Single-Mode Lasing from a Single 7 nm Thick Monolayer of Colloidal Quantum Wells in a Monolithic Microcavity

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In this work, monolithically-fabricated vertical cavity surface emitting lasers (VCSELs) of densely-packed, orientation-controlled, atomically flat colloidal quantum wells (CQWs) using a self-assembly method is reported and single-mode lasing from a record thin colloidal gain medium with a film thickness of 7 nm under femtosecond optical excitation is demonstrated. Specially engineered CQWs are used to demonstrate these hybrid CQW-VCSELs consisting of only a few layers to a single monolayer of CQWs and are achieved the lasing from these thin gain media by thoroughly modeling and implementing a vertical cavity consisting of distributed Bragg reflectors with an additional dielectric layer for mode tuning. Accurate spectral and spatial alignment of the cavity mode with the CQW films is secured with the help of full electromagnetic computations. While overcoming the long-pending problem of limited electrical conductivity in thicker colloidal films, such ultrathin colloidal gain media can be helpful to enable fully electrically-driven colloidal lasers.

1. Introduction

Lasers based on solution-processed materials have attracted intensive scientific and technological interest owing to their exceptional advantages including ease of fabrication, low-cost synthesis, and processability and precise control of emission wavelength enabling lasing readily in different colors. As a result, a wide range of solution-processed materials of different classes have been studied for lasing, including organic semiconductors,[1] perovskites,[2] and semiconductor nanocrystals (NCs).[3–5] Successful incorporation of such emitters into lasers thus far have helped expanding the use of lasers into new areas, such as medical and biological imaging,[6] chemical sensing,[7] lab-on-a-chip diagnostics,[8] and on-chip integrated photonic circuits.[9,10] Among solution-processable light-emitting materials studied for lasing applications, colloidal quantum wells (CQWs) have gained considerable attention recently owing to their attractive properties including large absorption cross-section,[11] giant oscillator strength,[12] large material gain coefficient,[13] giant gain cross-section,[14] and ultranarrow emission spectra stemming from pure carrier confinement in one dimension.[15] Several studies have been conducted to overcome fast Auger recombination (AR) in order to develop practical CQW lasers,[16–18] and type-II CQWs,[19] giant-shell CQWs,[17] and gradually interfaced core/shell CQWs,[18] have been introduced for this purpose. AR can also be in principle controlled by optimizing cavity designs and their interactions with the gain media. For example, coupling CQW layers to cavities with small mode volumes and high quality factors (Q-factors), such as micropillars, microdisks, and photonic crystal cavities, can enhance the spontaneous emission (SE) due to the Purcell effect (PE),[20] as a result providing means of overcoming fast AR. This issue requires careful design of cavities offering optimized optical mode overlap coefficients (Γ) and Q-factors.

Monolithic fabrication of ultrathin vertical cavity surface emitting laser (VCSEL) provides opportunities for enhancing Q-factor and enabling single-mode lasing. Single-mode lasers purified from noise of other lasing modes are advantageous in practical applications and typically enable lower lasing thresholds compared to multimode lasers. In addition, monolithic CQW laser fabrication is crucial for enhancing the VCSEL performance because parasitic scattering in the cavity originating from multicomponent setup is prevented along with the surface disruptions. Previously, CQW-VCSELs were fabricated by sandwiching drop-casted several-micrometer-thick CQW films
between two mirrors resulting in bulky devices suffering from heat generation, exhibiting small Q-factors, and multimode lasing. This further leads to increased surface roughness and formation of stacks, which in turn significantly increase the scattering losses within the cavity. For monolithic fabrication of CQW-VCSELs, deposition of uniform CQW films is required to tune the cavity thickness and the resonance wavelength as well as to obtain cavities with high Q-factors. Recently, our group has reported CQW film deposition with a self-assembly technique, which allows for thickness and orientation-controlled deposition of monolayers of CQWs on solid substrates with near unity surface coverage and sub-nm surface roughness across cm$^2$-large areas. Our method is compatible with well-developed micro- and nanofabrication technologies and can integrate CQWs with developed photonic components and in a dense footprint on a chip scale, which is critical for commercial mass-production. The uniform layering of top dielectric layers after CQW deposition and the monolithic chip-scale device fabrication needs the thermal stability of the underlying CQW film. CQW films with oleic acid ligands are appropriate for monolithic VCSEL fabrication because they are stable at temperatures up to 200 °C.

