaling laws.

In this article, we demonstrate a self-compressing broadening scheme for a multi-millijoule SWIR laser source. Driven by a  $2.1\ \mu\text{m}$  optical parametric chirped pulse amplifier [29], we generate a multi-octave spanning spectrum that self-compresses to 6.9 fs, which marks the single-cycle regime. With 2.35 mJ pulse energy at 10 kHz repetition rate and 138 mrad rms carrier-envelope-phase (CEP) fluctuations, the presented light source constitutes an excellent driver for high-harmonic generation with amplitude gating [5] or for the generation of tesla scale magnetic impulses [30,31]. To the best of our knowledge, our system constitutes the highest peak-power, multi-millijoule, multi-kilohertz, single-cycle and CEP stable SWIR laser system, see Fig. 1 for a comparison of related systems from the recent literature. Since high laser peak powers can be achieved by either increasing the pulse energy or by compressing the pulses and thus confining the energy in a shorter interval of time, it is necessary to compare the peak power of a system to the pulse duration to distinguish the technical approach. In Fig. 1, we compare the performance for SWIR systems with a repetition rate in the range from 1 kHz to 100 kHz. We estimate the peak power of the related literature systems, unless specified, by assuming that 90 % of the pulse energy is confined in the main peak of a  $\text{sech}^2$  pulse, which transfers to a conversion via  $P_{\text{peak}} = 0.88 \cdot \frac{0.9E_{\text{pulse}}}{\tau_{\text{FWHM}}}$ , where  $\tau_{\text{FWHM}}$  is the full-width at half-maximum pulse duration. This assumption might overestimate the performance of some systems. The assumption of a  $\text{sech}^2$  pulse shape is arbitrary, but the difference to other commonly used models, such as Gaussian pulses, is below 7 %. For our system, however, we use the exact value for the peak power extracted from the field-resolved data presented below.

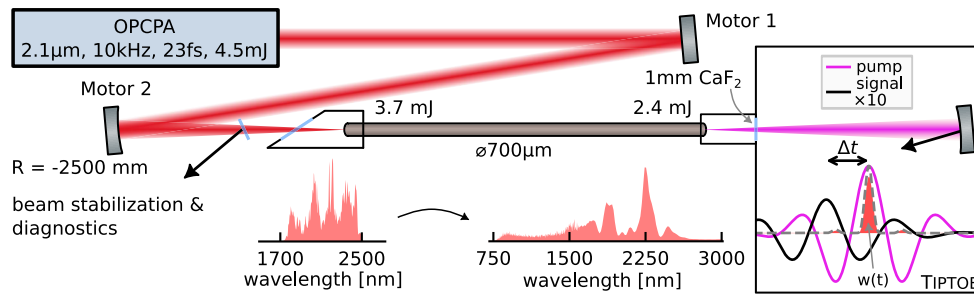


**Fig. 1.** Overview of state of the art laser systems in the SWIR region [12,21,23,26,29,32–51]. Green: CEP stable systems, blue: not CEP stable. The dashed lines indicate the pulse energy according to our conversion law. For systems with a peak power different than that, however, they do not apply.

## 2. Experimental approach

The laser front-end is a CEP-stable optical parametric chirped pulse amplifier reported elsewhere [29]. It is operated at 4.5 mJ pulse energy corresponding to 45 W average power at a central wavelength of  $\lambda_0 = 2.1\ \mu\text{m}$  with a pulse duration of 23 fs as confirmed via frequency-resolved optical-gating [52,53]. A portion of 3.75 mJ is focused with an  $R = -2500\ \text{mm}$  radius of curvature spherical mirror into a  $700\ \mu\text{m}$  core-diameter hollow fiber, see Fig. 2. The beam angle and position is actively stabilized (Aligna TEM Messtechnik GmbH) and the set point is used for alignment of the beam through the fiber. The latter is stretched between two internally developed

