

Graphene Q-switched, tunable fiber laser

D. Popa, Z. Sun, T. Hasan, F. Torrisi, F. Wang, and A. C. Ferrari^{a)}

Department of Engineering, University of Cambridge, Cambridge CB3 0FA, United Kingdom

(Received 30 October 2010; accepted 17 January 2011; published online 15 February 2011)

We demonstrate a wideband-tunable Q-switched fiber laser exploiting a graphene saturable absorber. We get ~ 2 μs pulses, tunable between 1522 and 1555 nm with up to ~ 40 nJ energy. This is a simple and low-cost light source for metrology, environmental sensing, and biomedical diagnostics. © 2011 American Institute of Physics. [doi:10.1063/1.3552684]

Q-switching and mode-locking are the two main techniques enabling pulsed lasers.¹ In mode-locking, the random phase relation originating from the interference of cavity modes is fixed, resulting in a single pulse,¹ with typical duration ranging from tens ps to sub-10 fs,² and a repetition rate corresponding to the inverse of the cavity round-trip time.² Many aspects, including the dispersive and nonlinear properties of the intracavity components, need to be precisely balanced in order to achieve stable operation.^{1,2} Q-switching is a modulation of the quality factor, Q , of a laser cavity,¹ Q being the ratio between the energy stored in the active medium and that lost per oscillation cycle¹ (thus, the lower the losses, the higher the Q). In Q-switching, the active medium is pumped while lasing is initially prevented by a low Q factor.¹ The stored energy is then released in a pulse with duration ranging from μs to ns when lasing is allowed by a high Q factor.¹ The time needed to replenish the extracted energy between two consecutive pulses is related to the lifetime of the gain medium, which is typically $\sim\text{ms}$ for erbium-doped fibers.¹ Thus the repetition rate of Q-switched lasers is usually low [$\sim\text{kHz}$ (Ref. 1)], much smaller than mode-locked lasers.^{1,2} On the other hand, Q-switching enables much higher pulse energies and durations than mode-locking.¹ Q-switching has advantages in terms of cost, efficient operation (i.e. rate of output pulse energy to input power), and easy implementation, compared to mode-locking, which needs a careful design of the cavity parameters to achieve a balance of dispersion and nonlinearity.^{1,2} Q-switched lasers are ideal for applications where ultrafast pulses (<1 ns) are not necessary, or long pulses are advantageous,^{3,4} such as material processing, environmental sensing, range finding, medicine, and long-pulse nonlinear experiments.³⁻⁵

Q-switching can be active (exploiting, e.g., an acousto-optic or electro-optic modulator¹), or passive [using, e.g., a saturable absorber (SA)¹]. Passive Q-switching features a more compact geometry and simpler setup than active, which requires additional switching electronics.¹ For Q-switching the SA recovery time does not need to be shorter than the cavity round-trip time, since the pulse duration mainly depends on the time needed to deplete the gain after the SA saturates,^{1,2} unlike mode-locking.² Doped bulk crystals,⁵ and semiconductor SA mirrors (SESAMs)^{3,6} are the most common SAs in passive Q-switching.¹ However, the use of doped crystals as SAs requires extra elements (mirrors, lenses) to focus the fiber output into the crystal.⁵ SESAMs

have limited operation bandwidth, typically few tens nm,⁷ thus are not suitable for broad-band tunable pulse generation. Broadband SAs enabling easy integration into an optical fiber system are thus needed to create a compact Q-switched fiber laser.

Single wall carbon nanotubes (SWNTs) and graphene are ideal SAs, due to their low saturation intensity, low-cost and easy fabrication.⁸⁻²³ Broadband operation is achieved in SWNTs using a distribution of tube diameters,^{8,18} while this is an intrinsic property of graphene, due to the gapless linear dispersion of Dirac electrons.^{19-22,24} Q-switching was reported using SWNTs: Ref. 25 achieved 14.1 nJ pulse energy and 7 μs width, while Ref. 26 reported 13.3 nJ and 700 ns. After our demonstration of a graphene-based mode-locked laser,¹⁸ various groups implemented graphene SAs in a variety of cavity designs.^{19-23,27-29}

Here, we demonstrate a fiber laser Q-switched by a graphene SA (GSA). The broadband absorption of graphene enables Q-switching over a 32 nm range, limited only by our tunable filter, not graphene itself. The pulse energy is ~ 40 nJ, for ~ 2 μs duration.