The poor confinement of the resonant mode in thin films of few monolayers could possibly lead to high optical losses, consequently making the lasing impossible. As a result, there are very few reports on lasing from thin films of colloidal NCs. Roh et al. achieved lasing with a threshold of 18.8 μJ cm$^{-2}$ from a quantum dot (QD) gain medium as thin as 50 nm using a distributed feedback resonator. Kim et al. reported single-mode lasing from a colloidal QD gain medium only 35 nm thick with a threshold of 20 mJ cm$^{-2}$. Achieving lasing from films of CQWs thinner than hundreds of nanometers, which has the advantage of giant modal gain compared to all other types of NCs (QDs, nanorods, and nanowires), has not been previously accomplished.

In this study, we demonstrate the single mode lasing by using femtosecond optical pumping pulses from precisely engineered monolithic VCSELs having dense ultrathin films of CQW monolayers obtained by self-assembly technique. Here, mode tuning films present in the monolithic VCSELs were designed to achieve high Q-factors as well as accurate spectral matching with the gain spectrum with the help of full electromagnetic simulations. Owing to the uniformity and thermal stability of our ultrathin CQW films used in these devices, the top DBRs could be deposited at temperatures above 150 °C, which ensures the fabrication of high Q-factor VCSELs. We observed a lasing threshold of 112 μJ cm$^{-2}$ owing to the resulting high Q-factor of ≈760 for a device having only 7 nm thick CQW layer, which is the thinnest lasing demonstration among all colloidal NCs reported to date to the best of our knowledge. We were also able to deposit multiple CQW monolayers uniformly in a cavity using self-assembly method and demonstrated lasing with a lower threshold of 78 μJ cm$^{-2}$ from a VCSEL with four CQW monolayers having total thickness of ≈28 nm. It is worth noting that the CQW films in our devices were as thin as those used for electrical operation of colloidal light-emitting diodes (LEDs). Further cavity optimization in addition to additional improvement in CQW performance may enable the realization of electrically-driven CQW lasers, which is a long-term goal.

2. Results and Discussion

CdSe/Cd$_{34}$Zn$_{66}$S core/giant alloyed hot-injection (HI) shell CQWs have been synthesized according to our recent recipe based on well-adjusted precursor and ligand concentrations. To produce highly emissive, spectrally narrow and environmentally stable near-unity efficiency core/shell CQWs, we have grown giant shell on seed CdSe CQWs by the HI technique. These novel HI-grown shell CQWs were shown to exhibit near-unity photoluminescence quantum yield (PL-QY) and high photo- and thermal-stability, and the device performance of these materials that have been recently reported with record high external quantum efficiency in LEDs and high optical gain and ultralow lasing thresholds. Figure 1a presents PL and absorption spectra of our CQWs. The inset of Figure 1a shows a representative transmission electron microscopy (TEM) image of these CQWs. The PL-QY of our CQWs used in this work was measured to be 97%. The two excitonic features in the absorbance spectra located at 632 and 574 nm correspond to the electron-heavy hole and electron-light hole transitions, respectively. The PL emission having a full-width at half-maximum (FWHM) of 25 nm peaks at 640 nm. The TEM image reveals that these CQWs have the average dimensions of (l × w × h) of 17.6 ± 1.6 × 16.9 ± 1.2 × 4.6 ± 0.5 nm$^3$, and their total thickness is 7 nm if the thickness of the capping ligands are taken into account.

