

Molecular Systems Design & Engineering

Building and designing systems from the molecular level

rsc.li/molecular-engineering



ISSN 2058-9689



PAPER

Alan E. Rowan, Anja R. A. Palmans, E. W. Meijer *et al.*
Catalytic single-chain polymeric nanoparticles at work: from ensemble towards single-particle kinetics

IChemE ADVANCING
CHEMICAL
ENGINEERING
WORLDWIDE



Cite this: *Mol. Syst. Des. Eng.*, 2018, 3, 609

Catalytic single-chain polymeric nanoparticles at work: from ensemble towards single-particle kinetics†

Yiliu Liu,^a Petri Turunen,^{bc} Bas F. M. de Waal,^a Kerstin G. Blank,^{id}^{bd} Alan E. Rowan,^{*bc} Anja R. A. Palmans^{id}^{*a} and E. W. Meijer^{id}^{*a}

Folding a single polymer chain around catalytically active sites to construct catalytic single chain polymeric nanoparticles (SCPNS) is a novel approach to mimic the activity and selectivity of enzymes. In order to relate the efficiency of SCPNs to their three-dimensional structure, a better understanding of their catalytic activity at an individual level, rather than at an ensemble level, is highly desirable. In this work, we present the design and preparation of catalytic SCPNs and a family of fluorogenic substrates, their characterization at the ensemble level as well as our progress towards analyzing individual SCPNs with single-molecule fluorescence microscopy (SMFM). Firstly, organocopper-based SCPNs together with rhodamine-based fluorogenic substrates were designed and synthesized. The SCPNs catalyze the carbamate cleavage reaction of mono-protected rhodamines, with the dimethylpropargyloxycarbonyl protecting group being cleaved most efficiently. A systematic study focusing on the conditions during catalysis revealed that the ligand acceleration effect as well as the accumulation of substrates and catalytically active sites in SCPNs significantly promote their catalytic performance. Secondly, a streptavidin–biotin based strategy was developed to immobilize the catalytic SCPNs on the surface of glass coverslips. Fluorescence correlation spectroscopy experiments confirmed that the SCPNs remained catalytically active after surface immobilization. Finally, single-SCPNS activity measurements were performed. The results qualitatively indicated that fluorescent product molecules were formed as a result of the catalytic reaction and that individual fluorescent product molecules could be detected. So far, no evidence for strongly different behaviors has been observed when comparing individual SCPNs.

Received 7th April 2018,
Accepted 24th May 2018

DOI: 10.1039/c8me00017d

rsc.li/molecular-engineering

Design, System, Application

A new approach to polymer-based catalytic systems has been recently disclosed, which is based on the folding of a polymer chain around catalytically active sites covalently bound to the polymer. In contrast to enzymes, these polymer-based catalysts are structurally heterogeneous, so that the catalytically active sites may reside in different micro-environments. As a result, differences in reactivity may arise due to this diversity of catalytic sites. It is therefore of great interest to investigate this possible diversity and we here design a system to study the catalytic activity at the single-polymer level. In the design, the single-chain polymer nanoparticles are connected to the surface, and substrates are transformed from a luminescent, silent state to a strongly fluorescent state. With these polymers and substrates synthesized it has become possible to study the catalytic activity of individual chains with single-molecule fluorescence microscopy. The new insights obtained in this work can be used to progress the search to an enzyme-mimic for bio-orthogonal chemistry in complex cellular media.

Introduction

Folding individual synthetic polymer chains into nanoparticles, which is reminiscent of nature's way of folding polypeptides into proteins, has emerged as a novel approach toward attaining defined polymer architectures.^{1–6} In general, these polymers are decorated with pendant groups that are capable of forming covalent or supramolecular bonds. After intra-chain (supramolecular) crosslinking between the pendant groups, the individual polymer chains are folded into single-chain polymeric nanoparticles (SCPNS). The incorporation of

^a Institute for Complex Molecular Systems, Laboratory of Macromolecular and Organic Chemistry, Eindhoven University of Technology, P.O. Box 513, 5600 MB, Eindhoven, The Netherlands. E-mail: a.palmans@tue.nl, e.w.meijer@tue.nl

^b Department of Molecular Materials, Institute for Molecules and Materials, Radboud University, 6525 AJ, Nijmegen, The Netherlands. E-mail: alan.rowan@uq.edu.au

^c The University of Queensland, Australian Institute for Bioengineering and Nanotechnology, Brisbane, Queensland 4072, Australia

^d Mechano(bio)chemistry, Max Planck Institute of Colloids and Interfaces, Potsdam-Golm Science Park, 14424 Potsdam, Germany

† Electronic supplementary information (ESI) available. See DOI: 10.1039/c8me00017d



substrates with different functional groups suitable for ensemble and SMFM experiments; 3) immobilization of active SCPNs on a surface, required for the SMFM experiments; 4) establishing a single-molecule experimental protocol and data analysis.

Results and discussion

Design and preparation of catalytic SCPNs

Organocopper-based SCPNs were chosen owing to their efficient catalytic performance in a broad range of reactions. Three ligand-containing polymers (P1–P3) and a catalytically inactive control polymer (P4) were designed (Fig. 1, Table 1). These polymers contain water-soluble polyether side-chains and BTA pendants, which enable them to form SCPNs in aqueous solution. P1–P3 were modified with phenanthroline ligands, which can bind Cu ions to form catalytically active sites, while the control polymer P4 has dodecyl chains instead. To allow surface immobilization, P2 and P3 were decorated with 1% of biotin-containing side chains. Moreover, P3 was labeled with the fluorescent dye Alexa Fluor® 488 to enable its visualization with fluorescence microscopy. The polymers were modularly synthesized using a strategy of post-polymerization modification.^{46,47} The precursor polymer poly(pentafluorophenyl acrylate) (pPFPA, M_n , SEC = 18.0 kD, $D = 1.28$, DP = 120) was synthesized by reversible addition-fragmentation chain-transfer (RAFT) polymerization and the thiocarbonyl end group was removed upon heating with excess azobisisobutyronitrile (AIBN) and lauroyl peroxide

(LPO) (Fig. S1†). P1–P4 were prepared *via* polymer analogous reactions of pPFPA by sequential addition of amines (Fig. S2–S4†). 5-Amino-*N*-(1,10-phenanthroline-5-yl)pentanamide (phen-C4-amine) and the enantiomerically pure, amino-functionalized BTA unit (BTA-amine) were synthesized, the other amines were commercially obtained. After full modification, the final polymers were obtained *via* dialysis to remove excess amines and byproducts.