and adjustable fiber mounts and has a length of 2.4 m. One mount (depending on the desired setup) is placed on a movable stage for adjusting the tension of the fiber by a spring. That mechanism guarantees that a thermal expansion of the fiber during operation is compensated. We find that the expansion is typically less than 0.5 millimeters in our setup, therefore, the movable part can also be placed at the entrance since the maximum travel distance is smaller than the Rayleigh length. The fiber is filled with 700 mbar of Neon at a static pressure. After the exit, a 1 mm  $\text{CaF}_2$  window allows for evacuating the subsequent beam path or for coupling the beam out for experiments at ambient air. After re-collimation, the beam is sent to a T1PTOE setup as well as to a  $2f$ - $3f$  interferometer for CEP control and measurement, besides other means of diagnostics. The input and output spectra are amplitude calibrated with a calibration lamp and shown in Fig. 2. The input spectrum is taken with an OceanOptics NIRQuest-512 (0.9 to  $2.5\ \mu\text{m}$ ), whereas the output spectrum is taken with three devices, OceanOptics HDX (Si) and NIRQuest-512 ( $0.8\ \mu\text{m}$ - $1.6\ \mu\text{m}$ , InGaAs), as well as Spectral Products SM301-EX (PbSe).



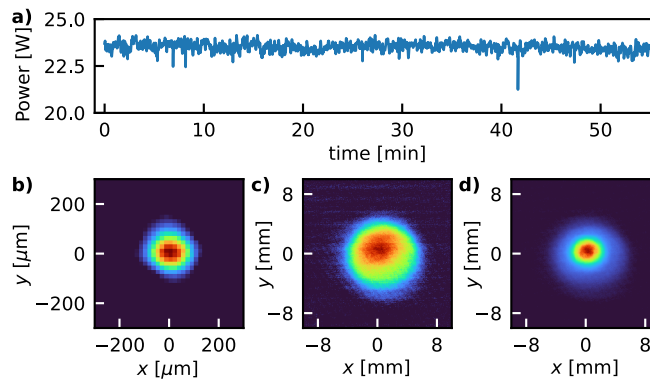
**Fig. 2.** Fiber broadening with the  $2.1\ \mu\text{m}$  OPCA laser. The black boxes indicate vacuum chambers that can be pumped and filled individually. The distances in the figure are not to scale. The spectra shown have a linear scale for the intensity amplitude. The T1PTOE box also includes other means of diagnostics, such as a  $2f - 3f$  interferometer or a powermeter.

We characterize the output of our system with the tunneling ionization with a perturbation for the time-domain observation of an electric field (T1PTOE) [54] which has proven its reliability in measuring laser pulses from the ultraviolet, visible or infrared regime [55], via ionization in gases [54], solids [56] or micro- and nano-structures [57–59]. As a first experimental application of the multi-octave spanning spectrum of our laser pulses, we investigate CEP effects in the spectral response of T1PTOE. In particular, we exploit the CEP dependent sensitivity of T1PTOE at harmonics of the central frequency [56,57,59] for an intrinsic calibration of the CEP owing to our broad spectrum.

The main principle of T1PTOE, as presented in Ref. [54] and outlined in the corresponding box in Fig. 2, is based on a strong pump beam,  $E_{\text{Pump}}$ , that generates an ion yield  $Q$  in a gas by sub-cycle field ionization during its strongest optical half-cycle. This ion yield is linearly perturbed by a second weak beam,  $E_{\text{Signal}}$ , delayed by a time  $\Delta t$ , which modulates the ion yield according to the interference of both beams during the sub-cycle ionization gate. The ion yield can be estimated via the ionization rate  $w(t)$ , see Supplement 1. For those measurements, we employ a Mach-Zehnder-type interferometer that contains perforated mirrors as dispersion free beam-splitters. The transmitted part of the beam ( $E_{\text{Signal}}$ ) contains a piezo-driven retro reflector and an optical chopper enabling lock-in detection of the ion yield modulation. For delay control, the interference pattern of a co-propagating frequency-stabilized HeNe laser is detected with a camera and stabilized by feedback to the piezo driver. The intensity ratio in the focus is  $I_{\text{Signal}}/I_{\text{Pump}} \approx 3 \times 10^{-4}$ , where Pump refers to the reflected beam in the interferometer. The perturbing field strength is only 1.7% of that of the pump.