Time-resolved PL (TRPL) data of the synthesized CdSe/Cd$_{34}$Zn$_{66}$S core/giant alloyed HI-shell CQWs are given in Figure 1b. The TRPL data were fitted with biexponential decay function and the resulting amplitude average lifetime is 24 ns. The atomic percentages of the Cd and Zn in the shell are found to be 34 and 66, respectively, from X-ray photoelectron spectra (XPS) of the CQWs that is presented in Figure 1c (for the calculation procedure, see the Supporting Information).

The optical gain properties of these CQWs were studied in our previous work, in which we showed the capability of amplified spontaneous emission from bi-CQW layers with a thickness of only 14 nm. This observation shows the possibility of lasing action from ultrathin CQW gain media that coupled into an effective cavity. In this work, we fabricated CQW-VCSELs with ultrathin CQW gain media and investigated their lasing characteristics. Here each device consists of a bottom distributed Bragg reflector (DBR) constructed with alternating layers of SiO$_2$ and TiO$_2$, a cavity formed by single or multiple layers of CQWs, a mode-tuning dielectric layer, and a top DBR containing SiO$_2$ and TiO$_2$ layers.

To achieve lasing in the cavity, the cavity resonant mode $\lambda_{res}$ should be spectrally aligned with the peak of CQW gain $\lambda_{max}$. Also, the optical field overlap $\Gamma$ with the gain medium should be maximized in order to achieve sufficient net modal gain given by $g_{modal} = \Gamma g_{material} - \sigma$, where $g_{material}$ is the material gain and $\sigma$ is the total modal loss, including self-absorption, local heating due to excitation and scattering resulting from roughness in our deposited alumina and DBR layers. The $\Gamma$-factor, and Q-factor as well as spectral behavior of the cavity were studied via numerical simulations based on finite-difference time-domain (FDTD) method (see the Experimental Section). All device parameters were carefully analyzed and optimized prior to the device fabrication for achieving the best possible performance in our numerical calculations. As shown in Figure 2a, $\lambda_{res}$ can be modulated...
Figure 1. a) Photoluminescence (PL) and absorption spectra of the CdSe/Cd$_{0.34}$Zn$_{0.66}$S core/giant alloyed HI shell-grown CQWs. Inset: High-resolution transmission electron micrograph of these CQWs (scale bar: 30 nm) b) time-resolved PL spectra (in solution), c) X-ray photoelectron spectra (XPS) of CdSe/Cd$_{0.34}$Zn$_{0.66}$S core/giant alloyed HI shell CQWs.

Figure 2. Product of Q-factor and Γ-factor and the corresponding resonant wavelength of our cavity a) as a function of the CQW film thickness, b) Q-factor and Γ-factor of the cavity separately for a monolayer of CQWs as a function of the mode tuning film (silica, alumina, and titania) thickness. c) The product of Q-factor and Γ-factor and the corresponding resonant wavelength of the cavity for a monolayer of CQWs as a function of the mode-tuning film (silica, alumina, and titania) thickness. d) Electric field intensity profile of the resonant mode inside a cavity with a monolayer of CQWs and 30 nm of alumina. e) Magnified plot of main lobe of the electric field intensity inside the cavity covering CQW and Al$_2$O$_3$ layers at the center shown in d).
continuously from $\approx 600$ to $\approx 680$ nm by increasing the thicknesses of the CQW films from a single monolayer up to 10 monolayers. This is the result of phase shift imparted by increasing of the cavity thickness. Adding a monolayer of CQWs with a thickness of $\Delta t = 7$ nm to the cavity redshifts the resonance wavelength by a relatively large amount of $\Delta \lambda_{\text{res}} = +13$ nm ($\Delta v_{\text{res}} = -11$ THz) at $v_{\text{res}} = 493$ THz. The shift $\Delta v_{\text{res}} = -v_{\text{res}} \Delta t/t_{\text{eff}}$ is proportional to effective cavity length $t_{\text{eff}}$, which is found to be $\approx 315$ nm (Figure 2e). The $\Gamma$ factor increases by the thickness of CQW film, however the shift of the cavity mode to the edge of DBR stop-band reduces the reflection of the DBR and subsequently the $Q$-factor of the cavity. For our DBR design, the $\Gamma \times Q$ product reaches to a peak value at about 640 nm and reduces when the $\lambda_{\text{res}}$ is further increased by increasing the CQW film thickness.

