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Photo-polymerizable, low shrinking modular construction kit with high efficiency based on vinylcyclopropanes†

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The successful development of a universal hydrogen bond (H-bond) concept, specific for bi-functional vinylcyclopropane (VCP) ester–amide derivatives is reported here. Thereby, uniformity within the intermolecular hydrogen bond strength as investigated by variable temperature NMR and FT-IR measurements provides an excellent control of high reactivity, nearly regardless of the chosen spacer-unit, offering the system a possible application as a modular construction kit for cross-linked networks with varied properties. Two entirely diversified VCP ester–amide systems were prepared. The performance benchmark of a flexible, high molar mass macro-monomer VCPPPG₂₀₀₀, implementing a polypropyleneglycol Genamine D01/2000 macro-spacer, and two rigid, low molar mass monomers VCP-*m*-phenyl and VCP-*p*-phenyl outlined significantly the strength of the system. Extremely low volume shrinkages between 1.4–4.5% and a wide range of E-moduli could be achieved. An easy control of adjusting final characteristics by varying the co-monomer content in co-networks was shown, especially without obtaining any significant disadvantage in the curing behavior. The VCP ester–amides offer the possibility to replace some currently used dimethacrylate resins within several high-end applications, especially where low volume shrinkage and high polymerization efficiency are required.

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Introduction

Bi-functional, cross-linkable resins featuring a variety of precise specifications are of great interest within a wide spectrum of applications, such as electronics,¹ coatings,^{2,3} lithography,^{4,5} dental applications,⁶ 3D microstructures⁷ and post-polymerization modifications.⁸ However, since the final properties of resins have to match precisely with the complex specifications, there is a demand for monomer-systems providing universality and high precision.^{1,9} A suitable method could be to apply low shrinking modular construction kits, merging rather specific techniques within one certain system.

Nevertheless, monomer-systems based on methacrylates pose serious constraints, due to a high discrepancy in photocuring behaviour, using cross-linkable resins with varied spacer-elements.^{10,11} Likewise the observed polymerization shrinkage during curing of methacrylate systems is too high, which can result in high internal compressive stress, leading

to microcracks, incomplete fillings, poor adhesion and a short life span of cured resin.¹² Based on these two issues methacrylate systems are not appropriate for low shrinking modular construction kits. An alternative concept, partly developed from the knowledge of low-molecular-weight gelators and supramolecular structures, establishes a further promise on weak forces, such as hydrogen bonds (H-bonds), presuming a partial self-assembly of the molecules. In particular, if a partial self-assembly is taking place selectively at a polymerizable group, a fast and selective curing, as well as targeted control can be assumed, likewise it is required for a modular kit.

Thereby, vinylcyclopropane (VCP) ester–amide derivatives can play a decisive role in the future. Recently we have shown in a special case, that intermolecular amide H-bonds could induce a very effective partial preorganization of bi-functional VCPs molecules, which strongly increased the polymerization behavior.¹³ Further, the radical ring-opening polymerization (RROP) of VCP units can significantly reduce the volume shrinkage on polymerization. Therefore, the shrinkage is lower than those recorded for *e.g.* for methacrylate resins, making VCPs attractive as low-shrinking monomers.^{14,15}

However, so far the advances within VCP resins have been mainly focused on particular characteristics, *e.g.* reducing the volume shrinkage and improving the reactivity, the ring-

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opening efficiency and mechanical properties, especially among researchers in the field of dental fillings.^{16–18}

Herein, we report the design of the first highly efficient modular construction kit based on bi-functional VCP ester-amides, extending the general concept of hydrogen-bonding to a universal tool for an extensive application of VCPs as low shrinking resin. The H-bond strength of two diverse VCP ester-amide derivatives was studied in detail, first for a flexible, high molar mass macro-monomer (VCPPPG₂₀₀₀, $M_w \sim 2230 \text{ g mol}^{-1}$), and secondly for a rigid, low molar mass monomer (VCP-*m*-phenyl, $M_w = 468.5 \text{ g mol}^{-1}$) by temperature dependent NMR- and FT-IR-measurements. Further, a detailed overview of mechanical strength by three point bending experiments, and thermal properties of the cured networks and co-networks by differential scanning calorimetry (DSC) and thermogravimetric (TGA) analysis is provided. In addition to the VCP-*m*-phenyl its constitution isomer of a *para*-phenyl substituted VCP ester-amide resulted in a partial-crystalline system. Thus, the cured and un-cured resins have been studied in detail by nuclear magnetic resonance (NMR) measurements in liquid and solid-state, by temperature dependent polarized microscopy as well as by temperature dependent X-ray diffraction (XRD).

