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# Inhibitor-conjugated harmonic nanoparticles targeting fibroblast activation protein†

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The recent progress in the engineering of nanosized inorganic materials presenting tailored physical properties and reactive surface for post-functionalization has opened promising avenues for the use of nanoparticles (NPs) in diagnosis and therapeutic intervention. Surface decoration of metal oxide NPs with ligands modulating circulation time, cellular uptake, affinity and extravasation through active targeting led to efficient cancer specific bioimaging probes. The most relevant cancer biomarkers studied so far include surface and transmembrane cancer cell receptors. More recently, tumor microenvironments and more specifically the fibroblastic element of the tumor stroma have emerged as a valuable target for diagnosis and treatment of several types of cancers. In this study, a low molecular weight ligand targeting fibroblast activation protein  $\alpha$  (FAP), which is specifically expressed by activated fibroblasts of the tumor stroma, was synthesized. This ligand demonstrated nanomolar inhibition of FAP with high selectivity with respect to prolyl oligopeptidase (PREP) and dipeptidyl peptidase (DPP) IV, as well as good biocompatibility toward a human lung tissue model. Bismuth ferrite (BFO) harmonic nanoparticles (HNPs) conjugated to this ligand showed target-specific association to FAP as demonstrated by reverse ELISA-type assay using Human Fibroblast Activation Protein alpha/FAP Alexa Fluor® 594-conjugated Antibody and multiphoton multispectral microscopy experiments. These functionalized HNPs may provide new nanocarriers to explore the role of FAP in tumorigenesis and to target the fibroblastic component of the tumor microenvironment.

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## Introduction

The ability to produce inorganic and organic nanoparticles (NPs) of tunable size and composition, combined with their high agent loading capacity and surface properties suitable for tailored chemical modifications, have generated high expectations that nanomaterials could provide unprecedented diagnosis and therapeutic tools.<sup>1</sup> In particular, inorganic nanostructures appear as highly promising candidates for cancer diagnosis and therapy due to their intrinsic optical and physical properties, as well as their capacity to be transformed into multifunctional platforms for the

co-delivery of multiple payloads.<sup>2</sup> In addition, the opportunity to activate modalities (*e.g.* photothermal and photodynamic effects) can enhance and complement the chemotherapeutic effects.<sup>3</sup> Major achievements in the field were reported with silica-based nanocarriers,<sup>4–7</sup> magnetic NPs,<sup>8,9</sup> gold nanocomposites<sup>10–15</sup> and quantum dots.<sup>16</sup> While conventional chemotherapies are generally poorly selective toward specific tissues, cells and intracellular compartments, inorganic NPs benefit from specific uptake by macrophages of the reticuloendothelial system or from the enhanced permeability and retention (EPR) effect.<sup>17</sup> Substantial advances in the design of NP-based delivery systems was achieved by the introduction of targeting ligands at the surface of the nanocarriers to promote specific interaction with cancer cell biomarkers such as prostate specific membrane antigen, folate receptor, vascular endothelial growth factor receptors,  $\alpha_v\beta_3$  integrin and epithelial growth factor receptors.<sup>18,19</sup> However, addressing malignant tumor cells only remains ineffective for several types of cancer. Evidence has accumulated that the microenvironment of a given tumor plays a crucial role in its progression, metastasis and response to chemotherapeutic intervention. In particular, the fibroblastic element of the tumor stroma, which is predominantly characterized by the abundance of activated fibroblasts often designated as cancer-associated fibroblasts (CAFs), appears as a promising target for both cancer diagnosis and treatment,<sup>20,21</sup>

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† Electronic supplementary information (ESI) available: Synthesis protocols for the preparation of compounds **4**, **6**, **8** and **Biotin-PEG<sub>3</sub>-FAPi**; analytical data (<sup>1</sup>H and <sup>13</sup>C NMR spectra for new synthesized compounds); analytical UPLC traces for compounds **5** and **Biotin-PEG<sub>3</sub>-FAPi**; procedure for the coating of BFO NPs with heterobifunctional PEG derivatives; STEM and FT-IR characterization of **BFO-PEG-FAPi**; qualitative analysis by ESI-HRMS; multiphoton multispectral microscopy. See DOI: 10.1039/c9ra05299b



including the development of targeted nanotheranostic systems.<sup>22,23</sup> The tumor-stroma symbiotic cross-talk was even proposed as a new cancer hallmark by Sverdlöv in 2018.<sup>24</sup> A specific characteristic of CAFs is the expression of fibroblast activation protein  $\alpha$  (FAP), a transmembrane protease which is not found in normal, healthy adult tissues, except in granulation tissue of healing wounds.<sup>25–27</sup> However, the mechanisms underlying the physiological and pathological functions of FAP are not fully elucidated.<sup>28</sup> The development of NP-based diagnostic and delivery systems targeting the proteolytic activity of FAP could contribute both to a better understanding of the role of FAP activity in modulating tumor behavior and to develop new anticancer treatments acting against the cancer-stroma symbiotic interactions.<sup>29,30</sup>

Recently, harmonic nanoparticles (HNPs), which are based on non-centrosymmetric metal oxide nanocrystals, have emerged as promising nanomaterials for bioimaging applications<sup>31</sup> due to several favorable properties,<sup>32–34</sup> including their efficient nonlinear response to excitation from the UV to mid-IR,<sup>35</sup> their long-term photostability under pulsed laser irradiation and their spectrally narrow emission signal.<sup>36</sup> Upon screening of different types of HNPs, bismuth ferrite (BiFeO<sub>3</sub>, BFO) HNPs demonstrated very high second harmonic efficiency<sup>37,38</sup> as well as excellent *in vitro* cell compatibility after surface modification with poly(ethylene glycol) (PEG) derived coatings.<sup>39</sup> These properties were exploited in various bioimaging applications, including stem cell tracking in tissue depth<sup>40</sup> exploiting multi-harmonic detection<sup>41</sup> and *in vitro* lung cancer cells imaging.<sup>42</sup> Other applications of BFO HNPs include contrast enhancement in magnetic resonance imaging and computed tomography,<sup>43</sup> and synergistic photothermal–photodynamic combination for solid tumor ablation.<sup>44</sup>

Herein, we present the design and production of BFO-based imaging HNPs targeting the fibroblastic elements of cancer tumor stroma by covalent conjugation at the surface of PEG-coated BFO NPs of a low molecular weight ligand containing a highly selective inhibitor of FAP. First, we synthesized a selective low nanomolar inhibitor of FAP containing cyanopyrrolidine and quinolinoyl subunits, as suggested by the SAR study reported by Jansen *et al.*<sup>45</sup> Further derivatization of this inhibitor with a 4-dibenzocyclooctyne (DIBO) moiety provided a targeting ligand for conjugation at the surface of coated BFO HNPs through copper-free azide–alkyne [3 + 2] cycloaddition (click reaction). Then, the resulting functionalized NPs were assessed for their association with human recombinant (hr) FAP, in the presence of a fluorescently labelled anti-FAP antibody, by co-localization of the second harmonic signal of the HNPs with the fluorescent signal from the antibody. This work enlarges the panel of multifunctional inorganic nanoparticles for cancer imaging by targeting the fibroblastic components of the tumor stroma rather than malignant tumor cell biomarkers.

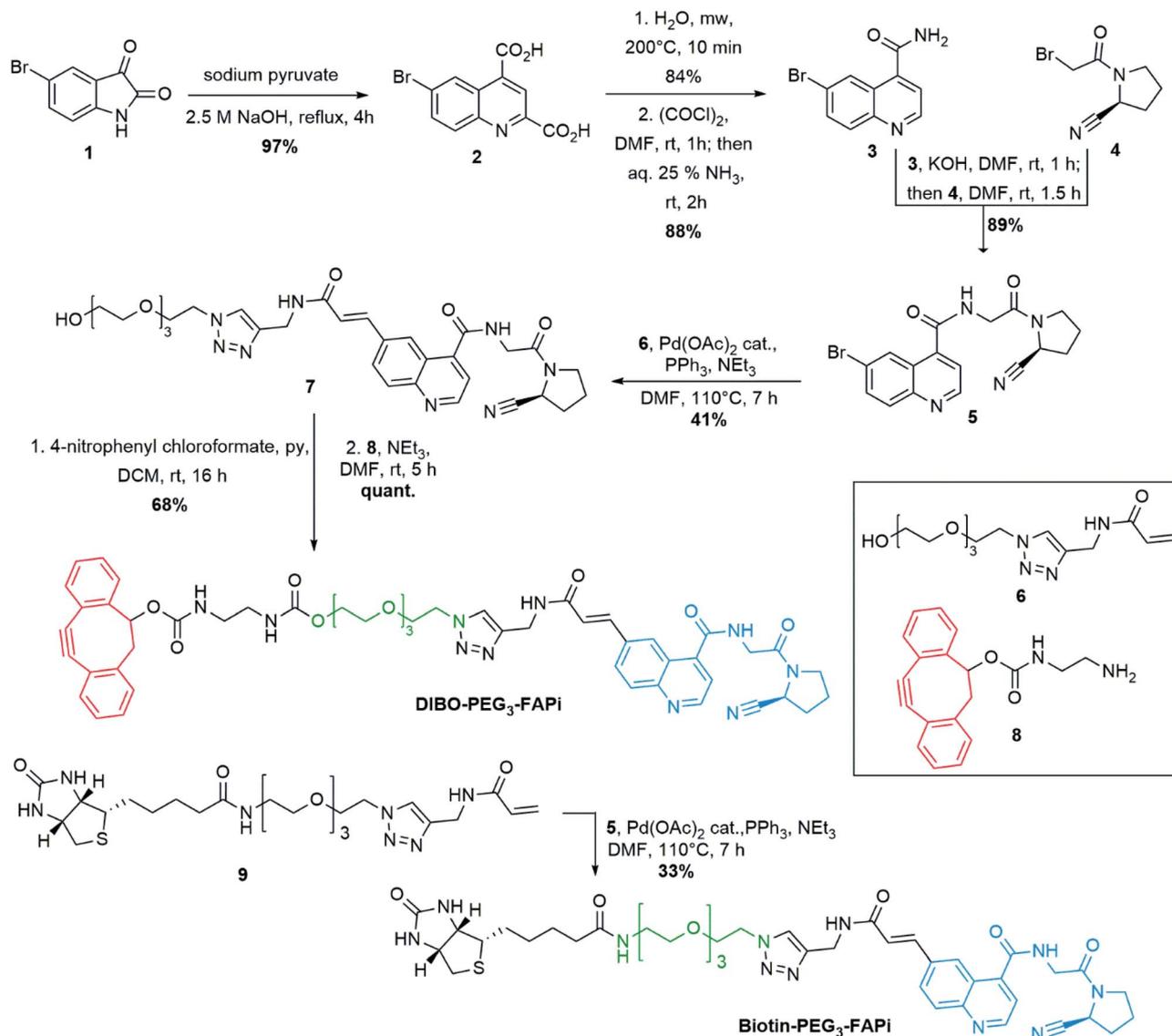
## Results and discussion

### Preparation of FAP inhibitor and targeting ligand for conjugation to BFO HNPs

Among the family of human prolyl-specific proteases, FAP presents the unique property to display both endo- and