Next, the polymers were complexed with Cu²⁺ ions. For this purpose, the polymers were dissolved in water, sonicated and subjected to a heating-cooling procedure. Then CuSO₄ was added to the solution. Taking P1 as an example, its folding behavior, complexation with Cu²⁺ ions (P1@Cu(II)) and SCPN formation were characterized using a combination of techniques. The binding of Cu²⁺ ions to the phenanthroline ligand was studied with UV-vis spectroscopy. As shown in Fig. 2a, the phenanthroline ligand shows a characteristic absorption band around 270 nm. Upon addition of CuSO₄ the absorption undergoes a red-shift to around 280 nm. A detailed plot of Cu²⁺ ratio *versus* shift in absorption reveals a 2 : 1 binding stoichiometry between the phenanthroline ligand and Cu²⁺ ions. In the CD spectra of P1, a negative Cotton effect was present at 223 nm before and after Cu²⁺ complexation without a change in value. Moreover, the CD temperature scans of P1 with or without CuSO₄ overlap (Fig. S5†). These observations indicate that (i) the BTA pendants form helically stacked dynamic aggregates driving the folding of polymer chains and (ii) Cu²⁺ complexation is not interfering with BTA aggregation.

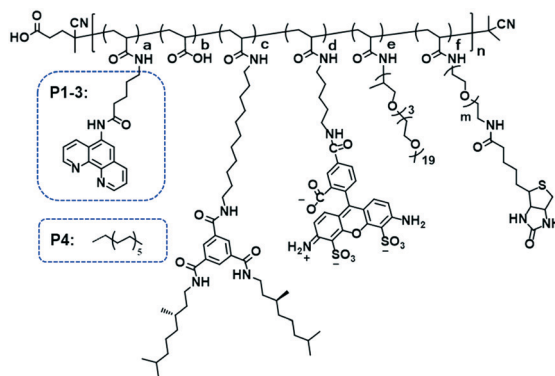


Fig. 1 Chemical structure of polymer P1–4. P1–P3 were modified with phenanthroline ligands, while the control polymer P4 has dodecyl chains instead. To allow surface immobilization, P2 and P3 were decorated with 1% of biotin-containing side chains. Moreover, P3 was labeled with the fluorescent dye Alexa Fluor® 488 to enable its visualization with fluorescence microscopy.

Table 1 Composition and SEC characterization of polymers P1–P4^a

Polymer	<i>a</i>	<i>b</i>	<i>c</i>	<i>d</i>	<i>e</i>	<i>f</i>	<i>n</i>	<i>m</i>	M_n (kD)	<i>D</i>
P1	0.08	0.04	0.10	—	0.78	—	120	—	31.4	1.18
P2	0.08	0.08	0.10	—	0.73	0.01	120	45	28.0	1.30
P3	0.08	0.10	0.10	0.01	0.70	0.01	120	45	27.8	1.20
P4	0.08	—	0.10	—	0.82	—	120	—	31.7	1.20

^a The values for *a*–*f* are determined *via* ¹⁹F-NMR. M_n is measured by SEC in DMF with 10 mM LiBr, relative to poly(ethylene oxide) standards.



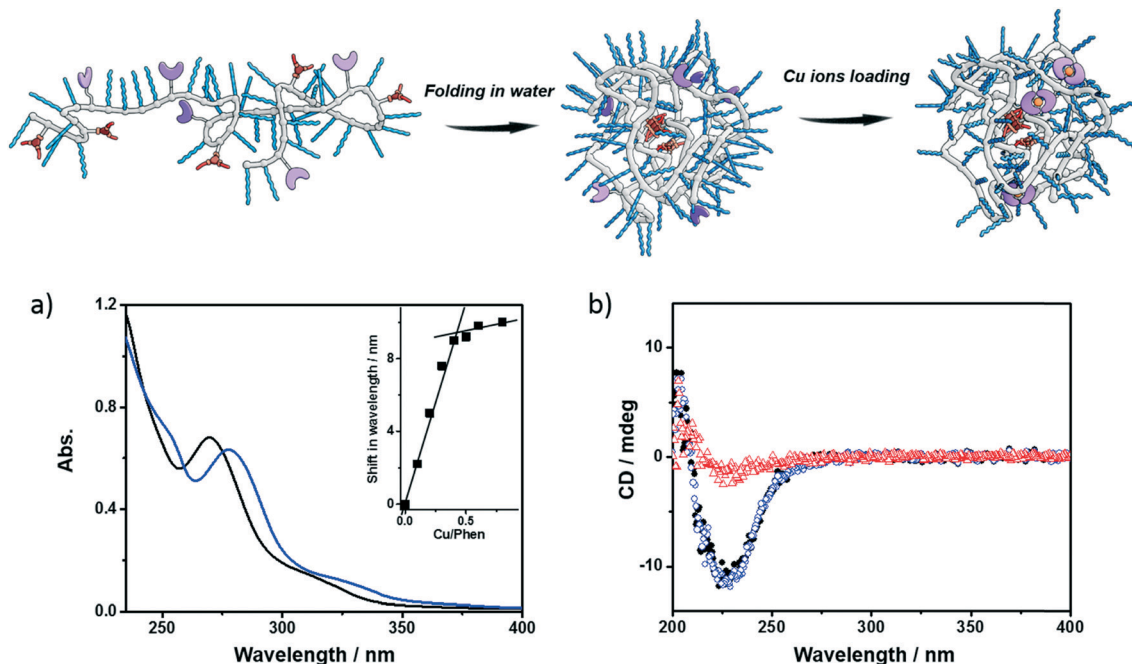


Fig. 2 Folding and Cu-loading of the polymers; a) UV-vis spectra of P1 (black, 0.6 mg mL^{-1}) and P1@Cu(II) (blue, 0.6 mg mL^{-1} , [phen] : [Cu] = 2 : 1); b) CD spectra of P1 (0.5 mg mL^{-1} , black dot: $20 \text{ }^\circ\text{C}$, red triangle: $90 \text{ }^\circ\text{C}$), P1@Cu(II) (blue square, $20 \text{ }^\circ\text{C}$, 0.5 mg mL^{-1} , [phen] : [Cu] = 2 : 1).