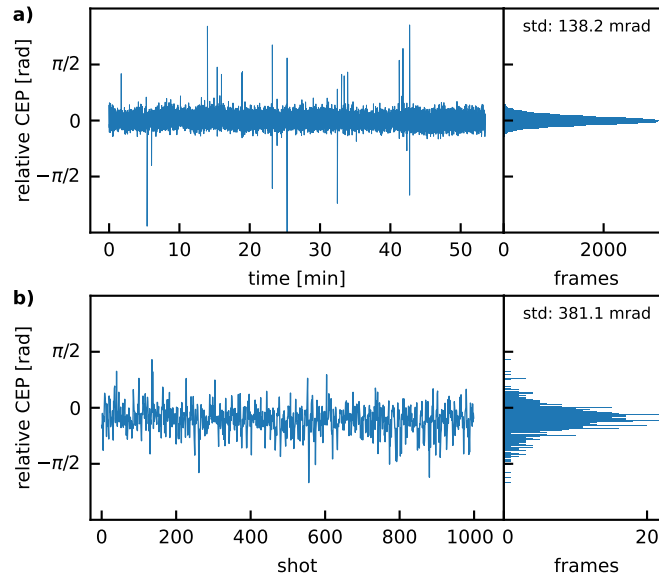
### 3. Power and CEP stability

To uncover the performance of our fiber broadening system, we investigate the power and CEP stability as well as the beam profile. We achieved a transmitted power of 23.5 W that is stable within a standard deviation of 0.226 W, corresponding to 0.96 %, as measured over a time interval of more than 50 minutes (Fig. 3(a)). The corresponding transmission is 63 %. The expected value we found from simulation with the code from Ref. [60] is around 90 % and deviates from our value due to in-coupling losses in our setup. The beam-profile of the fiber output in the focus of a  $f = 400$  mm spherical mirror measured with a micro-bolometer camera is shown in Fig. 3(b), together with the profile of the collimated beam (part c) and d)). In Fig. 3(c), we added a  $1\ \mu\text{m}$  longpass filter to show that the short wavelengths are closer on axis when compared to the full beam-profile in d). The spatial distribution of the spectral components has influence on the field-measurements, see discussion in the [Supplement 1](#).



**Fig. 3.** a) Power stability measurement yielding 23.5 W average power at the fiber exit with a standard deviation of 226 mW, which is 0.96 %. b) Focus of the fiber output with a  $f = 40$  cm spherical mirror on a Rigi Microbolometer camera. c) Collimated beamprofile on a Pyrocam IV, with  $1\ \mu\text{m}$  longpass filter and without (d)).

We confirmed the CEP stability of the fiber output by measuring  $2f$ - $3f$  interference fringes in an out-of-loop measurement, while stabilizing it with an in-loop PID control. Here, the  $2f$  spectral components generated via frequency doubling in  $\beta$ -barium borate overlap conveniently in the range of silicon detector with the third harmonic that is generated in the fiber. Thus, we could exploit readily available silicon based spectrometers for detection. The extracted relative CEP from the interference fringes is detected over an extended period of time as shown in Fig. 4(a) and its fluctuations amount to a standard deviation of 138.2 mrad. Here, we acquire a spectrogram every 64 ms, thus fast CEP modulations are not resolved. When measuring the single-shot CEP stability of our system, we find a value of 381.1 mrad, see Fig. 4(b). There, we use our triggered spectrometer (Avantes Avaspec ULS-4096CL-EVO-UA-10) to acquire a spectrum for a single laser shot. We set the integration time to  $90\ \mu\text{s}$ , shorter than the  $100\ \mu\text{s}$  time between two consecutive laser pulses to ensure only a single pulse is detected. However, we cannot measure the CEP for every consecutive shot due to a dead time of the device after an acquisition of  $698\ \mu\text{s}$  [61]. Thus, we expect that at least 7 pulses are skipped between each frame in the single-shot data. Nevertheless, we believe that the single-shot measurement confirms that the CEP stability is maintained from a near shot-to-shot basis to a one-hour-scale. To the best of our knowledge, this is the highest CEP stability of a multi-kilohertz, multi-millijoule, single-cycle  $2\ \mu\text{m}$  based system.