exoproteolytic activities, while other closely related proteases are exclusively exopeptidases, such as amino-dipeptidyl peptidase (DPP) IV, or exclusively endopeptidases such as prolyl oligopeptidase (PREP).<sup>25</sup> FAP, which is specifically associated to CAFs and stromal cells appears as a promising target for anti-cancer therapy. Due to its adhesive properties, DPP IV might also favor cancer progression. However, there is no evidence for the efficacy of inhibiting DPP IV in tumor growth reduction.<sup>46,47</sup> The abnormal activity of PREP was linked to neurological disorders in mammals. This enzyme was thus identified as a potential target in neurodegenerative disorders such as amnesia, Alzheimer's disease and depression.<sup>48,49</sup> In view of the similar hydrolytic mechanisms and kinetics characteristics of these three prolylpeptidases, the design of selective inhibitors is key to the development of therapeutic approaches based on their inhibition, to avoid undesired off-target activities. Importantly, *N*-acyl-Gly-Pro derived motifs were identified as key components to discriminate between FAP and the other prolyl peptidases DPP IV, VII, VIII, and IX.<sup>50</sup> However, until the demonstration by Jansen *et al.* that a (4-quinolinoyl)glycyl-2-cyanopyrrolidine motif could avoid cross-inhibition of both DPPs and PREP,<sup>45</sup> most of reported FAP inhibitors were not entirely specific.<sup>51,52</sup> We investigated the functionalization of the quinoline ring's 6-position of the (4-quinolinoyl)glycyl-2-cyanopyrrolidine core to introduce a suitable spacer between the subunit targeting FAP and the imaging BFO NPs. Based on previous work,<sup>42</sup> covalent conjugation to the surface of PEG-coated BFO NPs should be performed in (1 : 1) EtOH : H<sub>2</sub>O medium to ensure efficient copper-free azide–alkyne [3 + 2] cycloaddition. Therefore, a PEG<sub>3</sub> linker was selected to modulate the polarity of the targeting ligand and a terminal acrylamide was envisaged for cross-coupling to the 6-position of the quinoline ring (Scheme 1). Preparation of the 6-bromoquinoline core started from 5-bromoisatin which was submitted to a Pfizinger reaction<sup>53</sup> in the presence of sodium pyruvate to produce the dicarboxylic intermediate **2** in 97% yield. Microwave assisted selective decarboxylation,<sup>54</sup> followed by amide formation delivered 6-bromoquinoline-4-carboxamide (**3**) in high yield. Condensation with cyano-proline derivative **4** (synthesis described in ESI S-3†) afforded intermediate **5** which constitutes the central unit of the ligand targeting FAP. Introduction of the polar spacer was achieved by Pd-catalyzed Heck reaction<sup>55</sup> in the presence of the acrylamide derivative **6** (synthesis described in ESI S-5†). Activation of the terminal alcohol as 4-nitrophenyl carbonate allowed condensation with the DIBO derivative **8** (synthesis described in ESI S-8†) under mild conditions to generate **DIBO-PEG<sub>3</sub>-FAPi** (68% yield, 2 steps), ready to be conjugated to PEG-coated BFO NPs. To evaluate the FAP targeting ability of this ligand, a similar pathway was applied on compound **9** to introduce a biotin label in place of the strained cyclooctyne moiety, affording **Biotin-PEG<sub>3</sub>-FAPi** (synthesis described in ESI S-11†). Introduction of a biotin residue resulted in improved solubility of the targeting ligand and allowed evaluation of its inhibitory activity in absence of the sensitive cyclooctyne subunit.





Scheme 1 Synthesis of DIBO-PEG<sub>3</sub>-FAPi, targeting ligand for conjugation to the surface of imaging NPs. Synthesis of Biotin-PEG<sub>3</sub>-FAPi for evaluation of FAP targeting ability.

### Evaluation of the inhibitory activity on FAP, DPP IV and PREP

In order to assess the ability of the designed ligand to selectively target FAP, compounds 5 and **Biotin-PEG<sub>3</sub>-FAPi** were assessed for their inhibitory activity on human recombinant (hr) FAP, DPP IV and PREP (Table 1). Compound 5 showed potent inhibition of FAP and excellent selectivity with respect to PREP and DPP IV, which remained unaffected (selectivity index (SI) higher than 3300). Remarkably, addition of a PEG<sub>3</sub> spacer significantly increased the potency against FAP as **Biotin-PEG<sub>3</sub>-FAPi** reached IC<sub>50</sub> and K<sub>i</sub> values of 9.2 and 3.8 nM, respectively. Importantly, a sharp selectivity with respect to PREP and DPP IV was observed (SI<sub>FAP/DPP IV</sub> and SI<sub>FAP/PREP</sub> higher than 10 000), which confers to **Biotin-PEG<sub>3</sub>-FAPi** a highly favorable affinity/selectivity profile for the specific targeting of FAP.

In order to verify the compatibility of the targeting ligand in physiological environment, the effect of **Biotin-PEG<sub>3</sub>-FAPi** was evaluated on a model of human functional lung epithelium

(MucilAir™-HF) based on the co-culture of bronchial cells and lung fibroblasts.<sup>56</sup> MucilAir™-HF was submitted to repeated doses of **Biotin-PEG<sub>3</sub>-FAPi**, at 10 or 100 μM concentration, over

Table 1 Kinetic characteristics for the inhibition of human recombinant prolyl-oligopeptidases

Compound	hrPREP <sup>a</sup>	hrDPP IV <sup>a</sup>	hrFAP <sup>a</sup>	K <sub>i</sub> (nM)
	IC <sub>50</sub> (μM)	IC <sub>50</sub> (μM)	IC <sub>50</sub> (μM)	
5	>100	>100	0.03 ± 0.005	12 ± 1.8
<b>Biotin-PEG<sub>3</sub>-FAPi</b>	>100	>100	0.0092 ± 0.0006	3.8 ± 0.23

<sup>a</sup> Enzymes were incubated with increasing concentrations (5, 10, 20, 50 and 100 nM) of compounds 5 and **Biotin-PEG<sub>3</sub>-FAPi**, for 30 min at 37 °C. Then, the residual enzymatic activity was measured after exposure to 50 μM of the appropriate substrates (Z-Gly-Pro-AMC for hrFAP and hrPREP, H-Gly-Pro-AMC for hrDPP IV). The half maximal inhibitory concentrations (IC<sub>50</sub>) were graphically determined and the inhibition constants (K<sub>i</sub>) were calculated.<sup>57</sup>



16 days. The cytotoxicity was measured by dosing Lactate Dehydrogenase (LDH) released in the culture supernatant at regular time points (Fig. 1A). At 10  $\mu\text{M}$  concentration (3 basal applications per week), the targeting ligand did not affect cell viability in the tissue model. Cytotoxic effect started to be noticeable at higher doses (100  $\mu\text{M}$ ) from day 8. Similarly, the integrity and functionality of the epithelium, analyzed by trans-epithelial electrical resistance (TEER) and cilia beat frequency (CBF) measurements (Fig. 1B and C), were maintained upon repeated applications of **Biotin-PEG<sub>3</sub>-FAPi**, at 10  $\mu\text{M}$ . Damages to the epithelium were only observed at high doses (100  $\mu\text{M}$ ) as TEER was affected and cilia beat was lost after day 8. These data suggest the good compatibility of the designed ligand in a concentration range which is relevant for labelling and imaging applications.