Small-angle X-ray scattering (SAXS) and dynamic light scattering (DLS) were employed to evaluate the size of the polymeric nanoparticles formed. SAXS measurements indicate that P1 and P1@Cu(II) form nanoparticles with a radius of gyration (R_g) of 5.6 and 5.4 nm, respectively (Fig. S6†). The DLS results of P1 and P1@Cu(II) show a monodisperse distribution of particles with a hydrodynamic radius (R_h) of 7.4 and 8.2 nm, respectively (Fig. S7†). The scattering results are comparable to similar systems reported previously, suggesting that single-polymer chains fold into SCPNs indeed. P2–P4 were characterized in the same way as P1 and similar results were obtained (Fig. S7, Table S1†). P3, which is modified with Alexa Fluor® 488, was further studied with fluorescence correlation spectroscopy (FCS). The FCS experiments were performed at a concentration of 100 nM of P3 in aqueous solution. The diffusion of the P3-based fluorescent nanoparticles was found to be approximately one order of magnitude slower than a reference fluorophore (ATTO488, Fig. S8†). This indicates that the Alexa Fluor® 488 dye was successfully coupled to the polymers. Using the Stokes–Einstein equation, the R_h of P3 was determined to be $8.3 \pm 0.8 \text{ nm}$, which is consistent with the light scattering results.

Design and synthesis of fluorogenic substrates

A series of fluorogenic substrates were designed that are predicted to be good substrates for the Cu-phenanthroline functionalized SCPNs. Considering that these substrates should be suitable for both ensemble and single-SCPn kinetic experiments, a number of criteria need to be considered: i) the substrates are stable in aqueous solution, but

can undergo fast cleavage in the presence of Cu(I)-based organometallic catalysts, catalysts made out of the Cu(II)-based precursor; ii) the substrates are based on a fluorophore with high brightness, so that a high signal-to-noise ratio can be achieved; iii) the substrates contain only one cleavable group, in order to avoid the formation of reaction intermediates with different fluorescent properties; and iv) the substrates are hydrophobic and are converted into hydrophilic products, which could facilitate the SCPN-based catalysis. With these guidelines in mind, substrates S1–S4 were designed and synthesized (Fig. 3). Rhodamine 110 (Rh110) was chosen as the fluorophore owing to its high brightness. However, it carries two amino groups for further modification. To prepare a substrate with a single reactive site, one amino group of Rh110 was first modified with a morpholine-carbonyl group to obtain MC-Rh110.^{48–50} The urea bond formed is not reactive and hydrolytically stable. At the same time, MC-Rh110 remains sufficiently bright for SMFM experiments.⁴⁸ Further modification of the remaining amino group to generate carbamate bonds yields substrates S1–S4, which contain different alkyne or alkene groups. The carbamate bond of S1 was previously shown to be hydrolyzed in the presence of Cu(I),⁵¹ while the carbamate bond in S2 is labile in the presence of Pd(II) catalysts.^{52,53} The hydrolytic stability of the carbamate bonds in S3 and S4 in the presence of Cu(I) is not known.

SCPn-based catalysis in aqueous solution

The catalytic performance of SCPNs in the carbamate-cleavage reaction was first studied at the ensemble level in



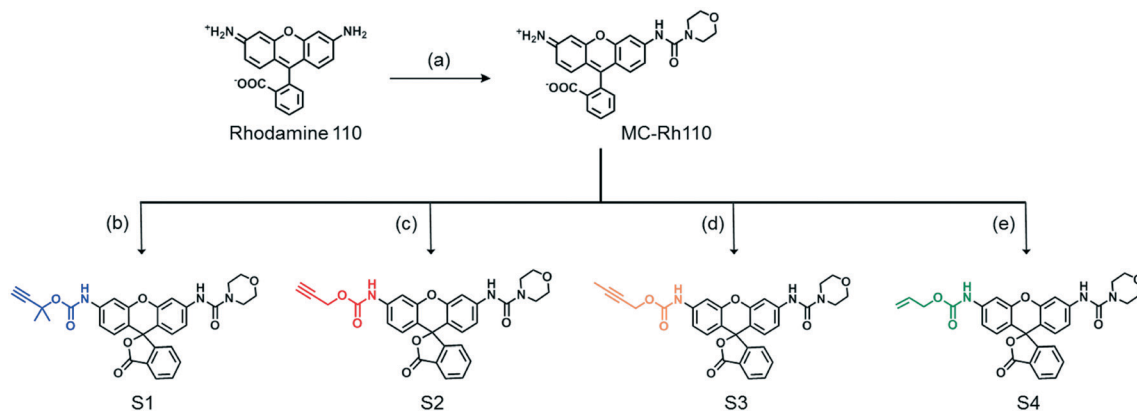


Fig. 3 Chemical structures and synthesis of the substrates **S1–S4**. (a) 4-Morpholinecarbonyl chloride; (b) triphosgene, 2-methylbut-3-yn-2-ol; (c) propargyl chloroformate; (d) triphosgene, but-2-yn-1-ol; (e) triphosgene, prop-2-en-1-ol.

aqueous solution. The progress of the reaction was monitored with fluorescence spectroscopy as well as liquid chromatography-mass spectroscopy (LC-MS) (Fig. S9 and S10†). Whereas the former provides information about the catalytic reaction in real time, LC-MS allows exact quantification of the substrate conversion at set time intervals. As a first step, a substrate activity screening was performed. **P1@Cu(I)** ([phen]:[Cu] = 2:1) was chosen as a representative catalyst. Generally, a stock solution of the substrate (in DMSO) was added into a **P1@Cu(II)** solution (2 μM) to reach

a final concentration of 30 μM . The reactions were initialized by adding sodium ascorbate (NaAsc, 1 mM) to generate **P1@Cu(I)** *in situ*. The fluorescence intensity was recorded at 520 nm for 30 min and substrate conversion was directly analyzed with LC-MS (Fig. S9b†). The fluorescence-time curves were normalized relative to the maximum conversion and are shown in Fig. 4. Substrates **S1** and **S2**, which contain end-functionalized alkyne moieties, are hydrolyzed, with the tertiary alkyne bearing **S1** showing the highest TOF ($\sim 200 \text{ h}^{-1}$). In contrast, **S3** and **S4**, which contain internal alkyne and