Considering the ultrathin cavity thickness and resulting large free spectral range, $\lambda_{\text{res}}$ needs to be tuned to reside in the FWHM of CQW emission. To work around this problem, additional dielectric layers were incorporated between the mirrors, which provides us with the ability to tune $\lambda_{\text{res}}$ while maintaining a high $\Gamma \times Q$ product at the desired wavelength. We studied the effect of adding an extra layer of SiO$_2$, Al$_2$O$_3$, and TiO$_2$ inside the cavity containing different numbers of CQW layers. As shown in Figure 2b (top panel), $\Gamma$ factor decreases as the thickness and refractive index of the mode-tuning layer increases. Increasing the thickness broadens the mode spatially and increasing the refractive index localizes the mode peak away from CQW film, and hence, reduces the effective mode overlap. The peak of $Q$-factor is related to the refractive index of the material, which happens at thinner films for the higher index. Figure 2c shows the calculated $\Gamma \times Q$ product for a cavity with a monolayer of CQWs using different mode-tuning materials. These calculations reveal that Al$_2$O$_3$ provides the best performance and the highest $\Gamma \times Q$.

Based on our computational results, we chose to deposit 30 nm of Al$_2$O$_3$ on top of a monolayer of CQWs to match $\lambda_{\text{res}}$ with the peak emission of our CQWs. The simulated $Q$-factor of $\approx 9000$ in this case shows a great potential for coupling the emission from CQW layer to the nanocavity, which can enhance the SE due to the PE. Figure 2d shows the electric field intensity of the resonant mode at 639 nm for the designed monolayer CQW cavity. Figure 2e shows one period of this mode with a length of 340 nm, which indicates an effective mode index of 1.88.

We fabricated four sets of VCSELs having one to four CQW monolayers in the cavity. To tune the resonant mode of each VCSEL within the FWHM of the CQW emission, an extra layer of Al$_2$O$_3$, which the thickness is optimized according to the number of CQW layers, was deposited. The CQW monolayers were deposited through our liquid–air interface CQW self-assembly method, which offers us the capability of depositing one CQW layer at a time with near-unity surface coverage and sub-nm surface roughness. The mode-tuning Al$_2$O$_3$ layer was laid down through atomic layer deposition at 150 °C. As the CQWs used in

Figure 3. a) Cross-section TEM image of CQW-VCSEL (top panel) and magnified image of a cavity (bottom panel) with a 7 nm thick (a monolayer-thick) CQW layer. b) Top-view SEM image of a self-assembled CQW monolayer. c) Simulated cavity transmission for CQW-VCSEL1, 2, 3, and 4, which possesses 7, 14, 21, and 28 nm thick CQW layers and exhibit cavity modes at 639, 649, 637, and 647 nm, respectively. The inset shows the magnified range containing the resonant modes. d) Spectra measured at a pumping energy density of 119 $\mu$J cm$^{-2}$ for CQW VCSELs (left panel) and design expectations from FDTD simulations (right panel).
this study have oleic acid ligand and are thermally stable up to 200 °C, maintaining their PL reversibly.[24] It is possible to use the high-temperature deposition techniques on top of the CQWs to obtain uniform film quality in top layers.