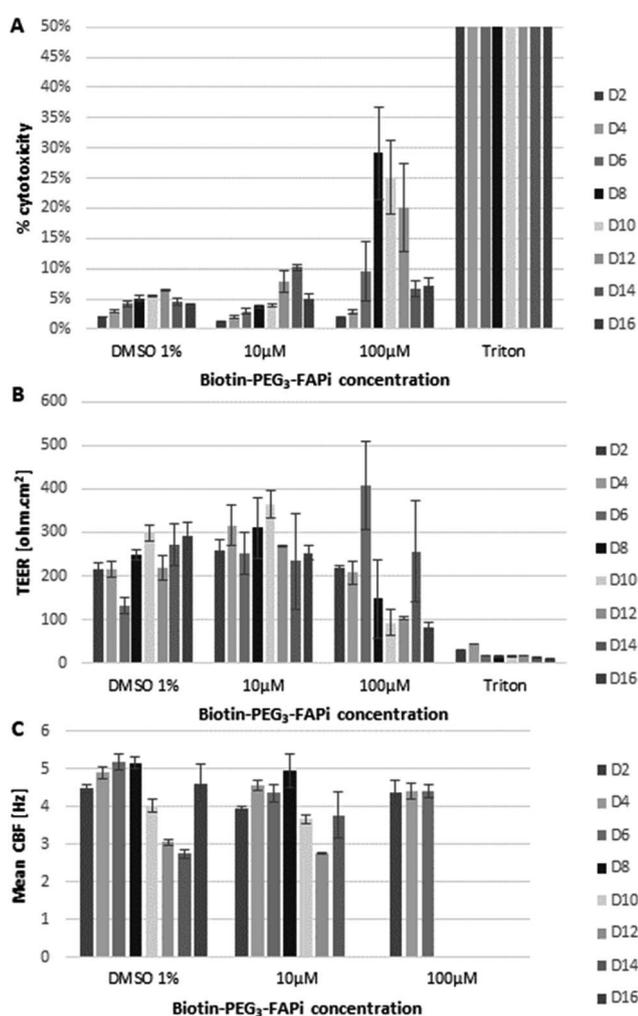


Fig. 1 Evaluation of the cytocompatibility of the targeting ligand on the human lung tissue model MucilAir™-HF. Tissue was exposed to repeated doses (10  $\mu\text{M}$  or 100  $\mu\text{M}$ ) of **Biotin-PEG<sub>3</sub>-FAPi** (day 0, 2, 4, 7, 9, 11 and 14) and measurements were made on different days (day 2, 4, 6, 8, 10, 12, 14 and 16). (A) Cytotoxicity was evaluated by dosing LDH in the culture supernatant. (B) Trans-epithelial electrical resistance (TEER) measurement. (C) Cilia beat frequency (CBF) measurement. Triton X-100 was used as positive control; DMSO (1%) was used as negative control.

## Conjugation of BFO NPs with targeting ligand

As previously established,<sup>39</sup> BFO NPs were coated with a 1 : 1 mixture of linear PEG derivatives presenting end amino and azido groups (for detailed protocol, see ESI S-29†). Resulting PEGylated NPs were ultrasonicated in the presence of **DIBO-PEG<sub>3</sub>-FAPi** (16 h, 40 °C) allowing covalent conjugation of the targeting ligand through a 1,2,3-triazole moiety, under mild conditions (Fig. 2A). Separation from unreacted ligand was operated by cycles of emulsification in EtOAc followed by centrifugation, to afford **BFO-PEG-FAPi** NPs which were characterized for their size and surface charge. Their mean hydrodynamic diameter, measured by dynamic light scattering, was  $50 \pm 15$  nm (Fig. 2B) and their zeta potential averaged  $-17.7 \pm 1.6$  mV. After coating and post-conjugation with the FAP targeting ligand, the average size of the functionalization layer was about 8 nm as measured on STEM images (ESI, Fig. S1†). The FT-IR spectrum of **BFO-PEG-FAPi** featured the apparition of bands at 1650 and 1375  $\text{cm}^{-1}$ , corresponding to alkene C=C stretching and triazole C-N stretching (ESI, Fig. S2†). These data support the conjugation of **DIBO-PEG<sub>3</sub>-FAPi** to the surface of **BFO-PEG** NPs through a click reaction. According to the principles of nanoparticles design for biomedical applications presented by Blanco *et al.*,<sup>58</sup> the size range and surface charge of **BFO-PEG-FAPi** NPs are expected to provide long-lasting systems in physiological environment.

## Association of BFO-PEG-FAPi NPs with FAP

The association of **BFO-PEG-FAPi** NPs with hrFAP, in comparison with **BFO-PEG** NPs was evaluated after incubation for 2 h at 37 °C of the functionalized or coated NPs with hrFAP, in the presence of an excess of Human Fibroblast Activation Protein alpha/FAP Alexa Fluor® 594-conjugated Antibody. The samples were centrifuged to remove the NPs from the supernatant, which was further analyzed on a fluorescence spectrophotometer ( $\lambda_{\text{ex}}/\lambda_{\text{em}} = 590/645$  nm) (Fig. 3). The fluorescence intensity of the solutions containing the anti-FAP antibody with hrFAP at the same concentrations than the ones used for the incubation experiments, was taken as reference. The incubation of **BFO-PEG-FAPi** NPs with hrFAP resulted in a significant decrease of the fluorescence intensity compared with the controls and to **BFO-PEG** NPs, indicating that the functionalized nanoparticles associated with FAP *via* a target-specific interaction.

The association of **BFO-PEG-FAPi** NPs to FAP was further analyzed by multiphoton microscopy. After incubation of hrFAP with **BFO-PEG** NPs or **BFO-PEG-FAPi** NPs for 2 hours at 37 °C, in the presence of the labeled anti-FAP antibody, the resulting conjugates were imaged by detection of the second harmonic (SH) signal from the NPs and of the fluorescent emission from the antibody, upon laser pulsed excitation at 840 nm (ESI, Fig. S3†). The ratio of conjugates emitting in both SHG (420 nm) and fluorescence (580–650 nm) channels to all structures emitting in the SHG channel was calculated in 4 different regions of each sample and averaged (Fig. 4). The number of conjugates generating both harmonic and fluorescence signals was significantly higher in the presence of **BFO-PEG-FAPi** NPs than in the presence of **BFO-PEG** NPs, thus confirming the



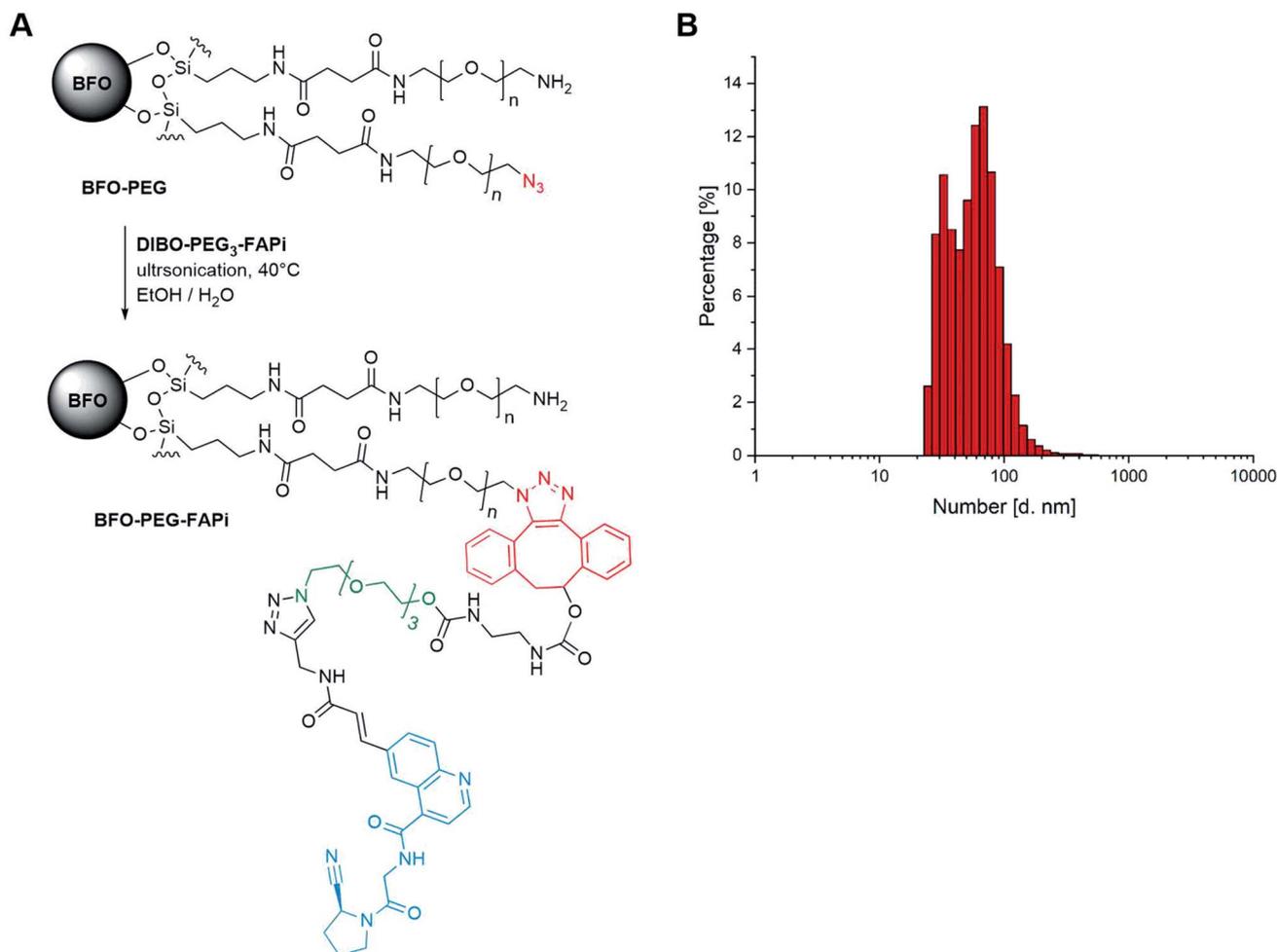


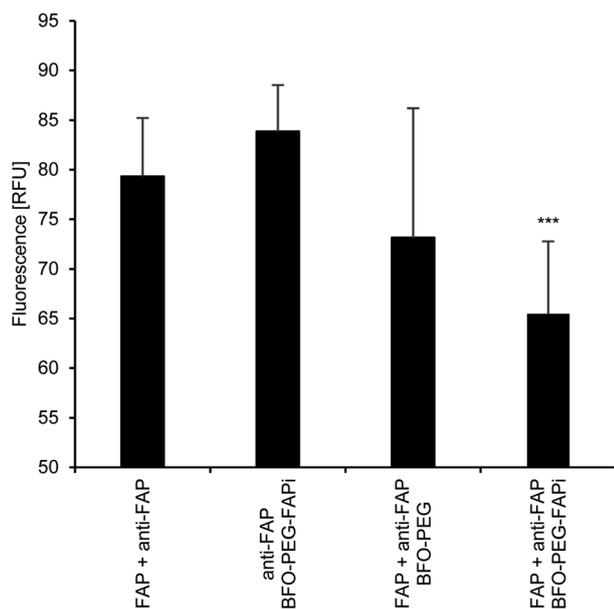
Fig. 2 (A) Synthesis scheme of functionalized BFO-PEG-FAPi NPs; (B) dynamic light scattering analyses of functionalized nanoparticles.

target-specific association of functionalized BFO HNPs with FAP.