Graphite flakes are exfoliated by mild ultrasonication with sodium deoxycholate (SDC).^{20,22,30} A dispersion en-

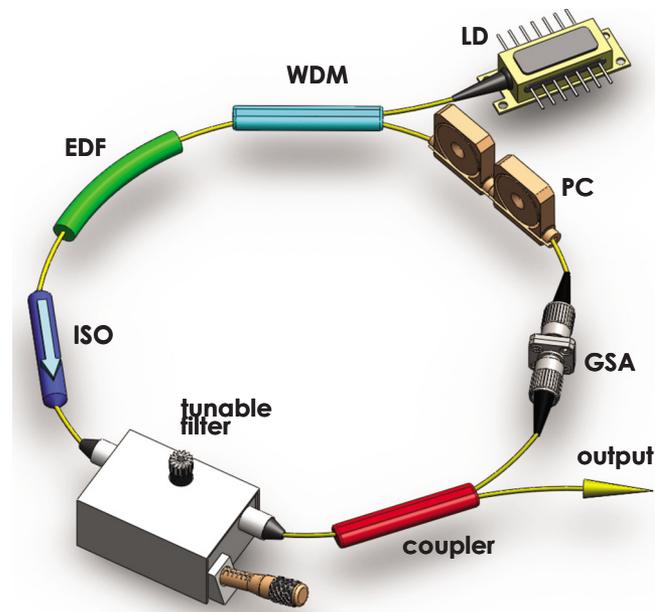


FIG. 1. (Color online) Schematic setup of our graphene-Q switched laser: laser diode (LD), wavelength division multiplexer (WDM), erbium doped fiber (EDF), isolator (ISO), graphene-based saturable absorber (GSA), and polarization controller (PC).

^{a)}Electronic mail: acf26@eng.cam.ac.uk.

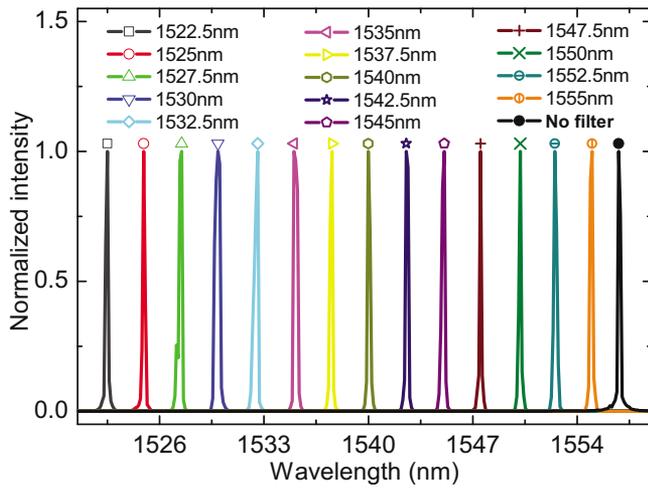


FIG. 2. (Color online) Output spectra for 14 tuning wavelengths. The curve with a filled circle corresponds to Q-switching without filter.

riched with single (SLG) and few layer graphene (FLG)²² is then mixed with an aqueous solution of polyvinyl alcohol (PVA). After water evaporation, a $\sim 50 \mu\text{m}$ thick graphene-PVA composite is obtained.^{18,20} This is then placed between two fiber connectors to form a fiber-compatible SA, then integrated into a laser cavity, Fig. 1, with 1.25 m erbium-doped fiber (EDF) as gain medium, pumped with a 980 nm laser diode (LD), coupled via a wavelength division multiplexer (WDM). An optical isolator (ISO) ensures unidirectional light propagation. An in-line tunable optical bandpass filter is inserted after the ISO. Our EDF can support lasing between 1520 and 1560 nm.³¹ The operation wavelength is selected by rotating the dielectric interference filter. The 20% port of an optical coupler provides the laser output. The rest of the cavity consists of a combination of single mode fiber (SMF) Flexcor 1060 and SMF-28. All fibers used in our cavity are polarization-independent, i.e., they support any light polarization, even if this changes as a result of outside perturbations (e.g., mechanical stresses, bending, or temperature). Thus, to improve the output pulse stability, we place in the cavity a polarization controller (PC), consisting of 2 spools of SMF-28 fiber acting as retarders. The total retardation induced by the PC is a function of the fiber geometry in the spool.³¹ This allows to maintain a given polarization state after each round trip. The total cavity length is $\sim 10.4 \text{ m}$. The operation is evaluated by a 14 GHz bandwidth photodetector and an oscilloscope. A spectrum analyzer with 0.07 nm resolution measures the output spectrum.

