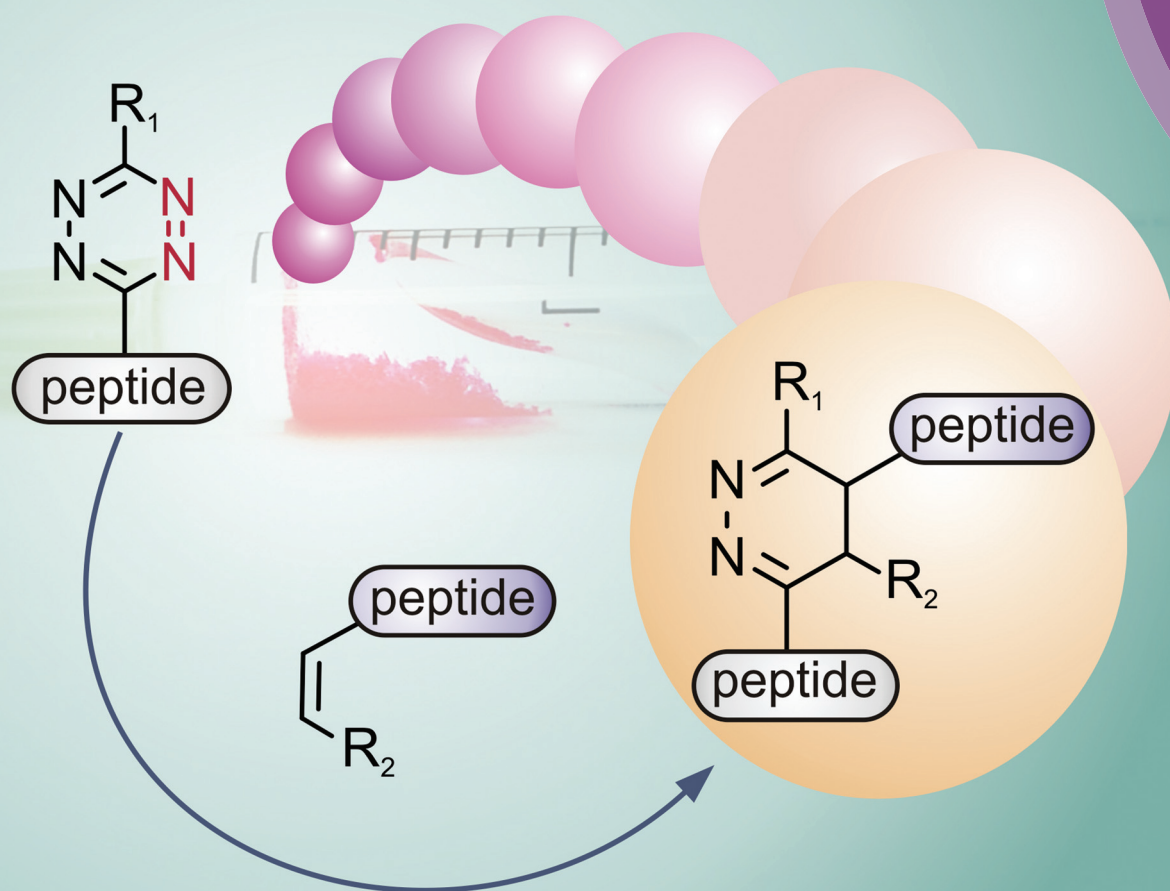


Organic & Biomolecular Chemistry

www.rsc.org/obc



ISSN 1477-0520



PAPER

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175 YEARS



Cite this: *Org. Biomol. Chem.*, 2016, **14**, 4809

On-resin Diels–Alder reaction with inverse electron demand: an efficient ligation method for complex peptides with a varying spacer to optimize cell adhesion†

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Solid phase peptide synthesis (SPPS) is the method of choice to produce peptides. Several protecting groups enable specific modifications. However, complex peptide conjugates usually require a rather demanding conjugation strategy, which is mostly performed in solution. Herein, an efficient strategy is described using an on-resin Diels–Alder reaction with inverse electron demand (DAR_{inv}). This method is compatible with the standard Fmoc/*t*Bu strategy and is easy to monitor. As a proof of concept a titanium binding peptide was modified with a cyclic cell binding peptide (RGD) by DAR_{inv} on a solid support applying different tetrazines and alkenes. The generated bulky DAR_{inv} linkers were employed to act as the required spacer for RGD mediated cell adhesion on titanium. *In vitro* studies demonstrated improved cell spreading on DAR_{inv}-conjugated peptides and revealed, in combination with molecular dynamics-simulation, new insights into the design of spacers between the RGD peptide and the surface. Performing the DAR_{inv} on resin expands the toolbox of SPPS to produce complex peptide conjugates under mild, catalyst free conditions with reduced purification steps. The resulting conjugate can be effectively exploited to promote cell adhesion on biomaterials.

Received 7th February 2016,
Accepted 19th April 2016

DOI: 10.1039/c6ob00314a

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Introduction

Peptide conjugates are perfect tools to selectively address various targets in medical applications and analytical investigations. Solid phase peptide synthesis (SPPS) provides a variety of orthogonal protecting groups, which facilitate the modification of peptides with fluorophores, drugs, carbohydrates, fatty acids or reporter molecules.^{1–3} Nevertheless, the efficient synthesis of multifunctional peptides with several complex bioactive moieties such as cyclic peptides is still challenging. Chemoselective click reactions offer the possibility to selectively ligate biomolecules without the use of further protecting groups in benign solvents resulting in high yields.^{4,5} Therefore click chemistry is a convenient method to upgrade traditional SPPS to obtain multifunctional peptides.^{6–8} It was shown that

copper(i) catalyzed azide–alkyne cycloaddition (CuAAC), oxime ligation and recently the Staudinger–phosphite reaction are applicable for on resin synthesis to obtain branched and functionalized molecules.^{9–12} Click reactions on a solid support reduce purification steps, enable automation or protect sensitive amino acids like L-3,4-dihydroxyphenylalanine (DOPA) from oxidation.¹⁰ Another bioorthogonal cycloaddition is the Diels–Alder reaction with inverse electron demand (DAR_{inv}) that can be performed in water without catalysts or hazardous sideproducts and reagents.¹³ This irreversible reaction between an alkene (dienophile) and a tetrazine (diene) has been successfully used to synthesize peptide conjugates by us and others for various medical applications.^{14–17} However, solubility problems or instability of tetrazines can lead to difficulties.^{17,18} An on-resin approach can overcome these limitations and decrease purification and cleavage steps, thus increasing the yield. DAR_{inv}-conjugates mostly consist of bulky linker-units between two conjugated molecules with different distances and flexibilities depending on the applied dienes and dienophiles. This could be beneficial for several *in vitro* investigations where distances are required to enable binding of ligands to proteins or reporter molecules.^{19–21} RGD-peptides bind to receptors in the cell-membrane and thus can mediate