While many of the proteases involved in cancer development and progression are expressed by both healthy and malignant cells, FAP holds the unique property to be mostly present at remodeling stroma in tumors and healing wounds.<sup>59</sup> This remarkable feature makes FAP a highly promising target for both diagnosis and therapeutic intervention. While several inhibitors of FAP based on a pyrrolidine-2-boronic acid scaffold were evaluated as potential anti-cancer drugs, safety concerns were raised during clinical trials due to the cross inhibition of other related enzymes.<sup>60–62</sup> Having previously disclosed the efficiency of PEGylated BFO NPs for selective imaging and cell tracking in tissue samples,<sup>40,42</sup> we focused on the design and synthesis of a targeting ligand which would exhibit selective and high affinity for FAP while holding a suitable spacer to be conjugated at the surface of BFO-PEG NPs. Functionalization of a (4-quinolinoyl)glycyl-2-cyanopyrrolidine core by cross-coupling on the 6-position of the quinoline ring allowed the introduction of a small PEG spacer to modulate the polarity of the ligand for further post-conjugation reaction in EtOH/H<sub>2</sub>O medium. Conjugation of this inhibitory scaffold to *N*-(2-

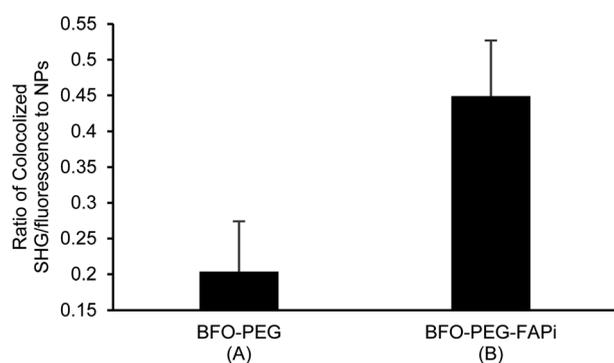
hydroxypropyl)methacrylamide copolymer was previously used to visualize and isolate FAP from cell lysates.<sup>63</sup> Interestingly, derivatization with a PEG spacer delivered **Biotin-PEG<sub>3</sub>-FAPi** which displayed effective binding to FAP ( $K_i = 3.8 \pm 0.20$  nM), high specific discrimination between FAP and its closely related proteases DPP IV and PREP (selectivity indexes above 10 000), and good biocompatibility with the human 3D tissue model MucilAir<sup>TM</sup>-HF.<sup>56</sup> Combination of a *N*-acyl-Gly-Pro fragment with a (4-quinolinoyl)glycyl-2-cyanopyrrolidine moiety and a polar PEG spacer resulted in sharp specificity toward FAP. In agreement with our previous report,<sup>42</sup> the use of a DIBO moiety allowed spontaneous azide to alkyne [3 + 2]-cycloaddition at the surface of PEG-BFO NPs, under mild conditions, for covalent coupling of the targeting ligand at the surface of the imaging NPs. While coated NPs did not show any significant interaction with FAP, functionalized BFO-PEG-FAPi NPs displayed efficient and stable association with FAP, assessed both by multiphoton microscopy and reverse ELISA-type assay using Human Fibroblast Activation Protein alpha/FAP Alexa Fluor® 594-conjugated Antibody. These results suggest the formation of a stable complex between BFO-PEG-FAPi NPs and FAP despite the potential reversibility of the covalent bond formation between





**Fig. 3** Association of BFO-PEG-FAPi NPs to FAP through target-specific interactions. HrFAP (FAP, 100 ng) was incubated for 2 h at 37 °C with BFO-PEG-FAPi NPs (100  $\mu\text{g mL}^{-1}$ ) or BFO-PEG NPs (100  $\mu\text{g mL}^{-1}$ ) in the presence of Human Fibroblast Activation Protein alpha/FAP Alexa Fluor® 594-conjugated Antibody (anti-FAP) (dilution 1 : 50). The samples were centrifuged (5 min, 10 000 rpm) and the supernatants were transferred into a flat bottom 96-well plate. Fluorescence was measured in a multi-well plate spectrophotometer ( $\lambda_{\text{ex}}/\lambda_{\text{em}} = 590/645 \text{ nm}$ ). HrFAP incubated with anti-FAP and BFO-PEG-FAPi NPs incubated with anti-FAP were used as controls. Results were analyzed using a Student's *t*-test. Fluorescence intensity of BFO-PEG-FAPi NPs incubated with FAP and anti-FAP was significantly lower than BFO-PEG NPs incubated with FAP and anti-FAP and the fluorescence intensity of the controls (all comparisons  $***p < 0.001$ ).

the cyanopyrrolidine moiety and FAP active site serine residue.<sup>51</sup> The possibility to exploit the simultaneous acquisition of second- and third-harmonic signals of BFO NPs for selective



**Fig. 4** Association of BFO-PEG NPs and BFO-PEG-FAPi NPs to hrFAP quantified by multiphoton multispectral microscopy. HrFAP (100 ng) was incubated for 2 h at 37 °C with BFO-PEG NPs (100  $\mu\text{g mL}^{-1}$ ) (A) or BFO-PEG-FAPi NPs (100  $\mu\text{g mL}^{-1}$ ) (B) in the presence of Human Fibroblast Activation Protein alpha/FAP Alexa Fluor® 594-conjugated Antibody (dilution 1 : 5). SH and fluorescence signals were epi-collected upon focused excitation at 840 nm. The ratio of the area of conjugates emitting in both SH and fluorescence channels to the area of SH emitting structures was averaged from four distinct regions of each sample (635  $\times$  635  $\mu\text{m}$ ). Error bars represent standard deviation from  $n = 4$  calculations.

imaging in complex tissue samples<sup>41</sup> combined with the stable association of BFO-PEG-FAPi NPs with FAP make these functionalized NPs promising nanodevices for cancer diagnosis applications.

## Conclusions

Based on a (4-quinolinoyl)glycyl-2-cyanopyrrolidine core, we designed and synthesized a high affinity and selective targeting ligand for FAP, which was covalently conjugated at the surface of BFO-PEG NPs. The resulting BFO-PEG-FAPi nanoconjugates demonstrated stable association with FAP, thus providing promising labelling and imaging tools to study and target the fibroblastic element of the tumor stroma rather than malignant tumoral cells.

## Experimental section

### Materials and methods

Reagent-grade solvents (Fluka, Riedel-de-Haën) and chemicals (Aldrich, Acros, Fluka, Sigma, Maybridge, TCI Chemicals, Apollo and Fluorochem) were used without further purification. BFO NPs (BF0011214) were obtained from TIBIO (Comano, Switzerland). All reactions were performed in flame-dried glassware under an inert atmosphere of nitrogen. All products were dried under vacuum ( $10^{-2}$  bar) before analytical characterization. Reactions were monitored by thin layer chromatography (TLC) on pre-coated aluminum plates SiO<sub>2</sub> 60 F254 (Merck, Darmstadt, Germany). The compounds were visualized by 254 nm light or stained with solutions of KMnO<sub>4</sub>, Pancaldi reagent [(NH<sub>4</sub>)<sub>6</sub>MoO<sub>4</sub>, Ce(SO<sub>4</sub>)<sub>2</sub>, H<sub>2</sub>SO<sub>4</sub>, H<sub>2</sub>O], ninhydrin or iodine vapors. Purifications were performed by flash chromatography (FC) on silica gel (Merck No. 9385 silica gel 60, 240–400 mesh, particle size 40–63  $\mu\text{m}$ ). NMR spectra were recorded on Bruker Avance III-400, Bruker Avance-400 or Bruker DRX-400 spectrometers (Bruker, Billerica, MA, USA) at room temperature (rt), unless otherwise stated. <sup>1</sup>H frequency is at 400.13 MHz, <sup>13</sup>C frequency is at 100.62 MHz. Chemical shifts are reported downfield from tetramethylsilane. <sup>1</sup>H signals are reported in ppm with the internal chloroform signal at 7.26 ppm or the internal DMSO signal at 2.50 ppm or the internal CH<sub>3</sub>CN signal at 1.94 ppm, as internal references. <sup>13</sup>C-NMR signals are reported in ppm with the internal chloroform signal at 77.16 ppm or the internal DMSO signal at 39.5 ppm or the internal CH<sub>3</sub>CN signal at 118.26 ppm, as internal references. The resonance multiplicity is described as s (singlet), d (doublet), t (triplet), q (quartet), quin (quintet), m (multiplet). Broad signals are indicated as br. Coupling constants (*J*) are given in hertz (Hz). IR spectra were recorded on a Jasco FT/IR-4100 spectrometer outfitted with a PIKE technology MIRacle™ ATR accessory as neat films compressed onto a Zinc Selenide window. The spectra are reported in cm<sup>-1</sup> (*w* = weak, *m* = medium, *s* = strong, *sh* = shoulder). The qualitative accurate masses were measured by ESI-TOF using the Xevo G2-S QTOF (Waters) and nanoESI-FT-MS using the Elite™ Hybrid Ion Trap-Orbitrap (ThermoFisher) mass spectrometer (more details in ESI S-30†). For all new compounds, the following data are provided:



accurate mass,  $^1\text{H}$  and  $^{13}\text{C}$  NMR spectra (with peak assignments), IR data. The purity of target compounds **5** and **Biotin-PEG<sub>3</sub>-FAPI** was tested on Waters Acquity UPLC® (gradient 98 : 2 to 2 : 98 · H<sub>2</sub>O containing 0.5% formic acid : ACN containing 0.45% formic acid in 4 minutes, flow rate 0.8 mL min<sup>-1</sup>) with column Acquity UPLC® BEH C18 1.7 μm. The purity of all new compounds is higher than 95%. Measurements of the dynamic light scattering and zeta potential were obtained using a Malvern NanoZ instrument (Malvern Instruments, Malvern, UK). Centrifugations were performed on HERAEUS Biofuge 13 centrifugator. Scanning transmission electron microscopy (STEM) was performed at the Interdisciplinary Centre for Electron Microscopy (CIME, EPFL, Lausanne, Switzerland) on a FEI Titan Themis 60-300 microscope.