Experimental section

Materials

m-Xylylenediamine (99%), *p*-xylylenediamine (99%), *trans*-1,4-dibromo-2-butene (99%), diethyl malonate (99%), potassium hydroxide (KOH, >85%), *N,N'*-dicyclohexylcarbodiimide (DCC, 99%), ethyl 4-(dimethylamino)benzoate (EDMAB, 99%), triethylamine (99%), camphorquinone (CQ, 97%), 1,3,5-trimethylbenzene (mesitylene, >99.8%) and dichloromethane (>99.5%) were supplied by Sigma Aldrich and used without further purification. 1-Hydroxybenzotriazole hydrate (HOBt, >97%, 12% water, Sigma Aldrich) was dried in vacuum prior use. Sodium (>99%, Sigma Aldrich) was purified by melting in dry xylene prior use. Genamine D01/2000 (99%) was provided by Clariant and used as received (determined by (MALDI-TOF-MS): $M_n = 1980 \text{ g mol}^{-1}$, $M_w = 2027 \text{ g mol}^{-1}$, $D = 1.02$). Urethane-dimethacrylate (UDMA), dodecanedioldimethacrylate (1,12-DMA) and a 6:4 mixture of bisphenol-A-glycidyl methacrylate (BisGMA) and triethylene-glycol-dimethacrylate (TEGDMA) have been provided by the Kettenbach GmbH (Eschenburg, Germany) and were used as received.

Analytical methods

¹H- (300 MHz) and ¹³C-NMR spectra (75 MHz) were recorded on a Bruker Ultrashield-300 spectrometer at room temperature in CDCl₃. The spectra were calibrated on the solvent signal ($\delta(^1\text{H}) = 7.26 \text{ ppm}$; $\delta(^{13}\text{C}) = 77.16 \text{ ppm}$). Variable ¹H temperature experiments at a concentration of 5 mM in CDCl₃ have been performed in a range from 20 °C to 60 °C. High-resolution solid state ¹³C-CP/MAS (cross polarization/magic angle spinning) NMR Spectra were measured by a Bruker spectro-

meter operating at 100 MHz, using a pulse experiment with a 30 s pulse delay for 4096 scans. High resolution mass spectra, coupled online to a HPLC system, were recorded on a Q-Exactive Orbitrap (Thermo Scientific). MALDI-TOF MS analysis were performed on a Bruker Reflex III instrument equipped with a N₂ laser ($\lambda = 337 \text{ nm}$) in linear and reflectron mode. The acceleration voltage was 20 kV. Dithranol (Sigma Aldrich, >98%) was used as matrix material and sodium trifluoroacetate (Sigma Aldrich, 98%) as ionization salt. The samples were prepared with the dried droplet method from THF solution by mixing matrix, polymer and salt in a ratio of 20:5:1 and applying approximately 1 μL to the target spot. Temperature dependent FT-IR spectra were recorded by a Nicolet Nexus 470 spectrometer with a heated attenuated total reflectance (ATR) unit. Chromatographic purifications were performed using a GRACE Reveleris X2 apparatus. Technical grade solvents, which have been distilled prior use, have been utilized. A dynamic-mechanical analyzer (DMTA IV, Rheometric Scientific) has been used to determine the E-moduli by three-point bending investigations, respectively at 25 °C. A Mettler thermal DSC analyzer (821c) was used for the thermal characterization of the monomer resins and cross-linked polymers. The DSC scans were recorded in nitrogen atmosphere and heating rates of 10 K min⁻¹. Thermogravimetric analysis were done with a TG 209 F1 (Netzsch) by heating the samples up to 800 °C, respectively under nitrogen atmosphere as well as under synthetic air. 85 μm corundum crucibles, heating rates of 10 K min⁻¹ and about 10–15 mg of each sample were applied. An AccuPyc 1330 gas pycnometer (Micromeritics) operating with helium gas was used to determine the density of monomers and cross-linked polymers. Temperature dependent XRD patterns were carried out on an X-ray powder diffractometer of the type PANalytical XPert Pro, applying a reaction chamber to measure from 25 to 90 °C in a 2 Theta range from 7 to 35, respectively.