Figure 3a shows the cross-sectional TEM image of one such VCSEL (top panel) and a magnified image of the cavity consisting of one monolayer of CQWs and Al₂O₃ mode-tuning layer (bottom panel). As shown in this image, the film in the fabricated device is highly uniform resulting in high cavity Q-factors. The scanning electron microscopy (SEM) image of a monolayer of self-assembled CQWs is shown in Figure 3b, which exhibits near-unity surface coverage. Figure 3c depicts the simulated transmission spectrum for the four fabricated CQW-VCSELs. According to our simulations, a cavity mode exists within a wide range of photonic bandgap in our designed structure. Hence, the cavity modes of all devices were successfully tuned to reside within the gain band of the CQWs. For a cavity with a monolayer of CQWs, adding 30 nm thick Al₂O₃ film causes a redshift in the mode from 595 to 639 nm. Similarly, for a cavity with 2 monolayers of CQWs, adding 30 nm of Al₂O₃ redshifts the mode from 608 to 649 nm. With the addition of 10 nm of Al₂O₃, the mode in the cavity having three monolayers of CQWs redshifts from 620 to 638 nm and, similarly, the mode redshifts from 632 to 647 nm in the case of 4 monolayers of CQWs.

Figure 3d shows the measured PL spectra of the fabricated CQW-VCSELs at a pump fluence of ≈119 µJ cm⁻² together with the simulated results. The experimentally observed lasing peak display an excellent match with our expectation from the numerical simulations. To find out the lasing threshold in each VCSEL, the cavities were excited with a pulsed laser at 400 nm with a pulse width of ≈110 fs and a repetition rate of 1 kHz, by varying the pump fluence (see the Experimental Section). The measured spectra from CQW-VCSELs having one to four monolayers of CQWs are presented in Figure 4a–d. Because of the ultrathin cavity and high Q-factor in all cases, a single longitudinal mode exists in all devices. The linewidths extracted from Gaussian fits for the measured lasing spectra are plotted as a function of the pump fluence in Figure 4e–h (bottom panel). The onset of lasing is marked by a sharp increase in the PL intensity at the lasing threshold. By fitting the data with a linear function, we obtained a threshold pump energy density of ≈112 µJ cm⁻² for the VCSEL having a monolayer of CQW layer (VCSEL1). To understand the threshold behavior of our CQW-VCSELs, we should consider the modal gain relationship \( g_{\text{modal}} = \Gamma g_{\text{material}} - \alpha \). When the thickness of the gain medium increases, the overlap increases linearly, which should result in a linear reduction of the threshold provided that the gain is constant over the spectrum. However, VCSELs have different resonance wavelengths having different levels of gain. As a result, the threshold does not reduce linearly. For example, in the case of VCSEL2 the threshold reduces down to about 107 µJ cm⁻², and 102 µJ cm⁻² for VCSEL3, while VCSEL4 exhibits a sharp reduction to 78 µJ cm⁻². The reason is that VCSEL3 has a resonant mode at shorter wavelength, at which it suffers from low gain and in spite of higher modal overlaps, its threshold reduction is not manifested as expected. The cavity linewidth of VCSEL1 narrows down to 0.8 nm at the lasing threshold and reaches a minimum value of 0.5 nm at the fluence of 114 µJ cm⁻². Then, the linewidth gradually increases as we further keep increasing the pump fluence. For example, the linewidth widens to ≈0.6 nm at the pump fluence of 119 µJ cm⁻². This broadening at high pump fluences can be attributed to the temperature fluctuation and/or refractive index modulation due to nonlinear effects under hard pumping, which may result in the cavity mode drift near the cavity resonance wavelength. Similar behavior has been observed for VCSEL2-4. The peak wavelengths extracted from Gaussian fits for the measured lasing spectra are shown in Figure 4e–h (bottom panel). For all VCSELs the peak wavelength deviation with increasing pump intensity is less than 1 nm, which shows excellent stability in emission wavelength. The small amount of deviation can be resulting from the refractive index modulation due to nonlinear effects, heating effects and noise. The lasing characterizations of the VCSELs are listed in Table 1 for all CQW film thicknesses studied. The VCSEL1 with only one monolayer of CQWs was not as stable as our
previous demonstration using thick samples,[17,24] under femtosecond excitation, and its emission endured only for several seconds. In the case of VCSEL4, the operation lifetime increased to about a minute.