### Synthesis protocols

Designation of the compounds refers to the chemical structures presented in Scheme 1.

#### Preparation of 6-bromoquinoline-2,4-dicarboxylic acid (2).

To a suspension of 5-bromoisatin (**1**) (1 equiv., 22.2 mmol, 5 g) in aqueous 2.5 M NaOH (53 mL), was added sodium pyruvate (1.2 equiv., 26.6 mmol, 2.92 g) and the reaction mixture was refluxed for 4 h. The mixture was cooled to rt and the pH was corrected to 2 with a solution of 6 M HCl in water. The precipitate was filtered and washed with water (100 mL) to afford **2** as a brown powder (21.5 mmol, 6.35 g, 97%).  $^1\text{H}$  NMR (400 MHz, DMSO-*d*<sub>6</sub>) δ 8.07 (1H, d, *J* = 9.0, 2.2 Hz), 8.19 (1H, d, *J* = 9.0 Hz), 8.54 (1H, s), 9.09 (1H, d, *J* = 2.2 Hz).  $^{13}\text{C}$  NMR (101 MHz, DMSO-*d*<sub>6</sub>) δ 122.8, 123.9, 126.7, 127.8, 132.6, 133.8, 135.8, 146.6, 149.1, 165.5, 166.5. HRMS *m/z* [M + H]<sup>-</sup> calcd for C<sub>11</sub>H<sub>5</sub>BrNO<sub>4</sub><sup>-</sup> 293.9407; found 293.9406. IR ( $\nu_{\text{max}}$ , cm<sup>-1</sup>): 2840 m, 2563 w, 1689 s, 1596 m, 1547 w, 1468 m, 1442 w, 1414 w, 1382 w, 1347 w, 1323 m, 1272 m, 1244 m, 1191 s, 1160 m, 1072 w, 1011 w, 923 m, 888 m, 837 m, 792 m, 746 w, 685 m.  $^1\text{H}$ -NMR,  $^{13}\text{C}$ -NMR spectra and signal assignments (ESI S-16†).

#### Preparation of 6-bromoquinoline-4-carboxamide (3).

A suspension of compound **2** (1 equiv., 0.84 mmol, 250 mg) in water (15 mL) was placed in a Pyrex pressure resistant tube and treated by microwave irradiation for 10 min at 200 °C. The mixture was slowly cooled down to rt and the precipitate was filtered and washed with water (15 mL) to afford intermediate 6-bromoquinoline-4-carboxylic acid as a brown powder (0.71 mmol, 179 mg, 84%). To a solution of this intermediate (1 equiv., 0.99 mmol, 250 mg) in anhydrous CHCl<sub>3</sub> (5 mL) was added dropwise (COCl)<sub>2</sub> (1 equiv., 0.99 mmol, 90 μL) and DMF (cat., 2 drops) and the mixture was stirred for 1 h at rt. Aqueous ammonia 25% (2 equiv., 1.98 mmol, 0.14 mL) was added dropwise and the mixture was stirred for 2 h at rt. Water (1 mL) was added and the volatiles were evaporated under reduce pressure. The pH was corrected to 11 with an aqueous solution of 3 M NaOH. The precipitate was filtered and washed with water (10 mL) to afford **3** as a brown powder (0.87 mmol, 218 mg, 88%).  $^1\text{H}$  NMR (400 MHz, DMSO-*d*<sub>6</sub>) δ 7.65 (1H, d, *J* = 4.3 Hz), 7.94 (1H, dd, *J* = 2.3, 9.0 Hz), 8.03 (1H, d, *J* = 9.0 Hz), 8.32 (2H, s), 8.46 (1H, d, *J* = 2.3 Hz), 9.01 (1H, d, *J* = 4.4 Hz).  $^{13}\text{C}$  NMR (101 MHz, DMSO-*d*<sub>6</sub>) δ 120.0, 120.5, 125.3, 127.6, 131.6,

132.8, 140.6, 146.6, 150.9, 168.0. HRMS *m/z* [M + H]<sup>+</sup> calcd for C<sub>10</sub>H<sub>8</sub>BrN<sub>2</sub>O<sup>+</sup> 250.9815; found 250.9814. IR ( $\nu_{\text{max}}$ , cm<sup>-1</sup>): 3343 m, 3122 w, 1945 w, 1910 w, 1668 s, 1622 m, 1577 m, 1496 m, 1449 m, 1398 m, 1337 s, 1300 w, 1200 w, 1135 w, 1060 w, 1011 w, 969 w, 862 s, 841 m, 827 m, 783 m, 757 w, 715 m, 685 m.  $^1\text{H}$ -NMR,  $^{13}\text{C}$ -NMR spectra and signal assignments (ESI S-19†).

**Preparation of (S)-6-bromo-N-(2-(2-cyanopyrrolidin-1-yl)-2-oxoethyl)quinoline-4-carboxamide (5).** To a solution of compound **3** (1.05 equiv., 1.98 mmol, 500 mg) in DMF (10 mL), KOH (2.5 equiv., 4.95 mmol, 56 mg) was added portion wise and the mixture was stirred for 1 h at rt. A solution of **4** (1 equiv., 1.89, 410 mg) in DMF (2 mL) was added dropwise and the mixture was stirred for 1.5 h at rt. The solvent was evaporated under reduced pressure and the residue was dissolved in DCM (10 mL). A sat. aqueous solution of NH<sub>4</sub>Cl (20 mL) was added to reach pH 5. The aqueous layer was extracted with DCM (3 × 10 mL). The combined organic layers were dried over MgSO<sub>4</sub>, filtered and concentrated *in vacuo*. The crude product was purified by FCC (DCM/MeOH 25 : 1) to afford **5** as a brown foam (1.76 mmol, 682 mg, 89%). Analytical UPLC: *R*<sub>t</sub> = 1.58 min.  $^1\text{H}$  NMR (400 MHz, chloroform-*d*) δ 2.19–2.43 (4H, m), 3.51–3.59 (1H, m), 3.69–3.76 (1H, m), 4.26 (1H, dd, *J* = 3.6, 17.8 Hz), 4.41 (1H, ddd, *J* = 2.8, 4.9, 17.8 Hz), 4.79 (1H, d, *J* = 5.2 Hz), 7.18 (1H, s), 7.54 (1H, d, *J* = 4.3 Hz), 7.83 (1H, dd, *J* = 2.2, 9.0 Hz), 8.00 (1H, d, *J* = 9.0 Hz), 8.49 (1H, d, *J* = 2.2 Hz) 8.95 (1H, dd, *J* = 2.3, 4.4 Hz).  $^{13}\text{C}$  NMR (101 MHz, chloroform-*d*) δ 25.2, 30.1, 42.6, 45.8, 46.9, 117.9, 119.6, 122.4, 125.7, 127.8, 131.7, 133.8, 139.8, 147.5, 150.3, 166.9, 167.0. HRMS *m/z* [M + H]<sup>+</sup> calcd for C<sub>17</sub>H<sub>16</sub>BrN<sub>4</sub>O<sub>2</sub><sup>+</sup> 387.0451; found 387.0450. IR ( $\nu_{\text{max}}$ , cm<sup>-1</sup>): 3297 w, 3058 w, 2954 w, 2168 w, 2156 w, 2022 w, 2880 w, 1655 s, 1586 w, 1537 w, 1493 w, 1430 m, 1407 m, 1305 m, 1345 w, 1263 m, 1194 w, 1157 w, 1043 w, 978 w, 913 w, 867 w, 846 w, 830 w, 783 w, 730 m, 723 m, 702 w.  $^1\text{H}$ -NMR,  $^{13}\text{C}$ -NMR spectra and signal assignments, UPLC trace (ESI S-21†).