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† Electronic supplementary information (ESI) available. See DOI: 10.1039/c6ob00314a



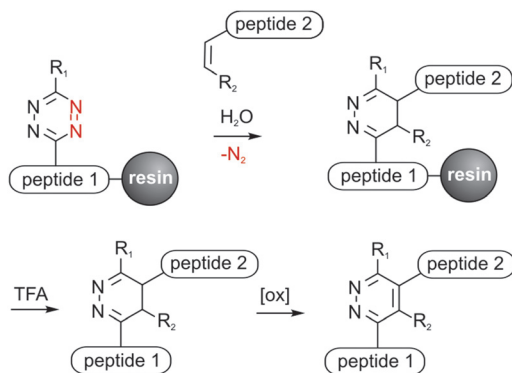


Fig. 1 General approach of the DAR_{inv} on resin. The hydrophobic tetrazine remains on the solid support to avoid solubility problems. After incubation with the dienophile, N_2 is released and the desired product is formed. The conjugate can be further modified on resin or cleaved off by TFA-treatment. The resulting mixture of the isomers can be oxidized.

cell-adhesion on surfaces to improve healing and osseointegration of *e.g.* titanium implants.^{22,23} It could be shown that a minimum distance between the RGD-peptide and the surface is crucial for successful cell adhesion.²⁴ Hence, DAR_{inv} -conjugated RGD-peptides present an efficient approach to promote cell attachment on surfaces such as titanium.

Herein, it is demonstrated that the DAR_{inv} can be expanded to exploit polymer-bound synthesis compatible with standard Fmoc/*t*Bu-SPPS (Fig. 1). The applicability of two tetrazines and four different dienophiles was investigated by ligating a cell-adhesive- and a titanium-binding-peptide to the solid support. The obtained products with a varying linker length were then applied to improve cell adhesion on coated titanium. Moreover, molecular dynamics (MD)-simulation was used to assess the DAR_{inv} -linker conformation and thus discuss the cell response in theory.

Experimental

Synthesis of TBP (1) and TBP 2

Titanium binding peptides (TBP (1) and TBP 2) were manually synthesized on a TentaGel S RAM (Iris Biotech) resin using the Fmoc/*t*Bu strategy. Deprotection of α -amino groups was performed twice with 30% (v/v) piperidine (Sigma Aldrich) in dimethylformamide (DMF, Biosolve) for 10 min. The peptide was elongated using 2 eq. hydroxybenzotriazole (HOBt, Novabiochem) and diisopropylcarbodiimide (DIC, IrisBiotech) in DMF and 2 eq. of Fmoc- β -Ala-OH, Fmoc-Pra-OH (Iris Biotech), Fmoc-DOPA(acetone)-OH (Novabiochem), Fmoc-Lys(Mtt)-OH (Bachem) or Fmoc-Lys(Dde)-OH (Iris Biotech), Fmoc-Cys(Trt)-OH or Boc-Cys(Trt)-OH (Iris Biotech, OrPEGen OPC) and Boc-Ser(*t*Bu)-OH (Iris Biotech). For activation of Fmoc-NH-(PEG)₂-COOH (13 atoms, Novabiochem) 1.5 eq. 1-[bis-(dimethylamino)methylene]-1*H*-1,2,3-triazolo[4,5-*b*]pyridinium 3-oxid hexafluorophosphate (HATU, Novabiochem) and equimolar amounts of *N,N*-diisopropylethylamine (DIPEA, Roth)

were used. As the Lys-side chain protecting group, 4-methyltrityl (Mtt), or *N*-(1-(4,4-dimethyl-2,6-dioxocyclohexylidene) ethyl) (Dde) was used. Selective cleavage of Mtt was performed with 2% trifluoroacetic acid (TFA, Sigma Aldrich) and 3% triisopropylsilane (TIS, Sigma Aldrich) in dichloromethane (DCM, Biosolve) (v/v/v) for 2 min and was repeated 30 times. Cleaving Dde was realized with 2% hydrazine (Sigma Aldrich) in DMF for 10 min. This step was repeated 12 times. Subsequently, the dienes **2b** (4-(6-(pyrimidin-2-yl)-1,2,4,5-tetrazin-3-yl)benzoic acid, synthesized according to Pipkorn *et al.*) and **2a** (5-[4-(1,2,4,5-tetrazin-3-yl) benzyl-amino]-5-oxopentanoic acid (Sigma Aldrich)) were coupled in 2 fold excess with HOBt/DIC for 16 h in DMF.¹³

Synthesis of c[RGDfk(Reppe)] (3), c[RGDfk(Alloc)] (4), c[RGDfE(Allyl)] (5) and c[RGDfk(maleimide)] (6)

The peptide was elongated on an acid labile 2-chlorotrityl-resin (Novabiochem) to obtain Fmoc-D-Phe-Lys(Dde)-Arg(Pbf)-Gly-Asp(*t*Bu)-resin. After Fmoc removal, the N-terminus was protected with triphenylmethyl chloride by DIPEA in DMF. Subsequently, Dde was cleaved from the Lys-side chain, as described, and modified with either the Reppe-dienophile (synthesized according to Pipkorn *et al.*) or 6-maleimidohexanoic acid by HOBt/DIC activation.¹³ After cleavage of the protected peptide from the resin with glacial acetic acid and 2,2,2-trifluoroethanol in DCM (1 : 1 : 8, v/v/v), the linear peptide was cyclized with HOBt/DIC in DCM for 16 h. Next, DCM was evaporated and the peptide was fully deprotected with TFA and a scavenger followed by purification. The peptides c[RGDfK(Alloc)] and c[RGDfE(Allyl)] were synthesized by the same protocol (Fmoc-D-Phe-Lys(Alloc)-Arg(Pbf)-Gly-Asp(*t*Bu)-resin, Fmoc-D-Phe-Glu(Allyl)-Arg(Pbf)-Gly-Asp(*t*Bu)-resin), however without side chain modification and trityl-protection.

Diels–Alder reaction with inverse electron demand (DAR_{inv}) on resin to yield peptides 8a–c

TentaGel resin loaded with **1a** and respectively **1b** was swollen in H_2O . Then the resin was incubated with 1.5 eq. of a 30 mM aqueous solution of peptides 3–6 at room temperature on a shaker in an open reaction vessel (N_2 release). When the reaction was complete, the resin was washed with water, DMF and DCM. If the N-terminus was protected by Fmoc, a deprotection step with piperidine, as stated above, followed. The peptide was cleaved from the resin and isolated as described.