Synthesis of polymer networks

To perform the photo initiation a blue light LED reactor with a light power of $\sim 2.0 \text{ mW cm}^{-2}$ at 465 nm was used as radiation source. 1 mol% of the photo-initiator mixture CQ:EDMAB in a molar ratio of 1:2 was used. The initiator was dissolved in little DCM and added to the resin. The DCM was evaporated in vacuum prior to curing. For DMTA measurements specimens were produced by filling a stainless steel profile with a dimension of $25 \times 3 \times 1 \text{ mm}$ with the monomer/initiator mixture. The profile was enclosed and fixed both on top and on the bottom with a 50 μm thick hostophan sheet and irradiated for 2 h. The cured specimens were pressed out carefully and stored for 24 h at room temperature prior to further testing. For density measurements the monomer initiator mixture was inserted into a 2 mL standard test tube, flushing the tubes with an argon stream for 30 s and irradiated for 2 h. Afterwards the samples were taken out of the glass tubes and stored for 24 h at 100 °C prior measuring the densities. The partial-crystalline VCP-*p*-phenyl was polymerized in 50 weight% solution of chloroform/toluene (1:1). The reaction mixture was



degassed by three freeze–pump–thaw cycles and polymerized within the LED reactor under stirring over 12 h. The polymerization was stopped by putting the test tube into liquid nitrogen; the samples were taken out of the glass tube, transferred into centrifuge tubes and extracted three times by over 25-fold excess of acetone. Afterwards, the resulting polymer was dried in vacuum at 100 °C for 24 h.

Determination of residual monomer amount

The quantification of the residual amount of uncured monomer was performed by $^1\text{H-NMR}$ measurements. Therefore the weighed and cured samples have been immediately immersed after radiation in 2.0 mL of CDCl_3 (as extraction fluid) in 5 mL bottles for 24 h. The residual monomer amounts have been calculated by comparing the integration areas of the respective $^1\text{H-NMR}$ proton signals to the proton signals of mesitylene, which has been added to the CDCl_3 as internal standard in $2.8444 \text{ mg mL}^{-1}$ amount. To achieve a high signal to noise resolution for each $^1\text{H-NMR}$ experiment a number of scans of 2048 has been chosen.

Synthesis of diethyl 2-vinylcyclopropane-1,1-dicarboxylate and 1-(ethoxycarbonyl)-2-vinylcyclopropanecarboxylic acid

Diethyl 2-vinylcyclopropane-1,1-dicarboxylate and the hydrolyzed diethyl 2-vinylcyclopropane-1,1-dicarboxylate were obtained as per published procedure.^{14,19}

General procedure to obtain bi-functional VCP ester–amide derivatives

In a round-bottom flask purged with argon 0.5 eq. (0.051 mol) of the diamine compound were dissolved/diluted in 200 mL of anhydrous DCM. The solution was transferred to a funnel, which was connected to 1000 mL three-necked reaction flask, charged with 200 mL of anhydrous DCM, 1.1 eq. (0.112 mol) of 1-(ethoxycarbonyl)-2-vinylcyclopropane-carboxylic acid, 1.1 eq. (0.112 mol) of DCC and 1.15 eq. (0.117 mol) of HOBt. The three-necked reaction flask was stirred for 15 min at room temperature, cooled to 0 °C and then 2.1 eq. (0.214 mol) of triethylamine were added. Afterwards the diamine solution with a drip rate of about 40 mL h^{-1} was added slowly. The reaction solution was stirred for 12 h at room temperature. Subsequently the reaction mixture was filtered, the pale yellow solution was washed two times with water (100 mL), once with 1 molar HCl solution (100 mL) and three times with neutral water (50 mL). After drying over magnesium sulfate, the solution was deposited on silica gel and purified by flash chromatography (general eluents: hexane and ethyl acetate). To the purified organic phase 1500–2500 ppm of butylated hydroxytoluene were added. The organic phase was evaporated in vacuum and the bi-functional VCP–amide derivative dried *in vacuo*. The average yield after purification was 87%. A detailed monomer characterization of the synthesized VCP₂₀₀₀, VCP-*m*-phenyl and VCP-*p*-phenyl monomers is provided in the ESI in Fig. S1–S8.†