The observed reduction in the lasing threshold with the increasing thickness of gain medium stems from the increased resonant mode overlap with the gain medium. Furthermore, in each of our VCSELs, most of the pump beam (more than 90%) was not absorbed by the ultrathin CQW film, but rather was transmitted through or reflected off the device, which would also require higher pumping energy. By simply increasing the CQW film thickness, in addition to Γ-factor enhancement, the pump beam absorption in the gain medium could be increased, thereby increasing the modal gain at the same pump power. All CQW-VCSELs exhibited relatively low lasing thresholds, thereby increasing the modal gain at the same pump power. All CQW-VCSELs exhibited relatively low lasing thresholds on the order of tens of µJ·cm⁻² which can be attributed to the spectral matching of the cavity mode and the maximized Γ- and Q-factors. Furthermore, a key factor for this high performance is the exploitation of H1 shell growth of CQWs, which simultaneously enables a near-unity PL-QY, reduces non-radiative channels, ensures smooth films, and enhances the stability.

3. Conclusion

In this study, we fabricated monolithic CQW-VCSELs and demonstrated single-mode lasing using densely-packed self-assembled films of CdSe/Cd₉₀.₃₄Zn₀.₆₆S core/giant alloyed H1-shelled CQWs as thin as 4.6 nm (if 2.4 nm organic ligands are not taken into account). Microcavities with the highest possible Q-factors as well as accurate spectral matching with the gain spectra were designed by employing numerical modeling. The high packing density of smooth films of the self-assembled CQWs, the high Q-factor of the cavity and the superior optical properties of core/shell CQWs synthesized with our hot injection technique have near-unity PL-QY enable low threshold single-mode lasing from ultrathin films of these CQWs. The demonstrated CQW films used in this work are sufficiently thin to possibly enable electrical carrier injection, given that they are combined with proper carrier-transporting layers. The findings of this work may pave the way for realization of ultrathin electrically-driven colloidal laser devices, providing critical advantages including single-mode lasing and high electrical conduction.

Supporting Information

Supporting Information is available from the Wiley Online Library or from the author.

Table 1. Lasing parameters of our CQW-VCSELs.

| VCSEL | Number of CQW layers | CQW film thickness [nm] | Al₂O₃ film thickness [nm] | Lasing threshold \(\phi_{th} \) [µJ·cm⁻²] | Lasing wavelength \(\lambda_{res} \) [nm] |
|-------|----------------------|-------------------------|--------------------------|-----------------------------------|-------------------------|
| 1     | 1                    | 7                       | 30                       | 112                               | 640                     |
| 2     | 2                    | 14                      | 30                       | 107                               | 649                     |
| 3     | 3                    | 21                      | 10                       | 102                               | 638                     |
| 4     | 4                    | 28                      | 10                       | 78                                | 648                     |

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Conflict of Interest

The authors declare no conflict of interest.

Keywords

colloidal quantum wells, liquid interface self-assembly, monolithic microcavity, single-mode lasing, vertical cavity surface-emitting laser

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[13] B. Guzelturk, M. Pelton, M. Olutas, H. V. Demir, \textit{Nano Lett.} \textbf{2019}, \textit{19}, 277.

[14] M. Li, M. Zhi, H. Zhu, W. Y. Wu, Q. H. Xu, M. H. Jhon, Y. Chan, \textit{Nat. Commun.} \textbf{2015}, \textit{6}, 8513.

[15] E. Lhuillier, S. Pedetti, S. Ithurria, B. Nadal, H. Heuclin, B. Dubertret, \textit{Acc. Chem. Res.} \textbf{2015}, \textit{48}, 22.

[16] E. Baghani, S. K. Oleary, I. Fedin, D. V. Talapin, M. Pelton, \textit{J. Phys. Chem. Lett.} \textbf{2015}, \textit{6}, 1032.