Monitoring of the Diels–Alder reaction with inverse electron demand (DAR_{inv}) on resin by photometry

Diene modified resin (approximately 1 μmol) was transferred into a 96-well plate and swollen in water. Subsequently, the water was removed and the dissolved dienophile (30 mM; 100 μL) was added. An absorption scan from 430–600 nm of the diene modified resin was carried out with a plate reader (Tecan, Infinite® 200 PRO series). If the absorption maximum at 540 nm was diminished, the reaction was complete (**8a**: 5 h, **8b**: 2 h, **8c**: 7 d).



Thiol-maleimide Michael addition to yield peptide 8d

Peptide **8d** was synthesized by ligation of TBP 2 (sequence: Ser-PEG-DOPA-Cys-DOPA-PEG-Pra-β-Ala) and c[RGDfK(maleimide)] (**6**) in H₂O/*t*BuOH for 20 h. The peptide was purified as described.

Cleavage of peptides from the resin and purification

Final and sample cleavage of the peptides was performed with TFA/scavenger (9 : 1 v/v) by shaking for 2 h at RT. Scavenger mixtures were used as follows: TIS/H₂O (1 : 1 v/v) for peptide **6** and TBP 2, thioanisole/thiocresol (1 : 1 v/v) and thioanisole/1,2-ethanedithiol (7 : 3 v/v) for **3–5**, conjugates **8a,b** and H₂O for peptides **1a,b** and **8c**. Peptides were precipitated and washed with diethyl ether. The dissolved peptide was analyzed and lyophilized. Isolation of the obtained peptides was carried out by RP-HPLC on a Phenomenex Jupiter Proteo column (90 Å/4 μm, 22 mm × 250 mm) using linear gradients of eluent B in eluent A (A: 0.1% TFA in H₂O, B: 0.08% TFA in ACN). The identity and purity of the isolated products was verified by MALDI-ToF (Bruker Daltonics) and ESI-HCT (high-capacity ion trap, Bruker Daltonics) mass spectrometry and analytical RP-HPLC (Varian VariTide RPC (200 Å, 6 μm), Phenomenex Jupiter Proteo (90 Å/4 μm and 300 Å, 5 μm) and Grace Vydac (300 Å, 5 μm)) with linear gradients of eluents A and B. The detected *m/z* signals were in agreement with the calculated molecular weights and the peptide products were obtained with a purity of ≥90% (Table 1).

Cellular assays

SaOS-2 cells (Sarcoma osteogenic, kindly provided by Prof. Scharnweber, Dresden, Germany) were cultured under a humidified atmosphere at 37 °C and 5% CO₂ in McCoy's 5A containing 15% heat inactivated fetal calf serum (FCS), 1% (v/v) glutamine and 1% (v/v) penicillin/streptomycin (by PAA, Lonza and Biochrome).

Titanium foil (Sigma Aldrich, thickness: 0.127 mm) was cut into round pieces and etched in H₂SO₄ (30%)/H₂O₂ (1 : 1, v/v) for 7 min. The slides were then washed with water and PBS buffer (Dulbecco's phosphate) and sterilized in 70% ethanol/water (v/v) under ultrasonic irradiation for 15 min. After

washing twice with PBS, the Ti-plates were incubated overnight at room temperature with a 1 μM solution of peptides dissolved in PBS as well as PBS (untreated Ti) and fibronectin (25 μg mL⁻¹ in PBS) as controls.

To investigate the initial cell behavior to the synthesized peptide coatings, cells were resuspended in media without FCS (to prevent adsorption of serum proteins) and seeded on coated and washed Ti-plates. After adhesion for 6 h, the plates were washed twice with PBS and subsequently the cells were fixed with 4% paraformaldehyde in PBS for 30 min. The plates were washed with PBS and the cells were permeabilized with 1% triton X-100 in PBS for 1 min. Next, the slides were washed and cells were stained with phalloidin-tetramethylrhodamine-isothiocyanate (phalloidin-TRITC, Sigma Aldrich) and HOECHST 33342 (Sigma Aldrich). Fluorescence microscopy (Axio Observer microscope, Zeiss) was performed with mounted Ti-plates on glass slides. For each plate, three representing pictures were taken at 20 fold magnification. The average cell area was determined by manually outlining each cell on one representing picture of each triplicate (using the Software Axio Vision 4.8, Zeiss). Data were analyzed from at least 3 independent experiments and presented as mean ± SEM.

Molecular dynamics of the linker

Molecular dynamics simulations were carried out using the Gromacs-4.6 package.²⁵ The simulation system consisted of the respective linker with a C-terminal amidated and N-terminal acetylated lysine on the RGD-peptide side and a C-terminal amidated and N-terminal acetylated lysine/cysteine on the titanium binding site. The linker was parameterized with antechamber²⁶ and acpype²⁷ for the GAFF force field. The charges were derived with the antechamber sqm method. The simulations were carried out under periodic boundary conditions in TIP3P water with 150 mM NaCl at 300 K for 50 ns after equilibration with NVT and NPT ensemble for 50 ps each. The trajectories were analyzed with the *g_dist* tool by measuring the C_α-C_α distance and plotting the values in a histogram with a bin size of 0.5 Å.

Table 1 Analytical data of purified peptides

Entry	Sequence	<i>M</i> [Da]	[<i>M</i> + H] ⁺	Elution ^a (%ACN)	Purity
3	c[RGDfK(Reppe)]	915.5	916.5	44 ^b	>95
4	c[RGDfK(Alloc)]	687.3	688.0	40	>95
5	c[RGDfE(Allyl)]	644.3	645.0	40	>95
6	c[RGDfK(maleimide)]	796.4	797.5	37	>95
8a	C-PEG-Y*-K(2a-c[RGDfK(Reppe)])-Y*-PEG-G*-βA-NH ₂	2349.1	2350.1	36	>90
8b^d	C-PEG-Y*-K(2b-c[RGDfK(Reppe)])-Y*-PEG-G*-βA-NH ₂	2326.1	2327.0	41 ^c	>90
8c	C-PEG-Y*-K(2a-c[RGDfK(maleimide)])-Y*-PEG-G*-βA-NH ₂	2230.0	2231.0	n.i.	n.i.
8d	S-PEG-Y*-C(c[RGDfK(maleimide)])-Y*-PEG-G*-βA-NH ₂	1933.9	1935.0	33	>95

Peptides were analyzed by MALDI-ToF MS (Bruker, Daltonics) and RP-HPLC. The peptide purity was evaluated by two different HPLC systems using different columns and gradients of eluent B (0.08 % TFA in ACN) in eluent A (0.1 % TFA in H₂O) on a Phenomenex Jupiter Proteo (90 Å, 5 μm)^a, Phenomenex Jupiter Proteo (300 Å, 4 μm)^b and VariTide (200 Å, 6 μm)^c; ^doxidized product isolated, PEG (polyethylene glycol), Y* (*l*-3,4-dihydroxyphenylalanine), G* (propargylglycine), βA (beta alanine), n.i. = not isolated.