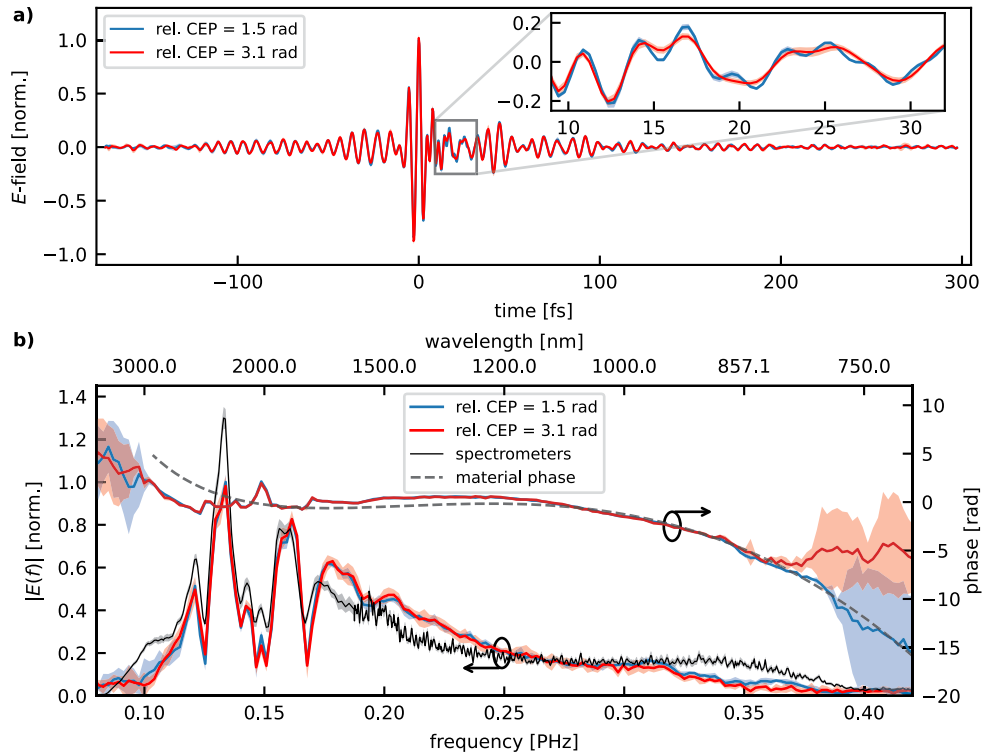


**Fig. 4.** **a)** The extracted phase of the  $2f$ - $3f$ -fringes after the fiber shows a standard deviation of 138 mrad over a time of more than 50 minutes. **b)** Single-shot CEP data exhibiting a standard deviation of 381.1 mrad for 1000 shots.

#### 4. Characterization of the time-domain electric field

The measurement of the pulse shape and duration is particularly challenging for super broadband spectra. For multi-octave spanning spectra, the measurement of the electric field of the laser pulse is the appropriate approach. Therefore, we perform Tiptoe-measurements [54] for the characterization of our system. The measured Tiptoe waveforms are shown in Fig. 5(a) for a relative CEP of the input pulses of 1.5 rad and 3.1 rad as stabilized with the  $2f$ - $3f$  interferometer. The full-width at half-maximum (FWHM) pulse duration of the intensity envelope is 6.9 fs, which is even shorter than the cycle-time associated with the central wavelength of the input of  $\approx 7$  fs. This value corresponds to a peak power of 0.26 TW. For the time-domain electric fields, we apply a Tukey-window in Fourier space to select the frequency components between 0.03 and 0.5 PHz. To sample such short pulses, we place an additional 0.3 mm fused silica plate before the Tiptoe setup when the chamber is pumped down to a millibar regime, and 0.6 mm fused silica if at room pressure. We do not observe a significant elongation of the main pulse in the Tiptoe measurement depending on the gas pressure when compensating for the dispersion, however, we do see the formation of side pulses due to the higher amount of dispersive media being present in the beam path. Additionally, we observe an increasing signal-to-noise ratio when reducing the pressure. We believe that this is an effect of the bias voltage (10.6 V) being more appropriate at lower pressures, similarly to Ref. [54], where a higher bias voltage was used in the ambient air measurements. The data shown in Fig. 5(a) is taken at a pressure of 10 mbar.

The CEP is shifted for both pump and signal beam synchronously. This simultaneous shift can, however, not be measured with Tiptoe since it is only sensitive to the CEP difference between both beams (c.f. Refs. [54,55,57,59] and Supplement 1). Thus, the CEP of the measured waveform is not supposed to change with the CEP of the laser pulses and only represents the actual electric field if the CEP of the pump beam is 0. Consequently, a calibration of the CEP of the pump is necessary which was done previously using a second harmonic field [54]. Profiting from the multi-octave bandwidth of our laser system, we can intrinsically calibrate the CEP by observing the resolved spectral amplitudes in the  $2f$  region of the central frequency  $f$ , in



**Fig. 5.** **a)** Time-domain electric field (blue) and measured waveform for a different CEP (red). **b)** spectral field amplitudes corresponding to the waveforms in a). Effects of the CEP of the laserpulses are visible between 0.3 and 0.38 PHz. Spectral amplitudes of the fiber output measured with spectrometers (black line). The arrows indicate the corresponding axis.

particular from 0.3 to 0.38 PHz. There, the resolved spectral amplitudes are enhanced for a CEP of the pump of 0. The results of a CEP scan and the observation of the expected modulations in the  $2f$  region, see supplement, identify the relative CEP of 1.5 rad with a pump beam CEP of 0. Thus, the waveform shown in that case represents the electric field of the signal beam.