Continuous wave operation starts at $\sim 43 \text{ mW}$ pump power; pulsed operation at $\sim 74 \text{ mW}$. The repetition rate is pump-dependent up to $\sim 200 \text{ mW}$ [Fig. 4(b)], a typical signature of Q-switching.¹ The output spectrum is tunable from ~ 1522 to 1555 nm. This is comparable to the 31 nm range reported for doped crystal Q-switched tunable lasers,⁵ but much larger than the 5 nm thus far achieved for SWNT Q-switched lasers.^{25,26} Our tuning range is limited by the filter and by the EDF gain, not by the GSA. Figure 2 shows the output spectra for 14 wavelengths at $\sim 2.5 \text{ mW}$ output power. Without filter, the laser exhibits Q-switching at 1557 nm. The full width at half maximum (FWHM) spectral width is $0.3 \pm 0.1 \text{ nm}$ over the whole tuning range, much shorter than thus far achieved for graphene mode-locked lasers.^{18-23,27-29}

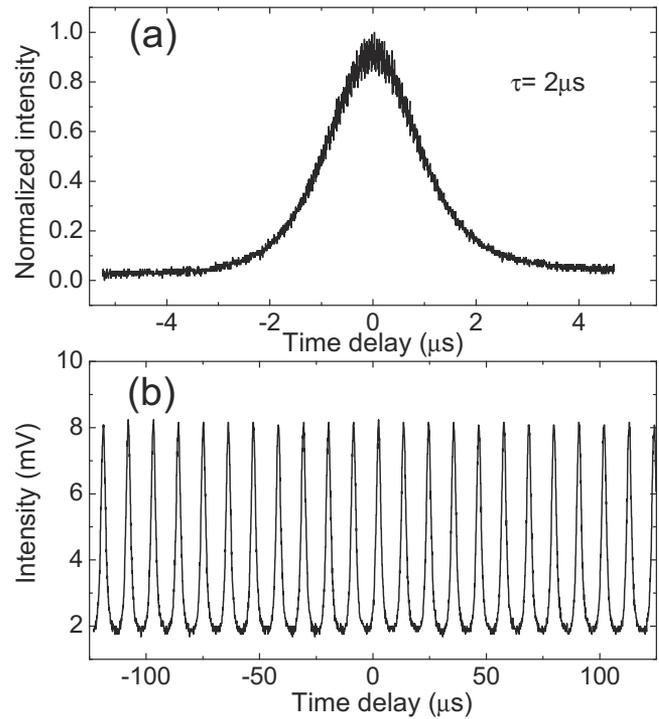


FIG. 3. (a) Single pulse envelope. (b) Typical pulse train for 2.8 mW output power.

Figure 3(a) plots a typical pulse envelope, having FWHM $\sim 2 \mu\text{s}$, comparable to fiber lasers Q-switched with other SAs (e.g., SESAMs,^{3,6} doped crystals,⁵ and SWNTs^{25,26}), but much longer than thus far achieved in graphene mode-locked fiber lasers.^{19-23,27-29} The output pulse duration has little dependence on wavelength, possibly due to the flat gain coefficient of our EDF.³¹ Figure 3(b) shows the pulse train for a typical laser output at 158 mW pump power.

The output power varies from 1 to 3.4 mW as a function of pump power [Fig. 4(a)]. The slope efficiency, i.e., the slope of the line obtained by plotting the laser output power against the input pump power,¹ is $\sim 2\%$. The repetition rate as a function of pump power varies from 36 to 103 kHz [Fig. 4(b)], with a 67 kHz change for a 2.4 mW output power variation. Unlike mode-locked lasers, where the repetition

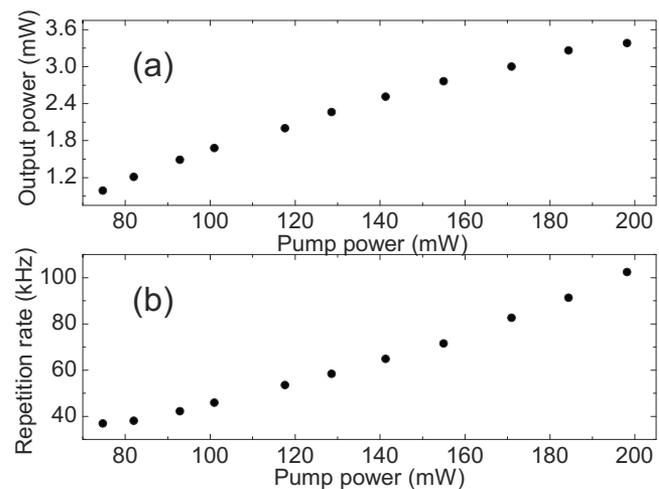


FIG. 4. (a) Output power and (b) repetition rate, as a function of input pump power at 1540 nm