Results and discussion

Peptide modification with tetrazines

The titanium binding peptide (TBP, **1**) was synthesized on a TentaGel resin to enable the DAR_{inv} on the resin in aqueous solution. The sequence was built up of spacer units such as polyethylene glycol (PEG) and β -Ala. DOPA (L-3,4-dihydroxyphenylalanine), which shows high affinity and stability to titanium and other surfaces, was used to anchor the peptide to the bio-material.²⁸ Functional groups were introduced by using Lys, Pra (L-propargylglycine) and Cys to allow the multifunctional modifications of TBP.²⁹ Two dienes (**2a,b**) were coupled to TBP after selective side chain deprotection of Lys (Fig. 2). The result of the reaction was studied by RP-HPLC and MALDI-ToF-MS analysis (Fig. 4d). The tetrazine stability in **1a** and **1b** was tested in solution and on resin over 7 d in an aqueous environment and yielded low fragmentation (see the ESI[†]). As a side reaction, the undesired DAR_{inv} between the alkyne functionality of Pra and the applied tetrazines was observed, mainly for peptide **1a** in solution. On resin, this reaction occurred only to a small extent. Cleavage of **1b** from the resin resulted in fragmentation induced by water, which was slightly increased after incubation of the resin in water for 7 d.³⁰ Nevertheless, possible degradation of tetrazines by the cleavage of the peptide from the resin is circumvented by the DAR_{inv} on resin.

Peptide modification with dienophiles

Different dienophiles were introduced to the cyclic cell binding peptide c[RGDFK]. The peptide was synthesized on an acid labile chlorotrityl resin (Fig. 3). The modification with different dienophiles was either realized by the incorporation of alkene containing amino acids (**4**, **5**) or by Lys-side chain modification (**3**, **6**). The dienophile functionalized peptides were cleaved under mild acidic conditions to retain side chain protecting groups. Subsequently, the selective head to tail

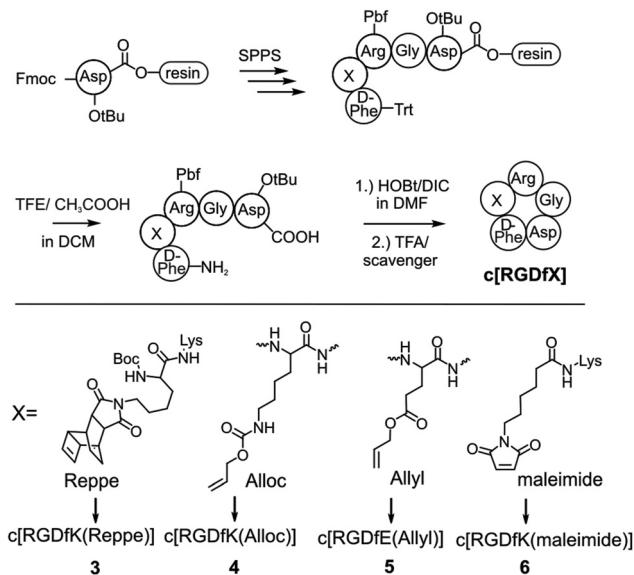


Fig. 3 Synthesis of dienophile modified cyclic RGD. The peptide was elongated on an acid labile chlorotrityl-resin. To yield **4** and **5**, Fmoc-Lys(Alloc)-OH and Fmoc-Glu(Allyl)-OH respectively were incorporated into the peptide. Peptides **3** and **6** were obtained by incorporation of lysine and subsequent modification with either Reppe-anhydride (Reppe) or 6-maleimidehexanoic acid (maleimide).

cyclization was performed in solution. Peptides were finally purified by RP-HPLC after complete deprotection by TFA. The Reppe-anhydride, which has already been used in previous studies, was coupled at the Lys-side chain of the c[RGDFK]-peptide to yield compound **3**.^{31–33} To facilitate the introduction of dienophiles during SPPS, we used terminal alkenes, which are present in common protecting groups like Alloc and Allyl. A 6-maleimidohexanoic acid modified peptide (**6**) was synthesized to explore the feasibility of this easily accessible dienophile for the DAR_{inv} on resin (Fig. 3).

Ligation of peptides by DAR_{inv} on resin

To perform the DAR_{inv} on the solid phase, the dienophile **3** was dissolved in H_2O ($c = 30 \text{ mM}$) and added to the resin bound peptides **1a** and **b**. A disadvantage of reactions on the solid phase is the often complicated monitoring of the reaction. However, herein we demonstrate a direct photometric measurement of the reaction process on resin. The DAR_{inv} was monitored by measuring the specific absorbance of the tetrazines at 540 nm directly on the resin by photometry (Fig. 4b and c). The cycloaddition with peptide **1b** was complete after 2 h. Tetrazine **2a** resulted in a slower reaction with full conversion after 5 h (see the ESI[†]), which can be explained by the presence of stronger electron withdrawing groups in **2b** compared to **2a**. Higher reactivity of tetrazines often results in lower stability.³⁴ During the DAR_{inv} on resin with peptide **3** we could not detect the differences in diene stability. However, for reactions with longer incubation times, as for peptide **6**, lower stability of tetrazine **2b** compared to **2a** was observed (see the ESI[†]). Furthermore, combining the DAR_{inv} with additional

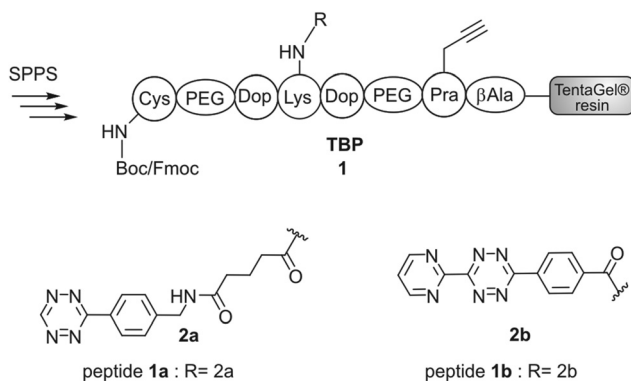


Fig. 2 Tetrazine-modified peptides. Modification with dienes of the scaffold peptide **1** on a PEG-based resin by standard HOBt/DIC activation. TBP (titanium binding peptide): DOPA (L-3,4-dihydroxyphenylalanine) for Ti-binding, Pra (L-propargylglycine) and Cys for further modification, β Ala (L- β -alanine) and PEG (polyethylene glycol) are used as the spacer to facilitate further modifications by additional bioorthogonal reactions.