The spectral amplitudes and phases of the T1PTOE data are shown in Fig. 5(b), together with spectrometer data (black line). Overall, a good agreement of both curves is found. The residual differences of the measured spectral amplitudes to the spectrometer data originate in the response function of T1PTOE and in the geometry of the interferometer, see Supplement 1. As apparent from Fig. 5(b), the spectral phase is flat over a broad spectral region but exhibits a curvature at higher frequencies. Its similarity to the expected spectral phase of the materials in the beam path from the fiber exit to the measurement point (black dashed line) confirms that the output phase at the fiber exit is flat. Thus, the presented broadening scheme is in principle self-compressing via means of soliton self-compression (see Supplement 1), where the fiber length aligns with the soliton fission length. The propagation through gas and material introduce a small amount of net dispersion that is responsible for the observation of the faster oscillations after the main pulse, see inset in Fig. 5(a). A polynomial fit yields a residual group delay dispersion of  $-3 \text{ fs}^2$  and a third order dispersion of  $35 \text{ fs}^3$ . Note that in our sign convention, the field at negative times arrives earlier at target than the one at positive times which defines the sign of the curvature of the spectral phase. Those oscillations are only detectable for correct CEP of the pump beam since they originate in the high-frequency part of the spectrum. For the spectral phase of the



material, we use the refractive index data of Refs. [62–64] for 0.7 m of Neon at 700 mbar, 1 mm of CaF<sub>2</sub>, and 0.3 mm of fused silica, respectively. We subtract the constant and linear terms to match the phase we found in the experiment. The data of Ref. [62] is only measured in the visible to near-infrared range and is extrapolated to our bandwidth which is a possible explanation for the deviation in the long wavelength range. Additionally, plasma effects on the spectral broadening are found to have minor influence, which is supported by the numerical studies of the fiber dynamics with and without plasma effect (see [Supplement 1](#)), but also with overall good agreement of the spectral phase with the material phase in the beam path.

Overall, the multi-octave bandwidth allows for an in-situ CEP calibration due to the observed modulations in the  $2f$  region. Even though the response phase of TPTOE has been discussed and observed before [56,57,59], an investigation of the modulations of the  $2f$  components has hitherto been hindered by too narrow band laser sources and therefore required the generation of a second harmonic [54]. With our light source, however, we intrinsically cover that region and can thus calibrate the CEP without any other calibration setups. Intensities in the range of  $10^{17}$  W/cm<sup>2</sup> are achievable with our system which may drive future strong field experiments even in the relativistic region, at a multi-kilohertz repetition rate.

## 5. Conclusion

We have demonstrated a high-power short-wave infrared spectral broadening scheme providing CEP stability, multi-millijoule pulse energy and high average power. The actual electric field waveform has been successfully characterized with TPTOE and due to the two octave bandwidth, an in-situ CEP calibration was possible. The self-compressing properties of the fiber eliminate the necessity of post-compression via dispersive mirrors which are prone to transient non-linearities even at moderate laser intensities that degrade the beam properties [65]. Additionally, the bandwidth limitations of dispersive mirrors can only be overcome by complex light-field synthesis setups [66–68] which our approach eliminates, resulting in a comparatively simple and robust pulse compression scheme. Our approach bypasses constraints for the polarization that are typically caused by dispersive mirrors and thus allows for spectral broadening of arbitrary polarization states. Even though there are new approaches with polarization independent chirped mirrors [69], the supported bandwidth is typically narrower than what is achievable with a self-compression approach as described in this article. We also demonstrated that a static pressure can be used in the fiber and since the window material dispersion and the dispersion in gases counteract each other, a small amount of material can be placed without significant performance degradation. Thus, the compression scheme is appealing to applications outside a vacuum environment.

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**Disclosures.** The authors declare no conflicts of interest.

**Data availability.** Data underlying the results presented in this paper are not publicly available at this time but may be obtained from the authors upon reasonable request.

**Supplemental document.** See [Supplement 1](#) for supporting content.

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