Results and discussion

The availability of targeted amendments to control selective characteristics by varying the spacer unit, without major transformation *e.g.* in reactivity is crucial for designing a modular construction kit for photo-polymerization purposes. Due to the existence of selective intermolecular H-bonds exactly at the VCP-unit, VCP ester–amide derivatives provide a clear advantage as the curing behaviour is actually independent of the chosen spacer unit. Thereby, Fig. 1A illustrates the universal concept of partial self-assembly for any VCP ester–amide derivative. Moreover, the proposed RROP mechanism of VCP ester–amides (Fig. 1B) clarifies the general efficiency of the intermolecular H-bonds, which are preferred because of N–H...O donor–acceptor interactions and due to chelating effect.

To provide a reasonable illustration, two entirely diversified VCP ester–amide systems were prepared. Thus, first a flexible, high molar mass macro-monomer VCP₂₀₀₀ and secondly two rigid, low molar mass monomers VCP-*m*-phenyl and VCP-*p*-phenyl have been synthesized and compared subsequently.

To study the H-bond relationship, both variable-temperature NMR- (Fig. 2) as well as FT-IR-measurements (see Fig. S9 and S10 in the ESI†) have been carried out, for VCP₂₀₀₀ and VCP-*m*-phenyl. Thereby the evaluation of the temperature dependency of the amide proton signal revealed the presence of H-bond. The corresponding signal was shifted to higher magnetic field, as with increased temperature a decreased deshielding effect is anticipated.^{20,21} The partial regression line of the chemical shift confirmed independently for both VCP ester–amides ($R^2 = 0.999$), that within the investigated temperature range of 20–60 °C the H-bonds were weakened, but not disrupted. Further, the corresponding slopes of the linear regressions were more or less identical, $-4.32 \times 10^{-3} \text{ ppm K}^{-1}$ for VCP₂₀₀₀ and $-4.85 \times 10^{-3} \text{ ppm K}^{-1}$ for VCP-*m*-phenyl,



Fig. 1 (A) Schematic aspect of the H-bond mediated partial self-assembly of VCP ester–amide units. The alignment is primary taking place at the functional group. (B) Proposed RROP mechanism of amide-based VCPs. The partial self-assembly and the enhanced orbital arrangements have to be taken into consideration.



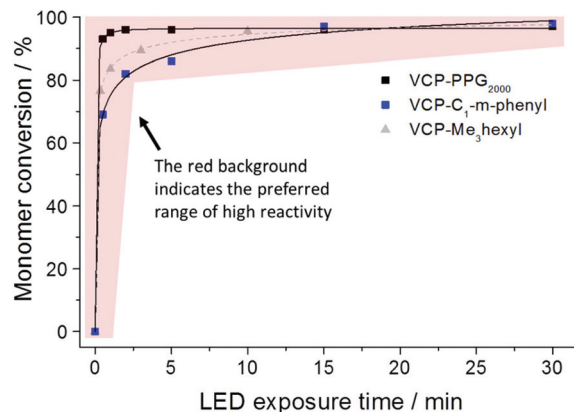


Fig. 3 Determined monomer conversion for the photo-polymerization of VCPPPP_{G2000}, VCP-*m*-phenyl and VCPMe₃hexyl (1 mol% initiator CQ : EDMAB in a molar ratio of 1 : 2). The curing was performed under a controlled atmosphere of nitrogen using a commercial LED source (2.013 mW cm⁻² for 465 nm).

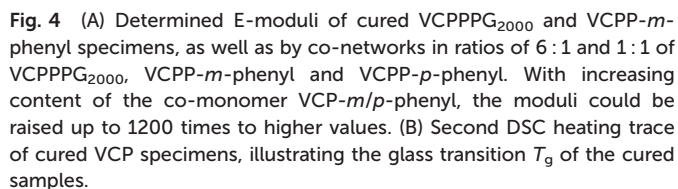
ester-amides an overall conversion of >96% could be observed. In comparison, VCP-resins without H-bonded interactions cannot reach high overall conversion even after prolonged curing times.¹⁴ Thus VCP ester-amides like VCPPPG₂₀₀₀, VCP-*m*-phenyl as well as the previous studied VCPMe₃hexyl have to be seen as pioneers defining a new, universal class of highly efficient and low shrinking resins. However, it is worth noting, that VCPPPG₂₀₀₀ reached a conversion of 92% after 30 s of exposure. In this particular case, the slightly enhanced conversion of VCPPPG₂₀₀₀ was explained by its low T_g of -55 °C (Table 1). Therefore, a high mobility and continuous diffusion to further radical centers is allowed.