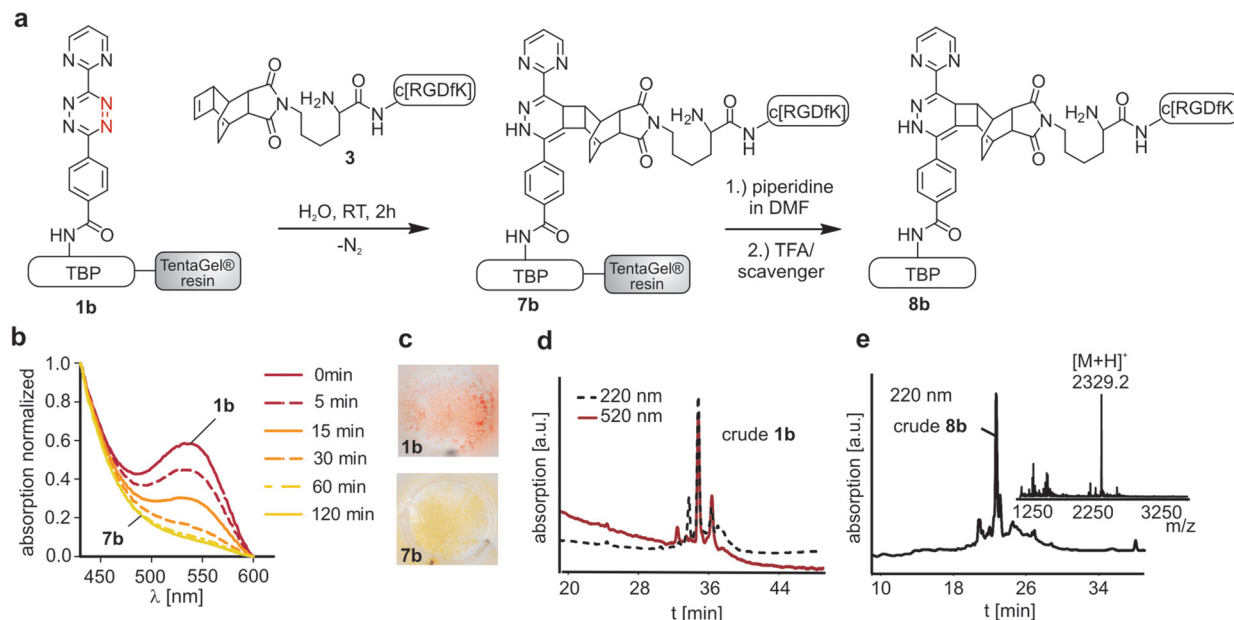


Fig. 4 DAR_{inv} on resin. (a) Reaction scheme of the DAR_{inv}. The tetrazine bearing resin was incubated with dienophile **3** dissolved in H₂O. After the reaction is complete, Fmoc can be cleaved from the N-terminus with piperidine in DMF. The peptide can be further modified or cleaved off from the resin by TFA and a scavenger. (b) Absorption scan of resin bound tetrazine during the DAR_{inv} to monitor the progress of the reaction (c) Photographs of modified TentaGel resin before (**1b**) and after the DAR_{inv} (**7b**). (d) Crude RP-HPLC chromatogram of tetrazine modified peptide **1b**. (e) Crude RP-HPLC chromatogram and MALDI-ToF MS of the DAR_{inv}-product **8b**.

reactions as the CuAAC lead to stability problems for compound **8b** originated from tetrazine **2b** as well (see the ESI†). Contrarily, **8a** was stable under the CuAAC on resin and was recently used to synthesize a multifunctional cell adhesive titanium coating.²⁹ This suggests that the choice of the tetrazine can strongly influence the stability of the resulting DAR_{inv}-product as shown also for metabolic degradation.¹⁴ After completion of the reaction, the dienophile containing peptide **3**, which was used in excess, was fully recovered by lyophilization. Cleavage from the resin and subsequent analysis showed the generation of the desired dihydropyridazine **8b** (Fig. 4e). Notably, the DAR_{inv}-product was stable under the conditions of the Fmoc-cleavage with piperidine in DMF, which underlines the compatibility with Fmoc/*t*Bu-based-SPPS.

Conjugate **8b** was partially oxidized to the corresponding pyridazine (Fig. 1). Complete oxidation can be achieved by incubation with isoamyl nitrite and acetic acid, which can be performed on resin and in solution (Fig. 5).³⁵

Terminal alkenes were successfully applied to react with tetrazines by a DAR_{inv}.^{36,37} Incubation of the tetrazine containing peptide **1a** and dienophile **4** or **5** respectively could show the formation of the desired DAR_{inv} product, which was however cleaved after cycloaddition (for the hypothesized mechanism see the ESI†). The reaction of these terminal alkene bearing peptides with tetrazine **1b** resulted only in low conversion of around 10% as a result of low reactivity and stability of the tetrazine (see the ESI†). Alternative tetrazines could be applied to overcome this problem to establish an easy introduction of dienophiles in SPPS. Moreover, the suggested mechanism could

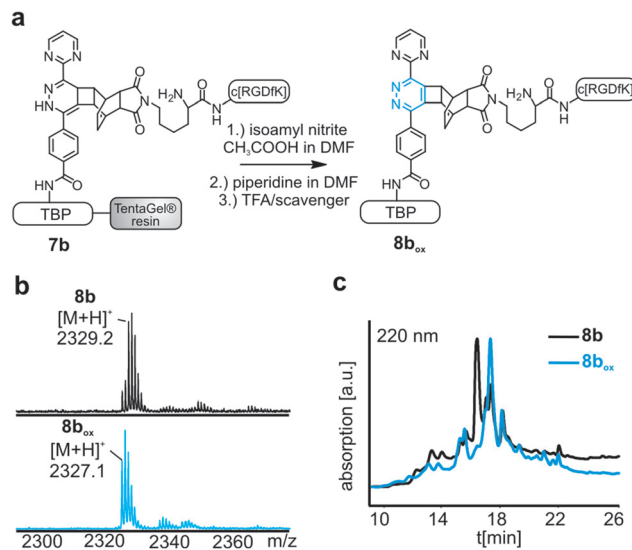


Fig. 5 Oxidation of the DAR_{inv} product (a) oxidation of **8b** on resin (**7b**) by isoamyl nitrite. (b) MALDI-ToF-analysis of the partially oxidized starting material **8b** and the oxidized product **8b_{ox}**. (c) RP-HPLC detected at 220 nm shows a shift of the dihydropyridazine completely converted to the corresponding pyridazine.