**Preparation of (S,E)-N-(2-(2-cyanopyrrolidin-1-yl)-2-oxoethyl)-6-(3-(((1-(2-(2-(2-(2-hydroxyethoxy)ethoxy)ethoxy)ethyl)-1H-1,2,3-triazol-4-yl)methyl)amino)-3-oxoprop-1-en-1-yl)quinoline-4-carboxamide (7).** To a solution of compound **5** (1 equiv., 0.72 mmol, 280 mg) and compound **6** (2 equiv., 1.1 mmol, 355 mg) in degassed DCM (2.8 mL) in a Pyrex pressure resistant tube, were added Pd(OAc)<sub>2</sub> (10 mol%, 0.07 mmol, 16 mg), PPh<sub>3</sub> (40 mol%, 0.29 mmol, 76 mg) and NEt<sub>3</sub> (5.5 equiv., 4 mmol). The solution was stirred for 7 h at 110 °C. The solvent was evaporated *in vacuo* and the crude product was purified by FCC (DCM/MeOH 25 : 1 then 10 : 1) to afford **7** as a white oil (0.29 mmol, 185 mg, 41%).  $^1\text{H}$  NMR (400 MHz, chloroform-*d*) δ 2.17–2.38 (4H, m), 3.48–3.62 (11H, m), 3.64–3.72 (2H, m), 3.70–3.76 (1H, m), 3.83 (t, *J* = 5.0 Hz), 4.26 (1H, dd, *J* = 4.8, 17.3 Hz), 4.37 (1H, dd, *J* = 5.4, 17.3 Hz), 4.50 (2H, t, *J* = 5.0 Hz), 4.55 (2H, dd, *J* = 3.7, 5.6 Hz), 4.79–4.84 (1H, m), 6.69 (1H, d, *J* = 15.7 Hz), 7.37 (1H, t, *J* = 5.6 Hz), 7.42 (1H, d, *J* = 4.3 Hz), 7.59 (1H, d, *J* = 15.6 Hz), 7.68 (1H, dd, *J* = 1.8, 8.9 Hz), 7.72 (1H, t, *J* = 5.2 Hz), 7.82 (1H, s), 7.94 (1H, d, *J* = 8.8 Hz), 8.40 (1H, d, *J* = 1.5 Hz), 8.79 (1H, d, *J* = 4.3 Hz).  $^{13}\text{C}$  NMR (101 MHz, chloroform-*d*) δ 25.3, 29.9, 35.1, 42.5, 45.9, 47.0, 50.4, 61.6, 69.5, 70.3, 70.5, 70.60, 70.62, 72.6, 118.4, 119.6, 123.0, 123.8, 124.4, 125.1, 129.1, 130.2, 134.3, 139.8, 141.6, 144.7, 148.8, 150.4, 165.9, 167.46, 167.51.



HRMS  $m/z$   $[M + Na]^+$  calcd for  $C_{31}H_{38}N_8NaO_7^+$  657.2756; found 657.2762. IR ( $\nu_{max}$ ,  $cm^{-1}$ ): 3307 w, 2929 w, 2873 w, 3057 w, 1711 w, 1658 m, 1577 w, 1535 w, 1428 w, 1410 w, 1366 w, 1310 w, 1266 w, 1210 w, 1196 w, 1120 m, 1103 m, 1062 m, 1036 w, 983 w, 934 w, 918 w, 862 w, 832 w, 732 s, 702 m, 665 w.  $^1H$ -NMR,  $^{13}C$ -NMR spectra and signal assignments (ESI S-23†).

**Preparation of ligand DIBO-PEG<sub>3</sub>-FAPi.** To a solution of compound 7 (1 equiv., 0.016 mmol, 10 mg) in anhydrous DCM (0.5 mL), were added 4-nitrophenyl chloroformate (2 equiv., 0.032 mmol, 6.5 mg) and anhydrous Py (2 equiv., 0.08 mmol, 6.5  $\mu$ L). The solution was stirred for 16 h at rt. The solvent was evaporated *in vacuo* and the crude product was purified by FCC (DCM/MeOH 10 : 1) to afford intermediate (*S,E*)-2-(2-(2-(2-(4-((3-(4-((2-(2-cyanopyrrolidin-1-yl)-2-oxoethyl)carbamoyl)quinolin-6-yl)acrylamido)methyl)-1*H*-1,2,3-triazol-1-yl)ethoxy)ethoxy)ethoxy)ethyl (4-nitrophenyl) carbonate as a white solid (0.011 mmol, 8.7 mg, 68%). To a solution of this intermediate (1 equiv., 0.063 mmol, 50 mg) in anhydrous DMF (2 mL), were added  $NEt_3$  (3 equiv., 0.19 mmol, 26  $\mu$ L) and DIBO derivative 8 (1.1 equiv., 0.069 mmol, 21 mg). The solution was stirred for 5 h at rt. The solvent was evaporated *in vacuo* and the crude product was purified by FCC (DCM/MeOH 15 : 1) to afford **DIBO-PEG<sub>3</sub>-FAPi** as a white oil (0.063 mmol, 61 mg, quant.).  $^1H$  NMR (400 MHz, acetonitrile- $d_3$ )  $\delta$  2.05–2.24 (4H, m), 2.75 (1H, dd,  $J = 4.0$ , 15.0 Hz), 3.07–3.18 (5H, m), 3.39–3.53 (11H, m), 3.62–3.69 (1H, m), 3.79 (2H, t,  $J = 5.1$  Hz), 4.03 (2H, t,  $J = 4.7$  Hz), 4.20 (2H, dd,  $J = 2.1$ , 5.7 Hz), 4.44 (2H, t,  $J = 5.0$  Hz), 4.51 (2H, d,  $J = 5.8$  Hz), 4.68–4.75 (1H, m), 5.28 (1H, s), 5.92 (1H, s), 6.26 (1H, s), 6.77 (1H, d,  $J = 15.7$  Hz), 7.19 (1H, t,  $J = 5.8$  Hz), 7.24–7.39 (7H, m), 7.44–7.55 (3H, m), 7.63 (1H, d,  $J = 15.7$  Hz), 7.76 (1H, s), 7.86 (1H, dd,  $J = 1.9$ , 8.9 Hz), 8.00 (1H, d,  $J = 8.8$  Hz), 8.54 (1H, d,  $J = 1.9$  Hz), 8.86 (1H, d,  $J = 4.3$  Hz).  $^{13}C$  NMR (101 MHz, acetonitrile- $d_3$ )  $\delta$  25.9, 30.6, 35.8, 41.6, 41.8, 43.0, 46.7, 47.0, 47.7, 50.9, 64.8, 70.0, 70.1, 71.0, 71.1, 71.2, 77.1, 110.7, 113.5, 120.1, 120.6, 121.8, 124.15, 124.20, 124.3, 125.0, 125.5, 126.78, 126.84, 127.1, 128.1, 128.2, 129.27, 129.34, 131.1, 131.2, 135.0, 140.1, 143.2, 145.8, 149.8, 151.8, 152.4, 153.3, 156.8, 166.3, 168.2, 168.5. Two  $C_q$  are not resolved. HRMS  $m/z$   $[M + Na]^+$  calcd for  $C_{51}H_{54}N_{10}NaO_{10}^+$  989.3917; found 989.3920. IR ( $\nu_{max}$ ,  $cm^{-1}$ ): 3319 w, 3061 w, 2922 w, 2878 w, 2159 w, 1716 m, 1657 m, 1521 m, 1443 w, 1431 w, 1288 w, 1260 m, 1121 m, 1105 m, 1053 w, 1037 w, 981 w, 916 w, 860 w, 832 w, 764 m, 733 s, 699 m, 1970 w, 1840 w.  $^1H$ -NMR,  $^{13}C$ -NMR spectra and signal assignments (ESI S-26†).

**Characterization of ligand Biotin-PEG<sub>3</sub>-FAPi.** Analytical UPLC:  $R_t = 1.32$  min.  $^1H$  NMR (400 MHz, chloroform- $d$ )  $\delta$  1.42–1.67 (6H, m), 1.93–2.07 (2H, m), 2.19–2.37 (4H, m), 2.49 (1H, d,  $J = 12.8$  Hz), 2.69 (1H, dd,  $J = 4.9$ , 13.0 Hz), 2.86–2.93 (1H, m), 3.30–3.51 (2H, m), 3.50–3.67 (11H, m), 3.74–3.82 (1H, m), 3.81–3.92 (3H, m), 4.14 (1H, t,  $J = 6.4$  Hz), 4.32 (2H, dd,  $J = 2.3$ , 5.5 Hz), 4.47–4.54 (2H, m), 4.58 (2H, t,  $J = 6.7$  Hz), 4.78–4.85 (1H, m), 5.16 (1H, s), 6.63 (1H, s), 6.75 (1H, d,  $J = 15.7$  Hz), 7.22 (1H, t,  $J = 5.5$  Hz), 7.47 (1H, d,  $J = 4.4$  Hz), 7.67 (1H, d,  $J = 15.6$  Hz), 7.78 (1H, dd,  $J = 1.8$ , 8.9 Hz), 7.91 (1H, s), 8.01–7.97 (1H, m), 8.03 (1H, d,  $J = 8.7$  Hz), 8.44 (1H, s), 8.56 (1H, t,  $J = 5.5$  Hz), 8.87 (1H, d,  $J = 4.3$  Hz).  $^{13}C$  NMR (101 MHz, chloroform- $d$ )  $\delta$  25.4, 25.6, 28.2, 28.4, 29.8, 34.9, 35.8, 39.5, 40.6, 42.4, 46.1, 47.1, 50.4, 55.6, 60.0, 61.9, 69.4, 70.0, 70.3, 70.6, 70.7, 118.3, 119.5, 123.1, 124.0,

124.8, 125.0, 129.6, 130.3, 134.2, 139.8, 141.7, 144.8, 149.0, 150.6, 163.9, 166.2, 168.2, 168.3, 173.6. HRMS  $m/z$   $[M + Na]^+$  calcd for  $C_{41}H_{53}N_{11}NaO_8S^+$  882.3691; found 882.3713. IR ( $\nu_{max}$ ,  $cm^{-1}$ ): 3300 (m), 2925 (m), 2870 (m), 2110 (w), 2099 (w), 1940 (w), 1887 (w), 1801 (w), 1656 (s), 1544 (m), 1440 (m), 1323 (m), 1307 (m), 1265 (m), 1088 (m), 1037 (m), 976 (w), 920 (w), 867 (w), 834 (w), 760 (w). Detailed synthetic procedure,  $^1H$ -NMR,  $^{13}C$ -NMR spectra and signal assignments, UPLC trace (ESI S-13†).

### Functionalization of BFO-PEG NPs

To a suspension of **BFO-PEG NPs** (4 mg) in EtOH (4 mL) was added distilled water (4 mL). A solution of **DIBO-PEG<sub>3</sub>-FAPi** (17.4  $\mu$ mol, 17 mg) in DMF (200  $\mu$ L) was added and the suspension was ultra-sonicated for 16 h at 40 °C. The mixture was then divided into eppendorfs and centrifuged (10 min, 13 000 rpm). The supernatant was discarded and the NPs were resuspended in EtOAc (1 mL) and centrifuged (10 min, 13 000 rpm). The procedure was repeated 3 times. **BFO-PEG-FAPi NPs** were stored in EtOH at a concentration of 1 mg mL<sup>-1</sup>. A sample (10  $\mu$ L) was diluted with distilled water (1 mL) and ultra-sonicated for 30 min, before being analyzed with a Malvern NanoZ instrument to determine mean hydrodynamic diameter by dynamic light scattering and zeta potential.