Hence, compared to methacrylate systems, VCP ester-amides show fast curing kinetics irrespective of the spacer unit, but is superior defining the final network characteristics as discussed in Fig. 4.

Thereby, we investigated the mechanical properties of cured VCP ester-amid specimens by three point bending experiments, for VCP_{PPG2000} and VCP-*m*-phenyl, as well as for the co-networks of VCP_{PPG2000}, VCP-*m*-phenyl and VCP-*p*-phenyl in different ratios (Fig. 4A). The mechanical pro-

Entry	Polymer	T_g^a °C	$T_{5\%}^b$ (N ₂)/°C	$T_{5\%}^c$ (air)/°C	Water uptake ^d /%	E_{mod}^e MPa
1	VCPPPG ₂₀₀₀	−55	340	239	1.3	0.15
2	VCP- <i>m</i> -phenyl	147	329	327	2.9	232.3
3	VCP- <i>p</i> -phenyl	189	339	321	2.3	—
4	UDMA	119	316	291	2.8	224.1

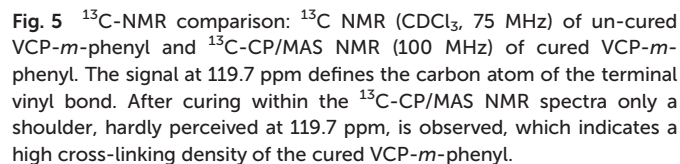
^a Determined by DSC. ^b Determined by TGA (50 mL min⁻¹ N₂ as purge gas). ^c Determined by TGA (50 mL min⁻¹ synthetic air as purge gas). ^d Determined by TGA as mass loss between 30–200 °C, after storage for 48 h at 50 °C in water, respectively. ^e After curing specimens of the dimension 25 × 3 × 1 mm³ by photo-polymerization and additional annealing for 12 h at 100 °C, respectively.



Further, we could easily control the E-moduli by varying the co-monomer content in the co-networks. With increasing content of the co-monomer VCP-*m/p*-phenyl, the moduli could be raised continuously from 0.19 ± 0.001 MPa for VCPPPP₂₀₀₀, up to a value of 232.3 ± 0.5 MPa for the cured VCP-*m*-phenyl. Thus, the moduli could be raised up to 1200 times to higher

Subsequently, the cured VCP ester-amide networks were analyzed by ^{13}C -CP/MAS solid-state spectroscopy. Due to the absence of the carbon atom of the terminal vinyl bond at 119.7 ppm an extraordinary high cross-linking density of the cured networks could be confirmed, respectively for VCP-*m*-phenyl (Fig. 5) and VCP-*p*-phenyl (S13 in the ESI†). Only a very small shoulder remained, hardly perceived, which indicated a negligible amount of remaining vinyl double bonds.

Further to depict the general potential of reducing the volume-shrinkage during polymerization by applying RROP of



VCP ester-amide resins, the corresponding state variables volume shrinkage, monomer- and polymer density for cured and uncured states were provided within Fig. 6. Very low volume changes between -4.5% to -1.4% have been determined for the VCP ester-amide resins, as well as for their co-networks in different ratios. It is remarkable, that the cured VCP-*m*-phenyl showed a reduction by 50% in volume shrinkage compared *e.g.* to UDMA, without compromising mechanical performance. VCP-PPG₂₀₀₀ showed an extremely low volume shrinkage (-1.4%), represented by its increased molecular weight as macro-monomer.^{9,27}

In this context we observed a significant difference between the cured isomers VCP-*m*-phenyl and VCP-*p*-phenyl. While VCP-*m*-phenyl showed a volume change of -4.5% , for VCP-*p*-phenyl a volume change of -1.9% was observed. In comparison to the amorphous VCP-*m*-phenyl, its isomer VCP-*p*-phenyl occurs as a heterogeneous system, combining a liquid resin with partial crystallinity. Thereby the DSC curve of VCP-*p*-phenyl showed a melting at $T_m = 67^\circ\text{C}$, whereas VCP-*m*-phenyl remained completely amorphous (Fig. 7A). In agreement with the DSC measurement, temperature variable polarizing microscopy showed at the same temperature range a transition to an isotropic melt, respectively for VCP-*p*-phenyl (Fig. 7B).