be probably applied as a decaging method as described recently for proteins or drugs, since the active chemotherapeutic RGD-peptide is released upon fragmentation.^{38,39} Maleimides are classical dienophiles in normal Diels–Alder reactions.⁴⁰ However, the DAR_{inv} between peptide **1a** and the



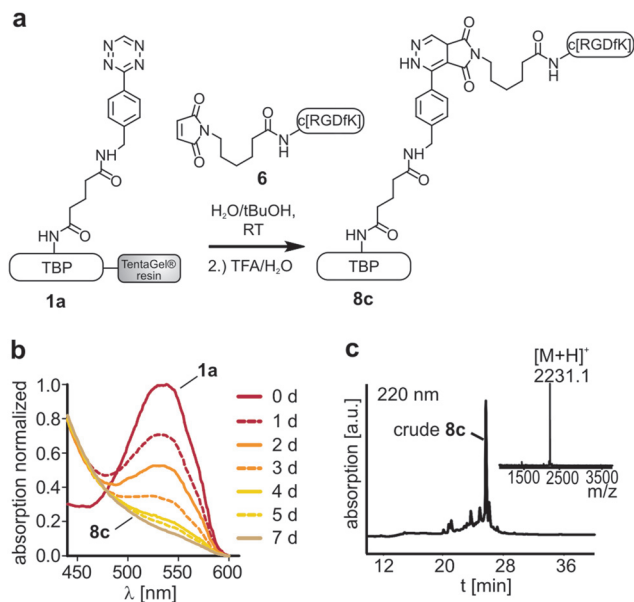


Fig. 6 DAR_{inv} on resin with c[RGDFK(maleimide)]. (a) Reaction scheme of the DAR_{inv}. The tetrazine bearing resin was incubated with the diene **6** dissolved in H₂O/tBuOH. The peptide product was cleaved off from the resin by TFA. (b) Absorption scan of resin bound tetrazine **1a** during the DAR_{inv} to monitor the progress of the reaction. (c) Crude RP-HPLC chromatogram and MALDI-ToF MS of the DAR_{inv} product **8c**.

maleimide modified RGD (**6**) resulted in the stable product **8c**. The reaction completed rather slowly after 7 d (Fig. 6). This reactivity towards tetrazines should be considered when using maleimide modified tetrazines or a combination of DAR_{inv} with a thiol-maleimide addition.^{41–43} Moreover, this finding gives rise to the potential use of maleimides as easily accessible dienophiles in the DAR_{inv}. However, alternative tetrazines should be tested to increase the reaction rate. The outcome of all the studied reactions between the different tetrazines and dienophiles is summarized in Table 2.

The DAR_{inv} on a solid support displays the following advantages. Tetrazines are often hardly soluble in aqueous solution. Therefore, the reaction has to be carried out in organic solvents or the synthesis strategy has to be changed by the introduction of solubility increasing molecules such as short PEG units.¹⁸ Performing the DAR_{inv} with a tetrazine containing

peptide bound to a solid phase overcomes this problem. Besides the advantages of water as a solvent for environmental and biochemical issues it is also favorable for kinetic reasons.⁴⁴ It is known that water can increase the proximity of hydrophobic molecules like the used diene and dienophiles.³¹ Hence the reaction can be accelerated.⁴⁵ Additionally, the water-swelling TentaGel resin facilitates the synthesis of rather long or branched peptides compared to the synthesis carried out on non-PEG-based resins. Furthermore, cleavage and purification steps are reduced since the crude resin bound peptides can be directly used for the ligation. This is especially beneficial for peptides that consist of rather cost intensive amino acids such as DOPA. DAR_{inv}-reagents as the dienophile carrying RGD peptide are easily recovered by lyophilization and can be thus used in excess without loss of yield owing to purification steps.

DAR_{inv}-linkers as spacers to improve cell adhesion

RGD-mediated cell adhesion is strongly dependent on the distance between the integrin ligand and the corresponding surface. Various spacers as polyglycine, polyproline, amino-hexanoic acid or PEG have been used so far.^{46,47} Using bulky dienes and dienophiles circumvents the introduction of additional spacer moieties. Hence we applied the DAR_{inv} with rather large dienes and dienophiles to efficiently introduce a distinct spacer. To test the influence of the varying lengths of tetrazines **2a** and **2b**, cell adhesion studies with the peptide conjugates **8a** and **8b** were performed. Furthermore, peptide **8d**, synthesized by maleimide-thiol Michael addition, was tested to compare the tested DAR_{inv}-linker with a shorter spacer (Fig. 7a). Negative controls, containing RAD were synthesized to proof the RGD-mediated cell adhesion (see the ESI†). The increased cell size is an indicator of successful cell attachment and is dependent on the RGD-surface distance.⁴⁷ Therefore, cell spreading was determined by measuring the average cell area on coated Ti. Interestingly, conjugate **8b** showed the significantly highest average cell size compared to all the tested coatings, although its spacer is shorter than that of peptide **8a** (Fig. 7b). By MD-simulation the maximum length (from C α of Lys in RGD to C α of Lys in TBP and respectively to C α of Cys in TBP **2**) as well as the most likely present spacer distance were calculated. These studies revealed that the longest spacer is in fact present in **8a** with 34 Å. However, this linker is more prone to collapse and interacts intramolecularly, thus it only reaches a distance of around 12 Å for the most stable conformations (Table 3, see also the ESI†). Peptide **8b** consists of a spacer with a maximum length of 30 Å. The mostly occurring distance is 20 Å, thus cell spreading is more advanced on **8b** than on **8a**. The linker of **8d** was, as anticipated, the shortest with a maximum length of 21 Å and the most stable conformations showed a distance of 16 Å. Although the most likely distance of **8d** is higher than that of **8a**, cell spreading is more pronounced on **8a**. This can be explained by the higher average length of 19 Å for **8a** compared to 16 Å for **8d** (Fig. 7d and Table 3).

Table 2 Summary of reactivity between different diene/dienophile combinations

Tetrazine	Dienophile			
	3	4	5	6
1a	Stable product 8a	Unstable product ^a	Unstable product ^a	Stable product 8c
1b	Stable product 8b	Low conversion ^a	Low conversion ^a	Low conversion ^a

^a Unstable products and low conversion of reactions are discussed in more detail in the ESI.