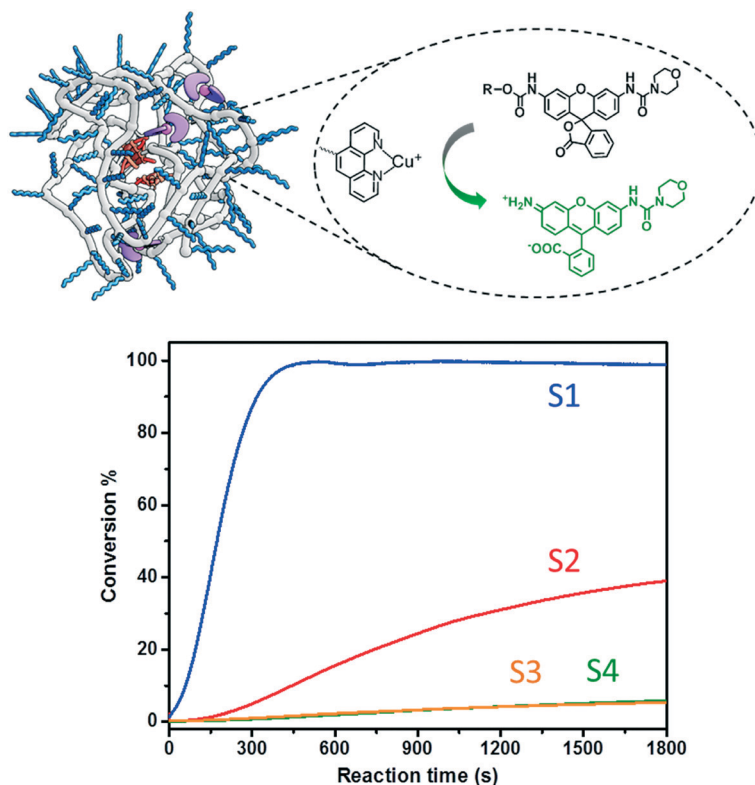


Fig. 4 Carbamate cleavage reaction of **S1–S4** catalyzed by **P1@Cu(I)** ([P1] = 2 μM , [CuSO₄] = 10 μM , NaAsc = 1 mM, substrate = 30 μM). The kinetic curves were recorded *via* fluorescence spectroscopy (ex. 495 nm; recorded at 520 nm) followed with normalization according to the conversions obtained from LC-MS.



alkene moieties, respectively, show almost no reactivity. It has been proposed that the formation of a Cu-acetylide intermediate is crucial in the catalytic mechanism. The terminal alkyne moieties of S1 and S2 bind Cu(I) to generate the Cu-acetylide. This leads to the formation of a Cu-stabilized propargyl cation thus cleaving the carbamate bond. The methyl groups of the tertiary propargyl moiety in S1 further stabilize the cation, thereby promoting the cleavage reaction.⁵¹

The ligand-containing SCPNs have several merits toward aqueous catalysis: α) ligand accelerated catalysis; β) hydrophobic interior; and γ) accumulation of catalytically active sites. Whereas the first feature is determined directly by the mechanism of the Cu-catalyzed reaction, the other two features lead to an enhancement of the local concentration of active sites and/or substrate molecules. To verify in how far these three features are responsible for enhancing the kinetics of the hydrolysis reaction, we performed systematic kinetic experiments on S1 and S2. As catalysts, we selected P1@Cu(I), which possesses contributions from α , β and γ , whereas P4 mixed with Phen@Cu(I) only possesses the α and β contribution. As reference Phen@Cu(I), P4 mixed with Cu(I) and only Cu(I) were used (Table 2). The outcome of these catalysis experiments is shown in Fig. 5 (and Fig. S9c†). The phenanthroline ligand indeed accelerates the Cu(I)-catalyzed carbamate-cleavage reaction. Moreover, the accumulation of catalytic active sites and substrates enhances the reaction to a large extent. In short, the more characteristics the system contains, the more effective the catalytic reaction becomes.

Table 2 Catalysts with different characteristics. α) ligand accelerated catalysis; β) hydrophobic interior; γ) accumulation of catalytically active sites

Entry (S1/S2)	Catalyst	Characteristics
1	P1@Cu(I)	α , β , γ
2	P4 & Phen@Cu(I)	α , β
3	Phen@Cu(I)	α
4	P4 & Cu(I)	β
5	Cu(I)	—

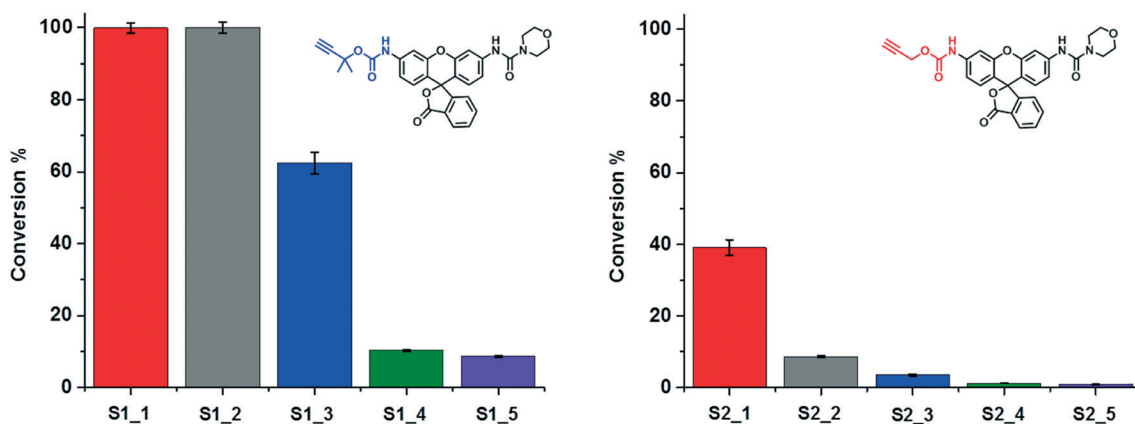


Fig. 5 A study on the characteristics of SCPN-based catalysis. The conversions of the reactions under different catalytic conditions (Table 2) were obtained through LC-MS.