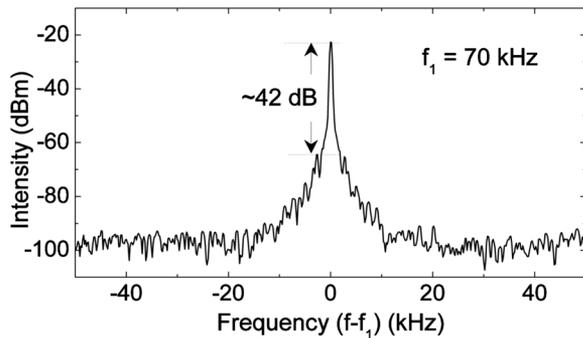


FIG. 5. Output rf spectrum measured around ~ 70 kHz at 1540 nm.

rate is fixed by the cavity length,¹ in Q-switched lasers this depends on pump power.¹ As this increases, more gain is provided to saturate the SA. Since pulse generation relies on saturation, the repetition rate increases with pump power.¹ The maximum output pulse energy is ~ 40 nJ for ~ 60 kHz repetition rate, similar to that achieved using other SAs.²⁶ Compared to graphene mode-locked fiber lasers,^{19–23,27–29} our pulse energy is ~ 6 times larger, but with less peak power, due to the larger pulse duration. It is also much larger than thus far achieved in SWNT Q-switched lasers.^{25,26} Even higher energies, thus peak powers, could be enabled by evanescent field interaction with GSA²⁸ and high-gain fibers (e.g., cladding-pumped fibers⁵ or large mode area fibers³).

The radio-frequency (rf) measurement of the output intensity at 70 kHz, corresponding to a period of ~ 143 μ s, is shown in Fig. 5. The peak to pedestal extinction is ~ 40 dB (10^4 contrast), confirming pulse stability.

In conclusion, we achieved Q-switching exploiting a graphene-based SA, using standard, telecom grade, fiber components. The wideband operation of graphene enables broad-band tunability. Such wideband Q-switched laser could provide a simple, low-cost, and convenient light source for metrology, environmental sensing, and biomedical diagnostics.

We acknowledge funding from EPSRC (Grant Nos. GR/S97613/01 and EP/E500935/1) ERC grant NANOPOTS, a Royal Society Brian Mercer Award for Innovation, King's College Cambridge, the Cambridge Integrated Knowledge Centre in Advanced Manufacturing Technology for Photonics and Electronics, the EU grants GENIUS and RODIN, and Nokia Research Centre, Cambridge. ACF is a Royal Society Wolfson Research Merit Award holder.

¹O. Svelto, *Principles of Lasers*, 4th ed. (Plenum, New York, 1998).

²U. Keller, K. J. Weingarten, F. X. Kartner, D. Kopf, B. Braun, I. D. Jung, R. Fluck, C. Honninger, N. Matuschek, and J. A. der Au, *IEEE J. Sel. Top.*

Quantum Electron. **2**, 435 (1996).

³R. Paschotta, R. Haring, E. Gini, H. Melchior, U. Keller, H. L. Offerhaus, and D. J. Richardson, *Opt. Lett.* **24**, 388 (1999).

⁴M. Siniava, M. Siniavsky, V. Pashinin, A. Mamedov, V. Konov, and V. Kononenko, *Laser Phys.* **19**, 1056 (2009).

⁵M. Laroche, A. M. Chardon, J. Nilsson, D. P. Shepherd, W. A. Clarkson, S. Girard, and R. Moncorge, *Opt. Lett.* **27**, 1980 (2002).

⁶S. Kivistö, R. Koskinen, J. Paajaste, S. D. Jackson, M. Guina, and O. G. Okhotnikov, *Opt. Express* **16**, 22058 (2008).

⁷O. Okhotnikov, A. Grudinin, and M. Pessa, *New J. Phys.* **6**, 177 (2004).

⁸F. Wang, A. G. Rozhin, V. Scardaci, Z. Sun, F. Hennrich, I. H. White, W. I. Milne, and A. C. Ferrari, *Nat. Nanotechnol.* **3**, 738 (2008).

⁹G. Della Valle, R. Osellame, G. Galzerano, N. Chiodo, G. Cerullo, P. Laporta, O. Svelto, A. G. Rozhin, V. Scardaci, and A. C. Ferrari, *Appl. Phys. Lett.* **89**, 231115 (2006).