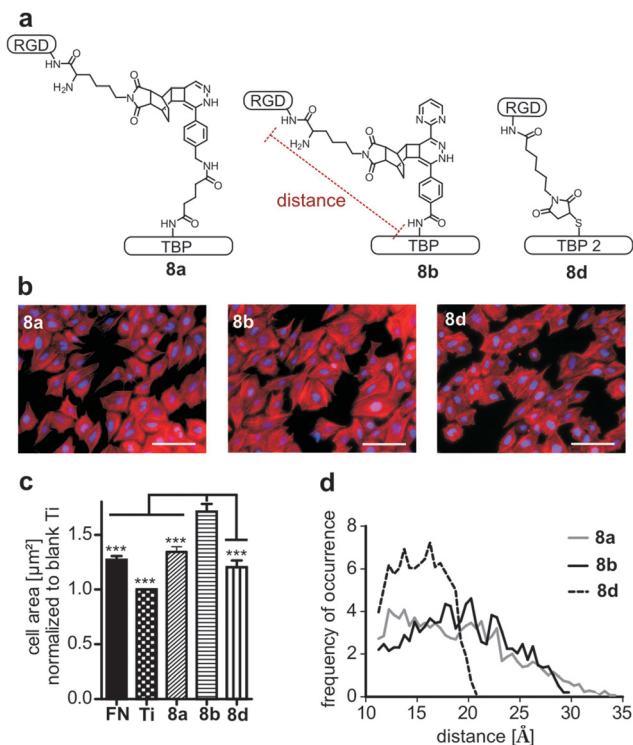


Fig. 7 Cell response to different peptide DAR_{inv} linkers (a) DAR_{inv} products with different spacings of the cell binding peptide c[RGDfK]. Peptide **8d** was synthesized by maleimide-thiol Michael addition. (b) Fluorescence microscopy of SaOS-2 cells adhered on peptides for 6 h, scale bar: 100 μm , the actin cytoskeleton is stained in red and nuclei in blue. (c) Average cell area after 6 h cell adhesion. Fibronectin (FN) coated Ti was used as positive and Ti (non-coated titanium) as negative control. Data are presented as mean \pm SEM of $n \geq 3$ (significant differences to **8b** were determined by one-way ANOVA ($***p < 0.001$)). (d) Normalized frequencies of occurrence of certain linker distances, determined by molecular dynamics (MD)-simulation.

Table 3 Summary of linker distances in **8a,b** and **d** calculated by MD-simulation

	Maximum distance [\AA]	Most frequent distance [\AA]	Average distance [\AA]
8a	34	12	19
8b	30	20	19
8d	21	16	15

Previous studies already have suggested that a longer, more rigid and extended spacer (e.g. polyproline) favors RGD-mediated cell adhesion, however until now this has not been assessed by MD-simulation.⁴⁶ In addition, the herein described MD-simulations support the assumption that too long spacers impair cell adhesion because of their pronounced flexibility.^{46,48} Nevertheless, a long and flexible linker should be favored over a too short spacer as their average length may be higher. These results highlight that estimating the distance of a RGD peptide to its surface only by the extended conformation of a linker molecule is not sufficient. It is rather

suggested that the three-dimensional length is evaluated. We suggest that a suitable linker consists of two short flexible ends, connected by a long rigid structure (see the ESI†). The flexible parts in the spacer may be important to freely orientate the RGD-peptide on the surface towards the cell membrane.

Conclusion

The Diels–Alder reaction with inverse electron demand (DAR_{inv}) could be successfully employed to ligate peptides on a solid support. This method describes an efficient strategy to complex peptide conjugates without using catalysts or additive reagents and it is easily monitored. Low solubility as well as yield loss and potential fragmentation through cleaving tetrazine modified peptides from the resin are thereby circumvented. Furthermore, maleimide was found to be an easily accessible dienophile alternative for DAR_{inv} ligations on resin. Large DAR_{inv} -reagents could be applied to generate a spacer that might be useful for several biochemical applications. MD-simulations gave insights into the theoretical conformations of different DAR_{inv} -linkers, thereby providing crucial information for the choice of a distinct diene–dienophile pair. Thereby, tetrazine **2b** in the DAR_{inv} product **8b** resulted in the most enhanced cell adhesion because of its less flexible structure. With the here presented DAR_{inv} on resin the toolbox of orthogonal reactions for SPPS is broadened.

Acknowledgements

We are thankful for the financial support from the DFG (TRR67, A4), the European Union and the Free State of Saxony as well as of the Graduate School BuildMoNa. For technical support in cell culture Kristin Löbner is acknowledged. The authors would like to thank Regina Reppich-Sacher for excellent assistance in mass spectrometry analysis. Ronny Müller and Christina Dammann are acknowledged for help in peptide synthesis.

References

- 1 A. Isidro-Llobet, M. Alvarez and F. Albericio, *Chem. Rev.*, 2009, **109**, 2455–2504.
- 2 D. Böhme and A. G. Beck-Sickinger, *J. Pept. Sci.*, 2015, **21**, 186–200.
- 3 V. M. Ahrens, R. Frank, S. Boehnke, C. L. Schutz, G. Hampel, D. S. Iffland, N. H. Bings, E. Hey-Hawkins and A. G. Beck-Sickinger, *ChemMedChem*, 2015, **10**, 164–172.
- 4 H. C. Kolb, M. G. Finn and K. B. Sharpless, *Angew. Chem., Int. Ed.*, 2001, **40**, 2004–2021.
- 5 D. Schumacher and C. P. R. Hackenberger, *Curr. Opin. Chem. Biol.*, 2014, **22**, 62–69.
- 6 E. M. Sletten and C. R. Bertozzi, *Angew. Chem., Int. Ed.*, 2009, **48**, 6974–6998.