Thus an increased density of VCP-*p*-phenyl (1.139 g cm^{-3}) was observed in comparison to VCP-*m*-phenyl (1.105 g cm^{-3}). A complete crystallization of VCP-*p*-phenyl is hindered, due to the general isomeric structure of VCP ester-amides (see Scheme S1 in the ESI†). Furthermore, once the crystalline phase is melted the reverse crystallization occurs within a longer time lag over weeks, thus within the XRD diffractogram no reflexes returned after cooling the resin back to room temperature (see Fig. S14 within the ESI†).

Generally, the volume change is calculated by the densities of the cured and uncured resins. As in most cases the density of the cured resin is higher, negative volume changes are observed. However, volume expandable resins are known



Fig. 7 (A) DSC curves of first heating cycles of un-cured VCP-*m/p*-phenyl resins. First the transition from a molecular glass to a resin ($T_g - 15^\circ\text{C}$) is passed, respectively for both VCPs. In the further heating VCP-*m*-phenyl shows a complete amorphous behavior, whereas for VCP-*p*-phenyl a melting (onset at 67°C) is observed. At a temperature range of $\sim 99^\circ\text{C}$ an auto-polymerization of both resins is observed. (B) Polarized microscopy images of the partial-crystalline VCP-*p*-phenyl resin. Beneath the melting transition both amorphous as well as crystalline areas can be observed. When the melting range is passed the VCP-*p*-phenyl resin becomes fully isotropic.

within the literature as well.^{28,29} Yet, their applicability is restricted, as the volume expansion is mainly based to the transition of a denser crystalline monomer structure compared to a less compressed, amorphous polymer structure. In fact, this attitude is lost, if the crystalline phase is melted or diluted by any further phase, like it can be observed also for the co-networks of VCP-*p*-phenyl (Fig. 5). In this sense heterogeneous systems, like VCP-*p*-phenyl, have to be seen as an intermediate stage between both physical states, offering a further alternative to reduce the volume shrinkage. Especially the connection of an easy processable wax-like structure and the low volume shrinkage exhibit further potential for composite formulations, and will be investigated in a following work.

Conclusions

A uniformity of the VCP ester-amide H-bond strength was shown, providing a universal concept of applying VCP ester-amides as suitable low-shrinking resins. With an excellent control of high reactivity, nearly regardless of the spacer-unit chosen, VCP ester-amides offer a clear advantage in comparison to methacrylate based systems. Thus with an appropriate variation of spacer-unit, selective characteristics, like *e.g.* optical properties or mechanical strength could be adjusted in a simple way, without obtaining any significant disadvantage within the curing behaviour. Based on this concept, new VCP ester-amide resins were presented, showing very low volume



Fig. 6 Spider chart showing the dependency between the monomer- and polymer density to the volume shrinkage. Only the partial-crystalline VCP-*p*-phenyl shows a disproportion within the correlation, based due to the denser partial-crystalline monomer structure of VCP-*p*-phenyl (1.139 g cm^{-3}).



changes between -1.4 to -4.5% , a broad variety in mechanical strength between 0.19 to 232 MPa and a high overall thermal stability even under an atmosphere of air. Thereby the network properties could be continuously adjusted by varying the comonomer content in co-networks. Moreover, due to the inevitable isomeric mixtures of VCP ester-amides, a heterogeneous system could be achieved, combining liquid phase with partial crystallinity, offering an intermediate stage between both physical states. The presented concept of applying VCP ester-amides as low shrinking resins for a modular construction kit is facile and variable. Thus VCP ester-amides offer vast potential for commercial use; especially as the number of combinations possibilities respectively for specified spacer units are yet not exploited.