¹⁰S. Y. Set, H. Yaguchi, Y. Tanaka, and M. Jablonski, *IEEE J. Sel. Top. Quantum Electron.* **10**, 137 (2004).

¹¹A. G. Rozhin, V. Scardaci, F. Wang, F. Hennrich, I. H. White, W. I. Milne, and A. C. Ferrari, *Phys. Status Solidi B* **243**, 3551 (2006).

¹²A. Martinez, K. Zhou, I. Bennion, and S. Yamashita, *Opt. Express* **18**, 11008 (2010).

¹³M. A. Solodyankin, E. D. Obraztsova, A. S. Lobach, A. I. Chernov, A. V. Tausenev, V. I. Konov, and E. M. Dianov, *Opt. Lett.* **33**, 1336 (2008).

¹⁴Z. Sun, A. G. Rozhin, F. Wang, V. Scardaci, W. I. Milne, I. H. White, F. Hennrich, and A. C. Ferrari, *Appl. Phys. Lett.* **93**, 061114 (2008).

¹⁵Z. Sun, T. Hasan, F. Wang, A. G. Rozhin, I. H. White, and A. C. Ferrari, *Nano Res.* **3**, 404 (2010).

¹⁶Z. Sun, A. G. Rozhin, F. Wang, T. Hasan, D. Popa, W. O'Neill, and A. C. Ferrari, *Appl. Phys. Lett.* **95**, 253102 (2009).

¹⁷T. R. Schibli, K. Minoshima, H. Kataura, E. Itoga, N. Minami, S. Kazaoui, K. Miyashita, M. Tokumoto, and Y. Sakakibara, *Opt. Express* **13**, 8025 (2005).

¹⁸T. Hasan, Z. Sun, F. Wang, F. Bonaccorso, P. H. Tan, A. G. Rozhin, and A. C. Ferrari, *Adv. Mater.* **21**, 3874 (2009).

¹⁹Z. Sun, D. Popa, T. Hasan, F. Torrisi, F. Wang, E. J. R. Kelleher, J. C. Travers, V. Nicolosi, and A. C. Ferrari, *Nano Res.* **3**, 653 (2010).

²⁰Z. Sun, T. Hasan, F. Torrisi, D. Popa, G. Privitera, F. Wang, F. Bonaccorso, D. M. Basko, and A. C. Ferrari, *ACS Nano* **4**, 803 (2010).

²¹F. Bonaccorso, Z. Sun, T. Hasan, and A. C. Ferrari, *Nat. Photonics* **4**, 611 (2010).

²²T. Hasan, F. Torrisi, Z. Sun, D. Popa, V. Nicolosi, G. Privitera, F. Bonaccorso, and A. C. Ferrari, *Phys. Status Solidi B* **247**, 2953 (2010).

²³D. Popa, Z. Sun, F. Torrisi, T. Hasan, F. Wang, and A. C. Ferrari, *Appl. Phys. Lett.* **97**, 203106 (2010).

²⁴A. K. Geim and K. S. Novoselov, *Nature Mater.* **6**, 183 (2007).

²⁵D. Zhou, L. Wei, B. Dong, and W. Liu, *IEEE Photon. Technol. Lett.* **22**, 9 (2010).

²⁶B. Dong, C. Liaw, J. Hao, and J. Hu, *Appl. Opt.* **49**, 5989 (2010).

²⁷H. Zhang, D. Y. Tang, L. M. Zhao, Q. L. Bao, and K. P. Loh, *Opt. Express* **17**, 17630 (2009).

²⁸Y. W. Song, S. Y. Jang, W. S. Han, and M. K. Bae, *Appl. Phys. Lett.* **96**, 051122 (2010).

²⁹A. Martinez, K. Fuse, B. Xu, and S. Yamashita, *Opt. Express* **18**, 23054 (2010).

³⁰Y. Hernandez, V. Nicolosi, M. Lotya, F. M. Blighe, Z. Y. Sun, S. De, I. T. McGovern, B. Holland, M. Byrne, Y. K. Gun'ko, J. J. Boland, P. Niraj, G. Duesberg, S. Krishnamurthy, R. Goodhue, J. Hutchison, V. Scardaci, A. C. Ferrari, and J. N. Coleman, *Nat. Nanotechnol.* **3**, 563 (2008).

³¹G. Agrawal, *Applications of Nonlinear Fiber Optics* (Academic Press, San Diego, CA, 2001).