- 7 S. Hofmann, S. Maschauer, T. Kuwert, A. G. Beck-Sickinger and O. Prante, *Mol. Pharmaceutics*, 2015, **12**, 1121–1130.
- 8 W. Tang and M. L. Becker, *Chem. Soc. Rev.*, 2014, **43**, 7013–7039.
- 9 C. W. Tornøe, C. Christensen and M. Meldal, *J. Org. Chem.*, 2002, **67**, 3057–3064.
- 10 H. O. Ham, S. H. Park, J. W. Kurutz, I. G. Szleifer and P. B. Messersmith, *J. Am. Chem. Soc.*, 2013, **135**, 13015–13022.
- 11 I. E. Decostaire, D. Lelievre, V. Aucagne and A. F. Delmas, *Org. Biomol. Chem.*, 2014, **12**, 5536–5543.
- 12 J. Bertran-Vicente, M. Schumann, P. Schmieder, E. Krause and C. P. Hackenberger, *Org. Biomol. Chem.*, 2015, **13**, 6839–6843.
- 13 R. Pipkorn, W. Waldeck, B. Diding, M. Koch, G. Mueller, M. Wiessler and K. Braun, *J. Pept. Sci.*, 2009, **15**, 235–241.
- 14 R. Selvaraj, B. Giglio, S. Liu, H. Wang, M. Wang, H. Yuan, S. R. Chintala, L. P. Yap, P. S. Conti, J. M. Fox and Z. Li, *Bioconjugate Chem.*, 2015, **26**, 435–442.
- 15 R. Hassert, M. Pagel, Z. Ming, T. Haupl, B. Abel, K. Braun, M. Wiessler and A. G. Beck-Sickinger, *Bioconjugate Chem.*, 2012, **23**, 2129–2137.
- 16 K. Braun, M. Wiessler, R. Pipkorn, V. Ehemann, T. Bauerle, H. Fleischhacker, G. Muller, P. Lorenz and W. Waldeck, *Int. J. Med. Sci.*, 2010, **7**, 326–339.
- 17 B. M. Zeglis, F. Emmetiere, N. Pillarsetty, R. Weissleder, J. S. Lewis and T. Reiner, *ChemistryOpen*, 2014, **3**, 48–53.
- 18 A. Niederwieser, A. K. Spate, L. D. Nguyen, C. Jungst, W. Reutter and V. Wittmann, *Angew. Chem., Int. Ed.*, 2013, **52**, 4265–4268.
- 19 M. Kantlehner, P. Schaffner, D. Finsinger, J. Meyer, A. Jonczyk, B. Diefenbach, B. Nies, G. Holzemann, S. L. Goodman and H. Kessler, *ChemBioChem*, 2000, **1**, 107–114.
- 20 D. M. Lewallen, D. Siler and S. S. Iyer, *ChemBioChem*, 2009, **10**, 1486–1489.
- 21 A. S. Ham, A. L. Klivanov and M. B. Lawrence, *Langmuir*, 2009, **25**, 10038–10044.
- 22 G. Vidal, T. Blanchi, A. J. Mieszawska, R. Calabrese, C. Rossi, P. Vigneron, J. L. Duval, D. L. Kaplan and C. Egles, *Acta Biomater.*, 2013, **9**, 4935–4943.
- 23 J. Auernheimer, D. Zukowski, C. Dahmen, M. Kantlehner, A. Enderle, S. L. Goodman and H. Kessler, *ChemBioChem*, 2005, **6**, 2034–2040.
- 24 J. H. Beer, K. T. Springer and B. S. Coller, *Blood*, 1992, **79**, 117–128.
- 25 B. Hess, C. Kutzner, D. van der Spoel and E. Lindahl, *J. Chem. Theory Comput.*, 2008, **4**, 435–447.
- 26 J. M. Wang, W. Wang, P. A. Kollman and D. A. Case, *J. Mol. Graphics Modell.*, 2006, **25**, 247–260.
- 27 A. Sousa da Silva and W. Vranken, *BMC Res. Notes*, 2012, **5**, 1–8.
- 28 H. Lee, N. F. Scherer and P. B. Messersmith, *Proc. Natl. Acad. Sci. U. S. A.*, 2006, **103**, 12999–13003.
- 29 M. Pagel, R. Hassert, T. John, K. Braun, M. Wiessler, B. Abel and A. G. Beck-Sickinger, *Angew. Chem., Int. Ed.*, 2016, **55**, 4826–4830.
- 30 R. Hoogenboom, B. C. Moore and U. S. Schubert, *J. Org. Chem.*, 2006, **71**, 4903–4909.
- 31 S. Horner, C. Uth, O. Avrutina, H. Frauendorf, M. Wiessler and H. Kolmar, *Chem. Commun.*, 2015, **51**, 11130–11133.
- 32 K. Braun, M. Wiessler, V. Ehemann, R. Pipkorn, H. Spring, J. Debus, B. Diding, M. Koch, G. Muller and W. Waldeck, *Drug Des., Dev. Ther.*, 2009, **2**, 289–301.
- 33 M. Wiessler, W. Waldeck, C. Kliem, R. Pipkorn and K. Braun, *Int. J. Med. Sci.*, 2010, **7**, 19–28.
- 34 M. R. Karver, R. Weissleder and S. A. Hilderbrand, *Bioconjugate Chem.*, 2011, **22**, 2263–2270.
- 35 H. S. Beckmann, A. Niederwieser, M. Wiessler and V. Wittmann, *Chemistry*, 2012, **18**, 6548–6554.
- 36 A. K. Späte, V. F. Schart, S. Schöllkopf, A. Niederwieser and V. Wittmann, *Chemistry*, 2014, **20**, 16502–16508.
- 37 Y. J. Lee, Y. Kurra, Y. Yang, J. Torres-Kolbus, A. Deiters and W. R. Liu, *Chem. Commun.*, 2014, **50**, 13085–13088.
- 38 R. M. Versteegen, R. Rossin, W. ten Hoeve, H. M. Janssen and M. S. Robillard, *Angew. Chem., Int. Ed.*, 2013, **52**, 14112–14116.
- 39 J. Li, S. Jia and P. R. Chen, *Nat. Chem. Biol.*, 2014, **10**, 1003–1005.
- 40 A. D. de Araujo, J. M. Palomo, J. Cramer, O. Seitz, K. Alexandrov and H. Waldmann, *Chemistry*, 2006, **12**, 6095–6109.
- 41 Z. Wu, S. Liu, M. Hassink, I. Nair, R. Park, L. Li, I. Todorov, J. M. Fox, Z. Li, J. E. Shively, P. S. Conti and F. Kandeel, *J. Nucl. Med.*, 2013, **54**, 244–251.
- 42 S. Liu, M. Hassink, R. Selvaraj, L. P. Yap, R. Park, H. Wang, X. Chen, J. M. Fox, Z. Li and P. S. Conti, *Mol. Imaging*, 2013, **12**, 121–128.
- 43 J. D. Thomas, H. Cui, P. J. North, T. Hofer, C. Rader and T. R. Burke Jr., *Bioconjugate Chem.*, 2012, **23**, 2007–2013.
- 44 G. K. van Der Wel, J. W. Wijnen and J. B. Engberts, *J. Org. Chem.*, 1996, **61**, 9001–9005.
- 45 G. Graziano, *J. Phys. Org. Chem.*, 2004, **17**, 100–101.
- 46 D. Pallarola, A. Bochen, H. Boehm, F. Rechenmacher, T. R. Sobahi, J. P. Spatz and H. Kessler, *Adv. Funct. Mater.*, 2014, **24**, 943–956.
- 47 J. W. Lee, Y. J. Park, S. J. Lee, S. K. Lee and K. Y. Lee, *Biomaterials*, 2010, **31**, 5545–5551.
- 48 U. Hersel, C. Dahmen and H. Kessler, *Biomaterials*, 2003, **24**, 4385–4415.