Surface immobilization of catalytic SCPNs

Having obtained insights into the substrate specificity and the kinetics of the SCPN catalyzed reactions, a first step was taken toward single-SCPN experiments. For observing individual SCPNs with a confocal microscope, the SCPNs need to be immobilized on the surface of glass coverslips. We selected an immobilization strategy based on the well-established streptavidin–biotin interaction. In detail, the glass surface was first modified with an amino silane (3-aminopropyl dimethylethoxy silane). In the next step, a hetero-bifunctional poly(ethylene glycol) containing an amino-reactive NHS ester and a biotin group (NHS-PEG₃₀₀₀Da-biotin) was used to obtain a biotin covered surface. The biotin-containing SCPNs were then immobilized on the surface using streptavidin as a bridge (Fig. 6a). P3 was employed to test the SCPN immobilization protocol. P3 contains biotin motifs on its side-chains and is labeled with Alexa Fluor® 488, which allows its visualization with confocal microscopy. Fig. 6b shows confocal images obtained for different surfaces incubated with P3 at concentrations of 50 nM, 5 nM, and 0.5 nM, respectively. Inspection of Fig. 6b shows that the immobilization is successful, as indicated by the concentration-dependent density of SCPNs.

The question then arises whether the surface-immobilized SCPNs are still catalytically active. Using fluorescence correlation spectroscopy (FCS), the carbamate-cleavage reaction was monitored by following the increase in fluorescent products that are released from the immobilized SCPNs and diffuse into the bulk solution (see ESI† for details, Fig. S12). P2@Cu(II) was chosen for these experiments instead of P3@Cu(II) to avoid any influence from the Alexa Fluor® 488 dye used for labeling the SCPNs. The immobilization of P2@Cu(II) followed exactly the same protocol as shown for P3. A solution containing S1 (10 μ M) and NaAsc (1 mM) was added onto a surface with or without P2@Cu(II), respectively. As shown in Fig. 7, hydrolysis of S1 is clearly visible on the P2@Cu(II) covered surface. This provides first evidence that at least a fraction of SCPNs remains catalytically active after surface immobilization.



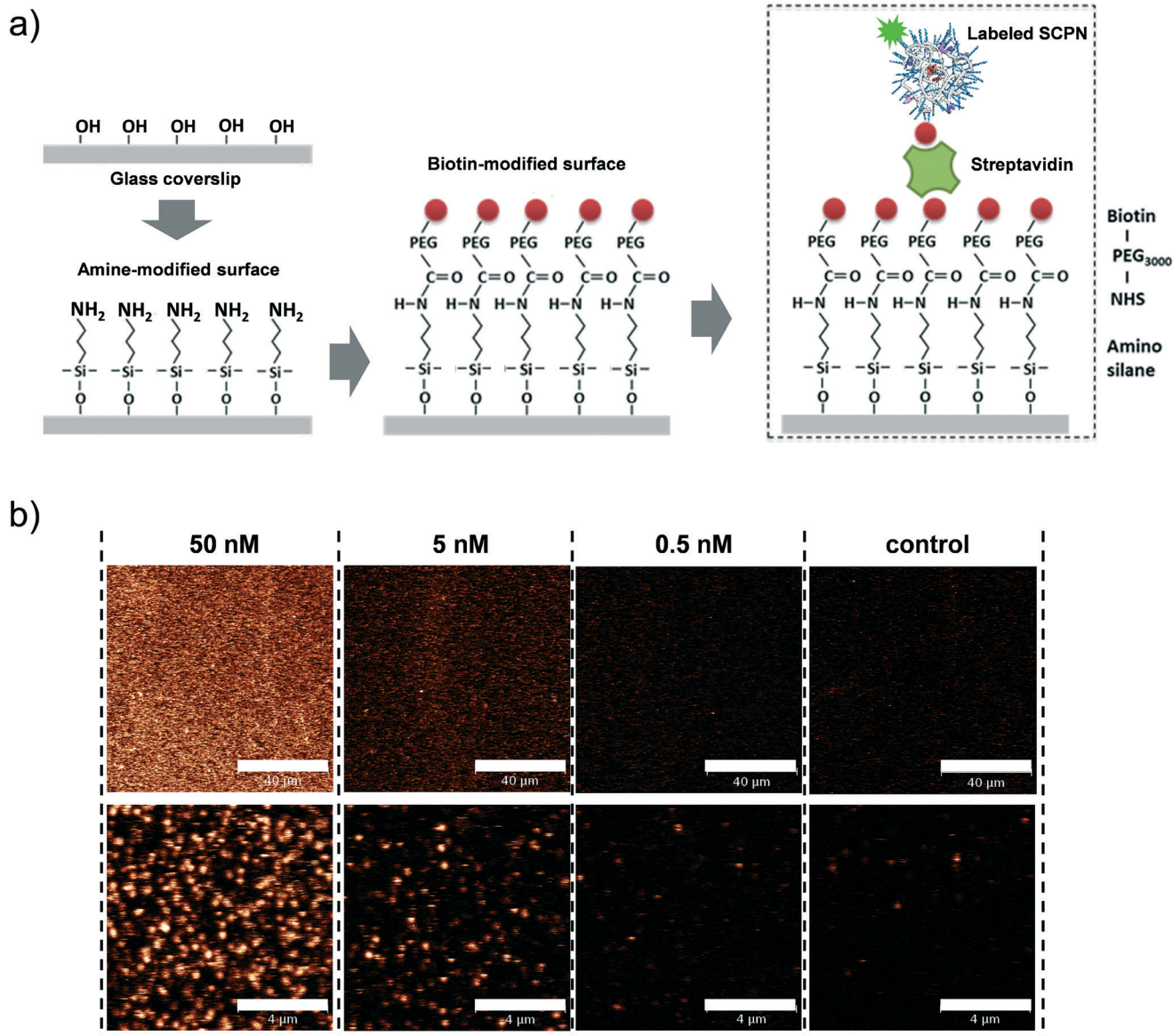


Fig. 6 Surface immobilization of biotinylated SCPNs. a) Experimental design. b) Confocal images showing biotinylated and Alexa Fluor® 488-labeled SCPN on a functionalized coverslip. The images show SCPNs immobilized in various concentrations and a corresponding control measurement where no SCPNs were added. The bar represents 40 and 4 μm in the top and bottom panel, respectively.