[17] Y. Altintas, K. Gungor, Y. Gao, M. Sak, U. Quiliyeva, G. Bippi, E. Mullet, E. H. Sargent, H. V. Demir, \textit{ACS Nano} \textbf{2019}, \textit{13}, 10662.

[18] N. Taghipour, S. Delikanli, S. Shendre, M. Sak, M. Li, F. Isik, I. Tanriover, B. Guzelturk, T. C. Sum, H. V. Demir, \textit{Nat. Commun.} \textbf{2019}, \textit{11}, 3305.

[19] B. Guzelturk, Y. Kelestemur, M. Olutas, Q. Li, T. Lian, H. V. Demir, \textit{J. Phys. Chem. Lett.} \textbf{2017}, \textit{8}, 5317.

[20] S. Gupta, E. Waks, \textit{Opt. Express} \textbf{2014}, \textit{22}, 3013.

[21] C. Dang, J. Lee, C. Breen, J. S. Steckel, S. Coe-Sullivan, A. Nurmiak, \textit{Nat. Nanotechnol.} \textbf{2012}, \textit{7}, 335.

[22] O. Erdem, S. Foroutan, N. Ghashghaii, B. Guzelturk, Y. Altintas, H. V. Demir, \textit{Nano Lett.} \textbf{2020}, \textit{20}, 6459.

[23] S. Foroutan-Barenji, O. Erdem, N. Ghashghaii, Y. Altintas, H. V. Demir, \textit{Small} \textbf{2020}, \textit{16}, 2004304.

[24] Y. Altintas, U. Quiliyeva, K. Gungor, O. Erdem, Y. Kelestemur, E. Mullet, M. V. Kovalenko, H. V. Demir, \textit{Small} \textbf{2019}, \textit{15}, 1804854.

[25] J. Roh, Y. S. Park, J. Lim, V. I. Klimek, \textit{Nat. Commun.} \textbf{2020}, \textit{11}, 271.

[26] H. Kim, K. S. Cho, H. Jeong, J. Kim, C. W. Lee, W. K. Koh, Y. G. Roh, S. W. Hwang, Y. Park, \textit{ACS Photonics} \textbf{2016}, \textit{3}, 1536.

[27] K. S. Cho, E. K. Lee, W. J. Joo, E. Jang, T. H. Kim, S. J. Lee, S. J. Kwon, J. Y. Han, B. K. Kim, B. L. Choi, J. M. Kim, \textit{Nat. Photonics} \textbf{2009}, \textit{3}, 341.

[28] T. H. Kim, K. S. Cho, E. K. Lee, S. J. Lee, J. Chae, J. W. Kim, D. H. Kim, J. Y. Kwon, G. Amarathunga, S. Y. Lee, B. L. Choi, Y. Kuk, J. M. Kim, K. Kim, \textit{Nat. Photonics} \textbf{2011}, \textit{5}, 176.

[29] A. A. Rossinelli, A. Riedinger, P. Marqués-Gallego, P. N. Knüsel, F. V. Antolínez, D. J. Norris, \textit{Chem. Commun.} \textbf{2017}, \textit{53}, 9938.

[30] B. Liu, Y. Altintas, L. Wang, S. Shendre, M. Sharma, H. Sun, E. Mullet, H. V. Demir, \textit{Adv. Mater.} \textbf{2020}, \textit{32}, 1905824.

[31] A. A. Rossinelli, H. Rojo, A. S. Mule, M. Aellen, A. Cicioni, E. De Leo, R. Schäublin, D. J. Norris, \textit{Chem. Mater.} \textbf{2019}, \textit{31}, 9567.

[32] Y. Kelestemur, Y. Shynkarenko, M. Anni, S. Yakunin, M. L. De Giorgi, M. V. Kovalenko, \textit{ACS Nano} \textbf{2019}, \textit{13}, 13899.