### Inhibition of hrFAP, hrDPP IV and hrPREP

The human recombinant enzymes were purchased from commercial sources: hrDPP IV and hrPREP (Enzo Life Sciences, Lausen, Switzerland), hrFAP (R&D systems, Abingdon, UK). The enzymatic activities were measured in flat bottom 96-well plates (Costar) containing in each well 0.01  $\mu$ g of the enzymes and 50  $\mu$ M of the appropriate substrates: Z-Gly-Pro-AMC for hrFAP $\alpha$  and hrPOP; H-Gly-Pro-AMC for hrDPP IV (both substrates from Bachem, Vionnaz, Switzerland) diluted in their respective assay buffers (50 mM Tris, 1 M NaCl, 1 mg mL<sup>-1</sup> BSA, pH 7.5, for hrFAP; 50 mM Tris, 1 mg mL<sup>-1</sup> BSA, pH 7.5, for hrPREP; 25 mM Tris and 1 mg mL<sup>-1</sup> BSA, pH 8.0 for hrDPP IV). The enzyme solutions were incubated for 30 min at 37 °C with increasing concentrations (5, 10, 20, 50 and 100 nM) of compounds 5 or **Biotin-PEG<sub>3</sub>-FAPi**. The residual enzymatic activity was determined by measuring fluorescence increase for 60 min at 37 °C in a fluorescence multi-well plate reader ( $\lambda_{ex}/\lambda_{em} = 360/460$  nm, Synergy HT). Experiments were conducted in triplicate wells and repeated twice. The half maximal inhibitory concentrations ( $IC_{50}$ ) were graphically determined and the inhibition constants ( $K_i$ ) were calculated.<sup>49</sup>

### Evaluation of cytocompatibility on MucilAir™-HF tissue samples

Airway cells were obtained from patients undergoing surgical polypectomy. All experimental procedures were explained in full, and all subjects provided informed consent. The study was conducted according to the declaration of Helsinki on biomedical research (Hong Kong amendment, 1989), and received approval from local ethics commission (Commission cantonale d'éthique de la recherche scientifique de Genève [CCER]). Airway epithelia co-cultured with fibroblasts were isolated from a mixture of



human airway cells from 14 different donors (MucilAir™-Pool-HF) and maintained at the air-liquid interface (ALI) in MucilAir™ culture medium (EP04MM), ready-to-use, chemically defined, serum-free (Epithelix Sàrl, Geneva, Switzerland), in 24-well plates with 6.5 mm Transwell® inserts (cat #3470, Corning Incorporated, Tewksbury, USA). **Biotin-PEG<sub>3</sub>-FAPi** was first solubilized in DMSO and diluted to reach targeted concentrations (*i.e.* 10 μM and 100 μM) in MucilAir™ culture medium (EP04MM) with a fixed 1% DMSO for each dilution.

**Trans-epithelial electrical resistance (TEER) measurement.** After addition of 200 μL of MucilAir™ culture medium to the apical compartment of the tissue cultures, resistance was measured across cultures with an EVOMX Volt-Ohm-Meter (World Precision Instruments, Sarasota, US). Resistance values (Ω) were converted to TEER (Ω cm<sup>2</sup>) by using the following formula: TEER (Ω cm<sup>2</sup>) = (resistance value (Ω) – 100(Ω)) × 0.33 (cm<sup>2</sup>), where 100 Ω is the resistance of the membrane and 0.33 cm<sup>2</sup> is the total surface of the epithelium.

**Cytotoxicity measurement.** For the lactate dehydrogenase assay, 100 μL from the basolateral medium was incubated with the reaction mixture of the Cytotoxicity Detection KitPLUS, following manufacturer's instructions (Sigma, Roche; ST Louis, USA). To determine the percentage of cytotoxicity, the following equation was used (*A* = absorbance values): Cytotoxicity (%) = (*A* (exp. value) – *A* (low control)/*A* (high control) – *A* (low control)) × 100. The high control value corresponds to a 10% Triton X-100 treatment applied to the culture for 24 hours.

**Cilia beat frequency measurement.** Cilia beating frequency set-up system consists of three parts: a Sony XCD V60 camera connected to an Olympus BX51 microscope and PCI card. The cilia beating frequency is expressed as Hz. 256 Images were captured at high frequency rate (125 frames per second) at room temperature, cilia beating frequency was then calculated using CiliaX software. It should be pointed out that CBF values may be subject to fluctuations due to parameters such as temperature, mucus viscosity or liquid applied on the apical surface of MucilAir™-HF.

#### Evaluation of the association of BFO-PEG-FAPi NPs with FAP

The protocol refers to the data presented in Fig. 3. Human Fibroblast Activation Protein alpha/FAP Alexa Fluor® 594-conjugated Antibody was purchased from R&D systems (Abingdon, UK). Incubations were carried out in hrFAP buffer (50 mM Tris, 1 M NaCl, 1 mg mL<sup>-1</sup> BSA, pH 7.5) in 1.5 mL Eppendorf® tubes at 37 °C. The following combinations were evaluated in a final volume of 200 μL: (1) anti-FAP antibody (1 : 50 dilution), hrFAP (100 ng) in buffer (control); (2) anti-FAP antibody (1 : 50 dilution), **BFO-PEG-FAPi** NPs (100 μg mL<sup>-1</sup>) in buffer (control); (3) anti-FAP antibody (1 : 50 dilution), hrFAP (100 ng), **BFO-PEG** NPs (100 μg mL<sup>-1</sup>) in buffer; (4) anti-FAP antibody (1 : 50 dilution), hrFAP (100 ng), **BFO-PEG-FAPi** NPs (100 μg mL<sup>-1</sup>) in buffer. The Eppendorf® tubes containing the mixtures were incubated under gentle shaking for 2 h. The mixtures were centrifuged (5 min, 10 000 rpm). The supernatants were collected and transferred to a flat bottom 96-well plate. The intensity of the fluorescent emission of the labelled anti-FAP antibody was measured on a fluorescence multi-well

plate reader ( $\lambda_{\text{ex}}/\lambda_{\text{em}} = 590/645$  nm, Synergy HT). Experiments were conducted in five replicates. Results were analyzed using a student's *t*-test (all comparisons \*\*\**p* < 0.001).

## Conflicts of interest

There are no conflicts to declare.

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## Notes and references

- 1 K. T. Nguyen and Y. Zhao, *Acc. Chem. Res.*, 2015, **48**, 3016.
- 2 G. Chen, I. Roy, C. Yang and P. N. Prasad, *Chem. Rev.*, 2016, **116**, 2826.
- 3 L. Zhang, Y. Li and J. C. Yu, *J. Mater. Chem. B*, 2014, **2**, 452.
- 4 C.-F. Wang, M. P. Sarparanta, E. M. Mäkilä, M. L. K. Hyvönen, P. M. Laakkonen, J. J. Salonen, J. T. Hirvonen, A. J. Airaksinen and H. A. Santos, *Biomaterials*, 2015, **48**, 108.
- 5 C. Argyo, V. Weiss, C. Bräuchle and T. Bein, *Chem. Mater.*, 2014, **26**, 435.
- 6 W. Q. Lim, S. Z. F. Phua, H. V. Xu, S. Sreejith and Y. Zhao, *Nanoscale*, 2016, **8**, 12510.
- 7 P. Kumar, P. Tambe, K. M. Paknikar and V. Gajbhiye, *J. Controlled Release*, 2018, **287**, 35.
- 8 A. Semkina, M. Abakumov, N. Grinenko, A. Abakumov, A. Skorikov, E. Mironova, G. Davydova, A. G. Majouga, N. Nukolova, A. Kabanov and V. Chekhonin, *Colloids Surf., B*, 2015, **136**, 1073.
- 9 J. Chomouka, J. Drbohlavova, D. Huska, V. Adam, R. Kizek and J. Hubalek, *Pharmacol. Res.*, 2010, **62**, 144.
- 10 A. Ghoshal, U. Goswami, A. K. Sahoo, A. Chattopadhyay and S. S. Ghosh, *ACS Biomater. Sci. Eng.*, 2015, **1**, 1256.
- 11 J. Zhu, L. Zheng, S. Wen, Y. Tang, M. Shen, G. Zhang and X. Shi, *Biomaterials*, 2014, **35**, 7635.
- 12 D. Dutta, A. K. Sahoo, A. Chattopadhyay and S. S. Ghosh, *J. Mater. Chem. B*, 2016, **4**, 793.
- 13 M. P. Melancon, M. Zhou and C. Li, *Acc. Chem. Res.*, 2011, **44**, 947.
- 14 A. A. Borran, A. Aghanejad, A. Farajollahi, J. Bara and Y. Omid, *Radiat. Phys. Chem.*, 2018, **152**, 137.
- 15 W. Zhao, A. Li, A. Zhang, Y. Zheng and J. Liu, *ChemMedChem*, 2018, **13**, 2134.
- 16 C. Ding, A. Zhu and Y. Tian, *Acc. Chem. Res.*, 2014, **47**, 20.
- 17 Y. Nakamura, A. Mochida, P. L. Choyke and H. Kobayashi, *Bioconjugate Chem.*, 2016, **27**, 2225.