Single-molecule activity measurements

Single-SCP activity measurements were performed to find out if individual turnovers of a single SCPN can be followed over time. In these experiments, P3@Cu(II) was used in combination with S1 as the substrate. As the complex of the phenanthroline ligand with Cu ions quenches the fluorescence of Alexa Fluor® 488 (Fig. S11[†]), this fluorescent label is used indirectly to visualize Cu-containing SCPNs on the surface. Therefore, we developed the following protocol to localize the SCPNs on the coverslip surface. P3-based SCPNs (20 nM) were first immobilized on the glass coverslip. A surface area of $10 \times 10 \mu\text{m}$ was scanned before and after incubation with CuSO_4 solution (concentration 1 mM). Fluorescent spots

that disappeared after the incubation with CuSO_4 were considered as locations of Cu-loaded SCPNs (Fig. 8a). It is important to note that the disappearance of the fluorescent signal can also originate from bleaching of Alexa Fluor® 488 during consecutive scans. The degree of bleaching was tested and it was found to be negligible under the experimental conditions used (Fig. S13[†]).

Fluorescence time traces were recorded at the positions of different SCPNs for 1 minute each after delivering a solution containing 10 μM S1 and NaAsc. One representative time trace is shown in Fig. 8b. Despite the relatively low signal-to-noise ratio, which is expected for MC-Rh110, the results suggest that fluorescent product molecules are produced on the surface and can be detected. When comparing the time





Fig. 7 Fluorescence correlation spectroscopy (FCS) to determine product accumulation in solution. The graph illustrates the rate of product accumulation for the sample containing immobilized SCPNs (blue) and for a control sample without SCPNs (red, auto-hydrolysis).

traces recorded at the position of one SCPN with the control (empty area which does not contain SCPNs), a small increase in the frequency of fluorescence bursts (with high intensity) was observed, suggesting that the SCPNs are turning over the fluorogenic substrate. However, the number of clearly resolved turnovers detected appears to be low (Fig. 8b), which can be rationalized by the relatively low average TOF of the catalytic SCPN system. Nevertheless, the fact that no individ-

ual SCPNs with a significantly higher activity was observed (Fig. S14 and S15[†]), may imply that structural heterogeneities of the SCPNs do not cause a dramatic diversity in their catalytic performance.

Although a detailed quantitative kinetic analysis proved difficult, the results above provide a practical platform that holds potential for further optimization. Catalytically-active SCPNs with organometallic sites other than copper complexes, *e.g.* palladium or ruthenium complexes, are possible candidates, which may have higher TOFs and overcome the fluorescence quenching problem. Besides developing new catalysts, we are continuing our efforts on optimizing the experimental setup to obtain higher SNR and throughput. For example, the possibility of using a wide-field fluorescence setup combined with nano-phonic metal structures, such as zero-mode waveguides (ZMWs),^{54–57} is under exploration. Wide-field fluorescence microscopy, which allows for monitoring many individual SCPNs on the surface simultaneously, may provide insights into possible heterogeneities between individual SCPNs in a more straightforward manner. Meanwhile, ZMWs are able to reduce the detection volume drastically, thus allowing the use of higher substrate concentrations while reducing the background signal. In addition, their ability to enhance the fluorescence signal could increase the signal-to-noise ratio even further.

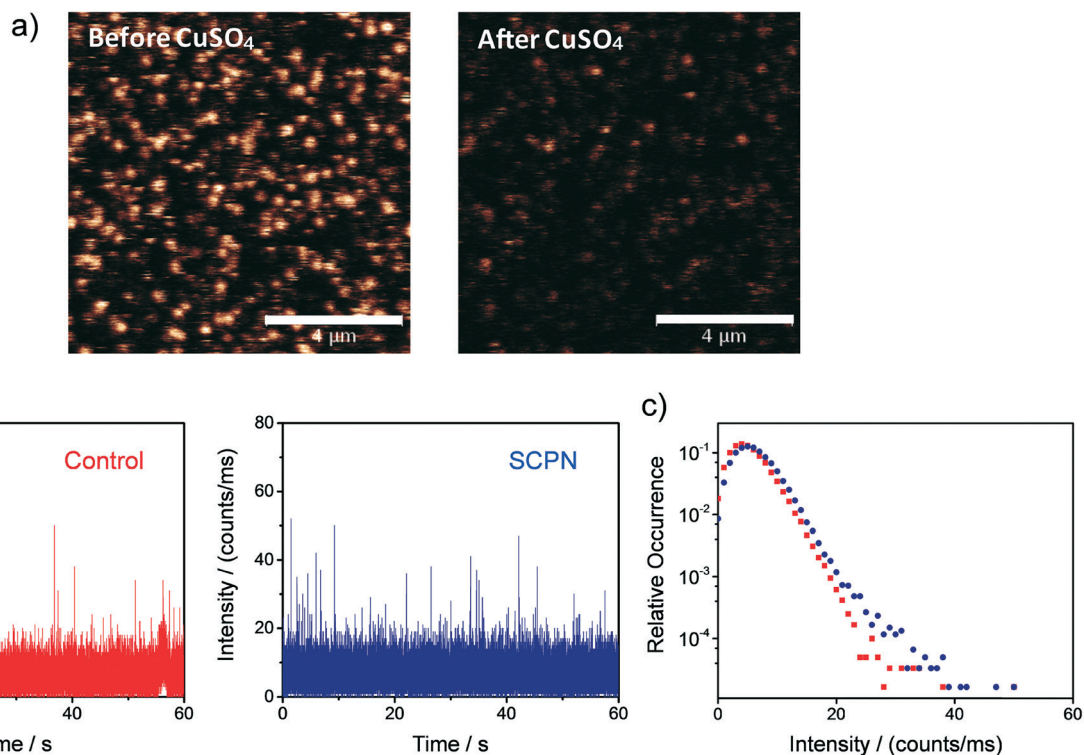


Fig. 8 Representative single-SCP activity measurement. a) Confocal images ($10 \times 10 \mu\text{m}$) showing the Alexa Fluor 488-labeled and biotin-functionalized SCPNs immobilized on a streptavidin-functionalized glass coverslip before (left) and after (right) the addition of CuSO_4 . The bar in the panel represents $4 \mu\text{m}$. b) Fluorescent intensity time traces and c) corresponding intensity distributions obtained when binning the data with 1 ms bin size. The data was recorded on the location of one individual SCPN (blue) and on an empty area on the surface (red; control).