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Notes and references

- 1 Y. Fuchs, O. Soppera and K. Haupt, *Anal. Chim. Acta*, 2012, **717**, 7.
- 2 Y. Jian, Y. He, L. Zhao, A. Kowalczyk, W. Yang and J. Nie, *Adv. Polym. Technol.*, 2013, **32**, 21331.
- 3 E. Andrzejewska, *Prog. Polym. Sci.*, 2001, **26**, 605.
- 4 C. Decker, *Macromol. Rapid Commun.*, 2002, **23**, 1067.
- 5 C. Gorsche, K. Seidler, P. Knaack, P. Dorfinger, T. Koch, J. Stampfl, N. Moszner and R. Liska, *Polym. Chem.*, 2016, **7**, 257.
- 6 K. L. van Landuyt, J. Snauwaert, *et al.*, *Biomaterials*, 2007, **28**, 3757.
- 7 A. S. Quick, A. de los S. Pereira, M. Bruns, T. Bückmann, C. Rodriguez-Emmenegger, M. Wegener and C. Barner-Kowollik, *Adv. Funct. Mater.*, 2015, **25**, 3735.
- 8 D. H. Seuyep Ntougam, G. A. Luinstra and P. Theato, *J. Polym. Sci., Part A: Polym. Chem.*, 2014, **52**, 2841.
- 9 K. S. Anseth, C. N. Bowman and N. A. Peppas, *J. Polym. Sci., Part A: Polym. Chem.*, 1994, **32**, 139.
- 10 I. Sideridou, V. Tserki and G. Papanastasiou, *Biomaterials*, 2002, **23**, 1819.
- 11 S. H. Dickens, J. W. Stansbury, K. M. Choi and C. J. E. Floyd, *Macromolecules*, 2003, **36**, 6043.
- 12 J. W. Stansbury, M. Trujillo-Lemon, H. Lu, X. Ding, Y. Lin and J. Ge, *Dent. Mater.*, 2005, **21**, 56.
- 13 P. Pineda Contreras, C. Kuttner, A. Fery, U. Stahlschmidt, V. Jerome, R. Freitag and S. Agarwal, *Chem. Commun.*, 2015, **51**, 11899.
- 14 P. Pineda Contreras, P. Tyagi and S. Agarwal, *Polym. Chem.*, 2015, **6**, 2297.
- 15 N. Moszner and U. Salz, *Macromol. Mater. Eng.*, 2007, **292**, 245.
- 16 N. Moszner, F. Zeuner, T. Völkel and V. Rheinberger, *Macromol. Chem. Phys.*, 1999, **200**, 2173.
- 17 F. Sanda and T. Endo, *J. Polym. Sci., Part A: Polym. Chem.*, 2001, **39**, 265.
- 18 V. Alupeu and H. Ritter, *e-Polym.*, 2002, **2**, 724.
- 19 V. Alupeu and H. Ritter, *Macromol. Rapid Commun.*, 2001, **22**, 1349.
- 20 K. D. Kopple, M. Ohnishi and A. Go, *J. Am. Chem. Soc.*, 1969, **91**, 4264.
- 21 V. Berl, M. Schmutz, M. J. Krische, R. G. Khoury and J.-M. Lehn, *Chem. – Eur. J.*, 2002, **8**, 1227.
- 22 M. T. Lemon, M. S. Jones and J. W. Stansbury, *J. Biomed. Mater. Res., Part A*, 2007, **83**, 734.
- 23 J. Lange, S. Toll, J.-A. E. Månson and A. Hult, *Polymer*, 1997, **38**, 809.
- 24 J. Tu, S. J. Tucker, S. Christensen, A. R. Sayed, W. L. Jarrett and J. S. Wiggins, *Macromolecules*, 2015, **48**, 1748.
- 25 S. G. Pereira, T. G. Nunes and S. Kalachandra, *Biomaterials*, 2002, **23**, 3799.
- 26 J. Park, G. Lee, H. Ooshige, A. Nishikata and T. Tsuru, *Corros. Sci.*, 2003, **45**, 1881.
- 27 J. E. Klee and U. Lehmann, *J. Org. Chem.*, 2009, **5**, 72.
- 28 H. Chiba, K. Kitazume, S. Yamada and T. Endo, *J. Polym. Sci., Part A: Polym. Chem.*, 2015, **54**, 39.
- 29 N. Moszner and U. Salz, *Prog. Polym. Sci.*, 2001, **26**, 535.