- 18 J. E. Rosen, L. Chan, D.-B. Shieh and F. X. Gu, *Nanomedicine*, 2012, **8**, 275.
- 19 R. Bazak, M. Hourri, S. El Achy, S. Kamel and T. Refaat, *J. Cancer Res. Clin. Oncol.*, 2016, **141**, 769.
- 20 J. Paulsson and P. Micke, *Semin. Cancer Biol.*, 2014, **25**, 61.
- 21 M. Li, M. Li, T. Yin, H. Shi, Y. Wen, B. Zhang, M. Chen, G. Xu, K. Ren and Y. Wei, *Mol. Med. Rep.*, 2016, **13**, 2476.
- 22 C. Fernandes, D. Soares and C. Mayur Yergeri, *Front. Pharmacol.*, 2018, **9**, 1230.
- 23 T. Cui, X. Li, Y. Shu, X. Huang, Y. Wang and W. Zhang, *Int. J. Pharm.*, 2018, **552**, 16.
- 24 E. Sverdlov, *BioEssays*, 2018, **40**, 1800079.
- 25 L. Juillerat-Jeanneret and S. Gerber-Lemaire, *Mini-Rev. Med. Chem.*, 2009, **2**, 215.
- 26 D. T. Fearon, *Cancer Immunol. Res.*, 2014, **2**, 187.
- 27 F. Zi, J. He, D. He, Y. Li, L. Yang and Z. Cai, *Mol. Med. Rep.*, 2015, **11**, 3203.
- 28 P. Busek, R. Mateu, M. Zubal, L. Kotackova and A. Sedo, *Front. Biosci.*, 2018, **23**, 1933.
- 29 A. Loktev, T. Lindner, W. Mier, J. Debus, A. Altmann, D. Jäger, F. Giesel, C. Kratochwil, P. Barthe, C. Roumestand and U. A. Haberkorn, *J. Nucl. Med.*, 2018, **59**, 1423.
- 30 K. W. Jackson, V. J. Christiansen, V. R. Yadav, R. Silasi-Mansat, F. Lupu, V. Awasthi, R. R. Zhang and P. A. McKee, *Neoplasia*, 2015, **17**, 43.
- 31 L. Bonacina, *Mol. Pharmaceutics*, 2013, **7**, 205.
- 32 W. P. Dempsey, S. E. Fraser and P. Pantazis, *BioEssays*, 2012, **34**, 351.
- 33 Y. Pu and D. Psaltis, *Appl. Opt.*, 2013, **52**, 567.
- 34 L. Bonacina, Y. Mugnier, F. Courvoisier, R. Le Dantec, J. Extermann, Y. Lambert, V. Boutou, C. Galez and J.-P. Wolf, *Appl. Phys. B*, 2007, **87**, 399.
- 35 J. Extermann, L. Bonacina, E. Cuna, C. Kasparian, Y. Mugnier, T. Feurer and J.-P. Wolf, *Opt. Express*, 2009, **17**, 15342.
- 36 L. L. Xuan, C. Zhou, A. Slablab, D. Chauvat, C. Tard, S. Perruchas, T. Gacoin, P. Villela and J. F. Roch, *Small*, 2008, **4**, 1332.
- 37 D. Staedler, T. Magouroux, R. Hadli, C. Joulaud, J. Extermann, S. Schwung, S. Passemard, C. Kasparian, G. Clarke, M. Gerrmann, R. Le Dantec, Y. Mugnier, D. Rytz, D. Ciepiewski, C. Galez, S. Gerber-Lemaire, L. Juillerat-Jeanneret, L. Bonacina and J.-P. Wolf, *ACS Nano*, 2012, **6**, 2542.
- 38 S. Schwung, A. Rogov, G. Clarke, C. Joulaud, T. Magouroux, D. Staedler, S. Passemard, T. Jüstel, L. Badie, C. Galez, J.-P. Wolf, Y. Volkov, A. Prina-Mello, S. Gerber-Lemaire, D. Rytz, Y. Mugnier, L. Bonacina and R. Le Dantec, *J. Appl. Phys.*, 2014, **116**, 114306.
- 39 D. Staedler, S. Passemard, T. Magouroux, A. Rogov, C. M. Maguire, B. M. Mohamed, S. Schwung, D. Rytz, T. Jüstel, S. Hwu, Y. Mugnier, R. Le Dantec, Y. Volkov, S. Gerber-Lemaire, A. Prina-Mello, L. Bonacina and J.-P. Wolf, *Nanomedicine*, 2015, **11**, 815.
- 40 L. Dubreil, I. Leroux, M. Ledevin, C. Schleder, L. Lagalice, C. Lovo, R. Fleurisson, S. Passemard, V. Kilin, S. Gerber-Lemaire, A. M. Colle, L. Bonacina and K. Rouger, *ACS Nano*, 2017, **11**, 6672.
- 41 A. Rogov, M. Irondelle, F. Ramos-Gomes, J. Bode, D. Staedler, S. Passemard, S. Courvoisier, Y. Yamamoto, F. Waharte, D. Ciepiewski, P. Rideau, S. Gerber-Lemaire, F. Alves, J. Salamero, L. Bonacina and J.-P. Wolf, *ACS Photonics*, 2015, **2**, 1416.
- 42 S. Passemard, D. Staedler, G. Sonego, T. Magouroux, G. S. Schneiter, L. Juillerat-Jeanneret, L. Bonacina and S. Gerber-Lemaire, *J. Nanopart. Res.*, 2015, **17**, 414.
- 43 R. Azimeh, X. Wensheng, L. Zhao, S. Wang, Y. Liu, Z. Wu, J. Wang and F. Si-She, *J. Biomed. Nanotechnol.*, 2018, **14**, 1159.
- 44 C. Yang, Y. Chen, W. Guo, Y. Gao, C. Song, Q. Zhang, N. Zheng, X. Han and C. Guo, *Adv. Funct. Mater.*, 2018, **28**, 1706827.
- 45 K. Jansen, L. Heirbaut, J. D. Cheng, J. Joossens, O. Ryabtsova, P. Cos, L. Maes, A.-M. Lambeir, I. De Meester, K. Augustyns and P. Van der Veken, *ACS Med. Chem. Lett.*, 2013, **4**, 491.
- 46 F. Kikkawa, H. Kajiyama, K. Shibata, K. Ino, S. Nomura and S. Mitztani, *Biochim. Biophys. Acta, Proteins Proteomics*, 2005, **1751**, 45.
- 47 H. C. Cheng, M. Abdel-Ghany, S. Zhang and B. U. Pauli, *Clin. Exp. Metastasis*, 1999, **17**, 609.
- 48 J. A. García-Horsman, P. T. Männistö and J. I. Venäläinen, *Neuropeptides*, 2007, **41**, 1.
- 49 J. Lawandi, S. Gerber-Lemaire, L. Juillerat-Jeanneret and N. Moitessier, *J. Med. Chem.*, 2010, **53**, 3423.
- 50 C. Y. Edosada, C. Quan, C. Wiesmann, T. Tran, D. Sutherlin, M. Reynolds, J. M. Elliott, H. Raab, W. Fairbrother and B. B. Wolf, *J. Biol. Chem.*, 2006, **281**, 7437.
- 51 S. E. Poplawski, J. H. Lai, Y. Li, Z. Jin, Y. Liu, W. Wu, Y. Wu, Y. Zhou, J. L. Sudmeier, D. G. Sanford and W. W. Bachovchin, *J. Med. Chem.*, 2013, **56**, 3467.
- 52 T.-Y. Tsai, T.-K. Yeh, X. Chen, T. Hsu, Y.-C. Jao, C.-H. Huang, J.-S. Song, Y.-C. Huang, C. H. Chien, J. H. Chiu, S.-C. Yen, H. K. Tang, Y.-S. Chao and W.-T. Jiaang, *J. Med. Chem.*, 2010, **53**, 6572.
- 53 J. N. Sangshetti, A. S. Zambare, I. Gonjari and D. B. Shinde, *Mini-Rev. Org. Chem.*, 2014, **11**, 225.
- 54 H. Zhu, R. F. Yang, L. H. Yun and J. Li, *Chin. Chem. Lett.*, 2010, **21**, 35.
- 55 I. P. Beletskaya and A. V. Cheprakov, *Chem. Rev.*, 2000, **100**, 3009.
- 56 K. Balogh Sivars, U. Sivars, E. Hornberg, H. Zhang, L. Brändén, R. Bonfante, S. Huang, S. Constant, I. Robinson, C. J. Betts and P. M. Åberg, *Toxicol. Sci.*, 2018, **162**, 301.
- 57 R. Z. Cer, U. Mudunuri, R. Stephens and F. J. Lebeda, *Nucleic Acids Res.*, 2009, **37**(suppl. 2), W441.
- 58 E. Blanco, H. Shen and M. Ferrari, *Nat. Biotechnol.*, 2015, **33**, 941.
- 59 R. T. Aimes, A. Zijlstra, J. D. Hooper, S. M. Ogbourne, M. L. Sit, S. Fuchs, D. C. Gotley, J. P. Quigley and T. M. Antalis, *Thromb. Haemostasis*, 2003, **89**, 561.
- 60 J. A. McIntyre and J. Castañer, *Drugs Future*, 2004, **29**, 882.



## Paper

- 61 J. Nemunaitis, S. J. Vukelja, D. Richards, C. Cunningham, N. Senzer, J. Nugent, H. Duncan, B. Jones, E. Haltom and M. J. Uprichard, *Cancer Invest.*, 2006, **24**, 553.
- 62 K. Narra, S. R. Mullins, H. O. Lee, R. Stzremkowski-Brun, K. Magalong, V. J. Christansen, P. A. McKee, B. Egleston, S. J. Cohen, L. M. Weiner, N. J. Meropol and J. D. Cheng, *Cancer Biol. Ther.*, 2007, **7**, 1691.
- 63 P. Dvořáková, P. Bušek, T. Knedlik, J. Schimer, T. Etrych, L. Kostka, L. Stollinová Šromová, V. Šubr, P. Šácha, A. Šedo and J. Konvalinka, *J. Med. Chem.*, 2017, **60**, 8385–8393.