Conclusion

We have shown the preparation of SCPNs that are capable of catalyzing fluorogenic reactions, as well as provide protocols for their surface immobilization and for monitoring their catalytic activity. The combined design of catalysts and substrates enabled us to obtain novel information about the substrate specificity of SCPNs and allowed to guide the activity of the catalytic reaction at the ensemble level. Similar to changes in reactivity as the result of mutations in enzymes, we expect that not every SCPN will be equally reactive.⁵⁸ Therefore, we are highly intrigued by the question on how to discriminate the diversity in reactivity at the individual level. The protocol for surface immobilization, which retains the catalytic activity of SCPNs, allows further SMFM measurements on individual SCPNs. In the course of our investigation, we identified the experimental challenges, mainly a low TOF of the catalysts combined with a low signal-to-noise ratio in SMFM measurements, which will allow us to improve the system in future experiments. So far, we have not found any evidence that some SCPNs are more reactive than the others. However, this work has laid a solid foundation for applying SMFM to further understanding the structure–function relation of catalytic SCPNs. We anticipate that these efforts will assist in directing the future of catalytic SCPNs, e.g. if sequence control in the polymer synthesis and regulation of polymer chain's precision folding are the gateway toward accessing enzyme-like activity and selectivity.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

Y. L., A. R. A. P. and E. W. M. acknowledge financial support from the Dutch Ministry of Education, Culture and Science (Gravity program 024.001.035) and the European Research Council (FP7/2007-2013, ERC Grant Agreement 246829). P. T., K. G. B. and A. E. R. acknowledge financial support from the Dutch Ministry of Education, Culture and Science (Gravity program 024.001.035), the Dutch National Research School Combination Catalysis Controlled by Chemical Design (NRSCC grant 2009-10016B, A. E. R.), as well as The Netherlands Organization for Scientific Research (NWO; VIDI grant 700.58.430, K. G. B.). Dr. L. Albertazzi is acknowledged for fruitful discussions. Gijs ter Huurne is acknowledged for assistance with the SAXS measurements. The ICMS animation studio is acknowledged for providing the artwork.

References

- M. Ouchi, N. Badi, J.-F. Lutz and M. Sawamoto, *Nat. Chem.*, 2011, 3, 917–924.
- S. Mavila, O. Eivgi, I. Berkovich and N. G. Lemcoff, *Chem. Rev.*, 2016, 116, 878–961.
- O. Altintas and C. Barner-Kowollik, *Macromol. Rapid Commun.*, 2012, 33, 958–971.
- M. Gonzalez-Burgos, A. Latorre-Sanchez and J. A. Pomposo, *Chem. Soc. Rev.*, 2015, 44, 6122–6142.
- A. M. Hanlon, C. K. Lyon and E. B. Berda, *Macromolecules*, 2016, 49, 2–14.
- J.-F. Lutz, J.-M. Lehn, E. W. Meijer and K. Matyjaszewski, *Nat. Rev. Mater.*, 2016, 1, 16024.
- J. Willenbacher, O. Altintas, V. Trouillet, N. Knöfel, M. J. Monteiro, P. W. Roesky and C. Barner-Kowollik, *Polym. Chem.*, 2015, 6, 4358–4365.
- N. D. Knöfel, H. Rothfuss, J. Willenbacher, C. Barner-Kowollik and P. W. Roesky, *Angew. Chem., Int. Ed.*, 2017, 56, 4950–4954.
- T. Terashima, T. Mes, T. F. A. De Greef, M. A. J. Gillissen, P. Besenius, A. R. A. Palmans and E. W. Meijer, *J. Am. Chem. Soc.*, 2011, 133, 4742–4745.
- I. Berkovich, S. Mavila, O. Iliashevsky, S. Kozuch and N. G. Lemcoff, *Chem. Sci.*, 2016, 7, 1773–1778.
- A. Sanchez-Sanchez, A. Arbe, J. Colmenero and J. A. Pomposo, *ACS Macro Lett.*, 2014, 3, 439–443.
- S. Thanneeru, S. S. Duay, L. Jin, Y. Fu, A. M. Angeles-Boza and J. He, *ACS Macro Lett.*, 2017, 6, 652–656.
- Y. Bai, X. Feng, H. Xing, Y. Xu, B. K. Kim, N. Baig, T. Zhou, A. A. Gewirth, Y. Lu, E. Oldfield and S. C. Zimmerman, *J. Am. Chem. Soc.*, 2016, 138, 11077–11080.
- K. Freytag, S. Säfken, K. Wolter, J. C. Namyslo and E. G. Hübner, *Polym. Chem.*, 2017, 8, 7546–7558.
- S. Mavila, I. Rozenberg and N. G. Lemcoff, *Chem. Sci.*, 2014, 5, 4196–4203.
- D. E. Bergbreiter, *ACS Macro Lett.*, 2014, 3, 260–265.
- G. Wulff, *Chem. Rev.*, 2002, 102, 1–28.
- N. E. Leadbeater and M. Marco, *Chem. Rev.*, 2002, 102, 3217–3274.
- L. J. Twyman, A. S. H. King and I. K. Martin, *Chem. Soc. Rev.*, 2002, 31, 69–82.
- D. Méry and D. Astruc, *Coord. Chem. Rev.*, 2006, 250, 1965–1979.
- J. Kofoed and J.-L. Reymond, *Curr. Opin. Chem. Biol.*, 2005, 9, 656–664.
- E. Huerta, B. van Genabeek, P. J. M. Stals, E. W. Meijer and A. R. A. Palmans, *Macromol. Rapid Commun.*, 2014, 35, 1320–1325.
- M. Artar, E. R. J. Souren, T. Terashima, E. W. Meijer and A. R. A. Palmans, *ACS Macro Lett.*, 2015, 4, 1099–1103.
- Y. Liu, S. Pujals, P. J. M. Stals, T. Paulöhr, S. I. Presolski, E. W. Meijer, L. Albertazzi and A. R. A. Palmans, *J. Am. Chem. Soc.*, 2018, 140, 3423–3433.
- M. A. J. Gillissen, T. Terashima, E. W. Meijer, A. R. A. Palmans and I. K. Voets, *Macromolecules*, 2013, 46, 4120–4125.
- G. M. ter Huurne, L. N. J. de Windt, Y. Liu, E. W. Meijer, I. K. Voets and A. R. A. Palmans, *Macromolecules*, 2017, 50, 8562–8569.
- N. Hosono, A. M. Kushner, J. Chung, A. R. A. Palmans, Z. Guan and E. W. Meijer, *J. Am. Chem. Soc.*, 2015, 137, 6880–6888.



- 28 P. J. M. Stals, C.-Y. Cheng, L. van Beek, A. C. Wauters, A. R. A. Palmans, S. Han and E. W. Meijer, *Chem. Sci.*, 2016, 7, 2011–2015.
- 29 W. E. Moerner and D. P. Fromm, *Rev. Sci. Instrum.*, 2003, 74, 3597–3619.
- 30 P. Tinnefeld and M. Sauer, *Angew. Chem., Int. Ed.*, 2005, 44, 2642–2671.
- 31 T. G. Terentyeva, J. Hofkens, T. Komatsuzaki, K. Blank and C.-B. Li, *J. Phys. Chem. B*, 2013, 117, 1252–1260.
- 32 V. I. Claessen, H. Engelkamp, P. C. M. Christianen, J. C. Maan, R. J. M. Nolte, K. Blank and A. E. Rowan, *Annu. Rev. Anal. Chem.*, 2010, 3, 319–340.
- 33 P. Chen, X. Zhou, M. Andoy, K.-S. Han, E. Choudhary, N. Zou, G. Chen and H. Shen, *Chem. Soc. Rev.*, 2014, 43, 1107–1117.
- 34 D. Wöll, H. Uji-i, T. Schnitzler, J. Hotta, P. Dedecker, A. Herrmann, F. C. De Schryver, K. Müllen and J. Hofkens, *Angew. Chem., Int. Ed.*, 2008, 47, 783–787.
- 35 Q. T. Easter and S. A. Blum, *Angew. Chem., Int. Ed.*, 2017, 56, 13772–13775.
- 36 Q. T. Easter and S. A. Blum, *Angew. Chem., Int. Ed.*, 2018, 57, 1572–1575.
- 37 G. K. Hodgson, S. Impellizzeri and J. C. Scaiano, *Chem. Sci.*, 2016, 7, 1314–1321.
- 38 Z. Ristanović, J. P. Hofmann, G. De Cremer, A. V. Kubarev, M. Rohnke, F. Meirer, J. Hofkens, M. B. J. Roeflaers and B. M. Weckhuysen, *J. Am. Chem. Soc.*, 2015, 137, 6559–6568.
- 39 P. Turunen, A. E. Rowan and K. Blank, *FEBS Lett.*, 2014, 588, 3553–3563.
- 40 J. D. Ng, S. P. Upadhyay, A. N. Marquard, K. M. Lupo, D. A. Hinton, N. A. Padilla, D. M. Bates and R. H. Goldsmith, *J. Am. Chem. Soc.*, 2016, 138, 3876–3883.
- 41 B. Dong, Y. Pei, F. Zhao, T. W. Goh, Z. Qi, C. Xiao, K. Chen, W. Huang and N. Fang, *Nat. Catal.*, 2018, 1, 135–140.
- 42 K. Velonia, O. Flomenbom, D. Loos, S. Masuo, M. Cotlet, Y. Engelborghs, J. Hofkens, A. E. Rowan, J. Klafter, R. J. M. Nolte and F. C. de Schryver, *Angew. Chem., Int. Ed.*, 2005, 44, 560–564.
- 43 N. S. Hatzakis, H. Engelkamp, K. Velonia, J. Hofkens, P. C. M. Christianen, A. Svendsen, S. A. Patkar, J. Vind, J. C. Maan, A. E. Rowan and R. J. M. Nolte, *Chem. Commun.*, 2006, 2012–2014.
- 44 N. S. Hatzakis, L. Wei, S. K. Jorgensen, A. H. Kunding, P.-Y. Bolinger, N. Ehrlich, I. Makarov, M. Skjot, A. Svendsen, P. Hedegård and D. Stamou, *J. Am. Chem. Soc.*, 2012, 134, 9296–9302.
- 45 T. G. Terentyeva, H. Engelkamp, A. E. Rowan, T. Komatsuzaki, J. Hofkens, C.-B. Li and K. Blank, *ACS Nano*, 2012, 6, 346–354.
- 46 A. Das and P. Theato, *Chem. Rev.*, 2016, 116, 1434–1495.
- 47 M. A. Gauthier, M. I. Gibson and H.-A. Klok, *Angew. Chem., Int. Ed.*, 2009, 48, 48–58.
- 48 L. D. Lavis, T.-Y. Chao and R. T. Raines, *ACS Chem. Biol.*, 2006, 1, 252–260.
- 49 T. G. Terentyeva, W. Van Rossom, M. Van Der Auweraer, K. Blank and J. Hofkens, *Bioconjugate Chem.*, 2011, 22, 1932–1938.
- 50 Z.-Q. Wang, J. Liao and Z. Diwu, *Bioorg. Med. Chem. Lett.*, 2005, 15, 2335–2338.
- 51 A. A. Kislukhin, V. P. Hong, K. E. Breitenkamp and M. G. Finn, *Bioconjugate Chem.*, 2013, 24, 684–689.
- 52 A. Isidro-Llobet, M. Álvarez and F. Albericio, *Chem. Rev.*, 2009, 109, 2455–2504.
- 53 J. Li and P. R. Chen, *Nat. Chem. Biol.*, 2016, 12, 129–137.
- 54 P. Zhu and H. G. Craighead, *Annu. Rev. Biophys.*, 2012, 41, 269–293.
- 55 M. J. Levene, J. Korlach, S. W. Turner, M. Foquet, H. G. Craighead and W. W. Webb, *Science*, 2003, 299, 682–686.
- 56 H. Rigneault, J. Capoulade, J. Dintinger, J. Wenger, N. Bonod, E. Popov, T. W. Ebbesen and P.-F. Lenne, *Phys. Rev. Lett.*, 2005, 95, 117401.
- 57 D. Gérard, J. Wenger, N. Bonod, E. Popov, H. Rigneault, F. Mahdavi, S. Blair, J. Dintinger and T. W. Ebbesen, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 2008, 77, 45413.
- 58 P. A. Romero and F. H. Arnold, *Nat. Rev. Mol. Cell Biol.*, 2009, 10, 866–876.

