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Hook, Andrew L. and Scurr, David J. and Anderson, Daniel G. and Langer, Robert and Williams, Paul and Davies, Martyn C. and Alexander, Morgan R. (2013) High throughput discovery of thermo-responsive materials using water contact angle measurements and time-of-flight secondary ion mass spectrometry. *Surface and Interface Analysis*, 45 (1). pp. 181-184. ISSN 1096-9918

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# High throughput discovery of thermo-responsive materials using water contact angle measurements and time-of-flight secondary ion mass spectrometry

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**Switchable materials, that alter their chemical or physical properties in response to external stimuli, allow for temporal control of material-biological interactions, thus, are of interest for many biomaterial applications. Our interest is the discovery of new materials suitable to the specific requirements of certain biological systems. A high throughput methodology has been developed to screen a library of polymers for thermo-responsiveness, which has resulted in the identification of novel switchable materials. In order to elucidate the mechanism by which the materials switch, ToF-SIMS has been employed to analyse the top 2 nm of the polymer samples at different temperatures. The surface enrichment of certain molecular fragments was identified by ToF-SIMS analysis at different temperatures, suggesting an altered molecular conformation. In one example a switch between an extended and collapsed conformation was inferred.**

**Keywords:** thermo-responsive; stimuli; switchable; ToF-SIMS; water contact angle; polymer microarray

## Introduction

Controlled capture and release from surfaces of biomolecules and biomolecular assemblies, such as eukaryotic cells, has been the focus of numerous studies and has been achieved using thermo-responsive hydrogels such as poly(N-isopropyl acrylamide) (pNIPAM).<sup>[1-5]</sup> This polymer has been extensively used to temporally control cell attachment by exploiting its transition between a swollen and collapsed state by altering the temperature above and below the lowest critical solution temperature (LCST).<sup>[1]</sup> Alternatives to pNIPAM based thermoresponsive hydrogels have been explored such as polymers containing the ethylene glycol moiety,<sup>[6, 7]</sup> for example, using 2-(2-methoxyethoxy)ethyl methacrylate (MEO<sub>2</sub>MA) and oligo(ethylene glycol) methacrylates.<sup>[8-10]</sup> To enlarge the scope of biological and physical applications where switchable materials can be applied, a broadened library of thermoresponsive materials is of interest. Recently, polymer microarrays have become a key tool for the discovery of novel polymers.<sup>[11-13]</sup> High throughput surface characterisation (HTSC) has also been developed on this platform and has enabled the elucidation of structure-function relationships.<sup>[11, 14-17]</sup> Recently, a study used polymer microarrays to screen for temperature responsive

materials based upon the thermal release of attached eukaryotic cells.<sup>[18]</sup> We use a different approach to identify thermo responsive materials, carrying out a direct screen of water contact angle switching to identify thermo-responsive materials rather than implying switchability through cell detachment. High throughput water contact angle (WCA) measurements were utilised to identify polymers with thermo-responsive properties from a library of 279 unique materials in a polymer microarray format. The surface sensitivity and molecularly specificity of time-of-flight secondary ion mass spectrometry (ToF-SIMS) was exploited to investigate temperature dependant conformational changes at the surface of the ‘hit’ polymers.<sup>[19, 20]</sup>

## Experimental

### Polymer Polymerisation

Polymer microarrays were formed using a XYZ3200 pin printing workstation (Biodot) as described previously.<sup>[15]</sup> Slotted metal pins (946MP8B, Arrayit) with a tip diameter of 295  $\mu\text{m}$  were used to transfer approximately 4 nL of polymerisation solution onto poly(2-hydroxyethyl methacrylate) (pHEMA) dip-coated substrates<sup>[21]</sup> before slides were irradiated with a long wave UV source for 1 min, resulting in an average polymer spot size of 435  $\mu\text{m}$ . For formation of polymer coupons, 8  $\mu\text{l}$  of polymerisation solution was dispensed in triplicate onto a pHEMA coated substrate or onto a 1.5 x 1.5 cm silicon wafer for ToF-SIMS samples. Polymer coupons were polymerised in an argon atmosphere ( $\text{O}_2 < 1300$  ppm) by photopolymerisation with a long wave UV source for 10 min. Polymerisation solution was composed of 75% (v/v) monomer (Sigma), 24% (v/v) DMF and 1% (w/v) photoinitiator 2,2-dimethoxy-2-phenylacetophenone. Samples were subsequently dried at  $< 50$  mTorr for 7 days. The monomers are shown in Figure S11.

### Water contact angle measurements

Sessile WCA measurements were taken of each polymer as previously described.<sup>[16]</sup> The temperature of an aluminium stage was regulated using a FBC 735 Temperature Controller (Fisherbrand) and held the samples at a constant temperature for 30 mins before WCA measurements were taken.

### Time of flight secondary ion mass spectrometry

ToF-SIMS analysis was performed on an ION-TOF IV instrument (IONTOF GmbH). Measurements were taken at temperatures of 5  $^{\circ}\text{C}$  and 40  $^{\circ}\text{C}$ . A pulsed 25 kV  $\text{Bi}_3^+$  primary ion source was used at a target current of approximately 1 pA to raster 2 randomly selected  $100 \times 100 \mu\text{m}$  areas of the coupon to collect both positive and negative secondary ions. Charge compensation of the samples was accomplished with a pulsed electron floodgun. The mass of secondary ions was determined using a time-of-flight mass analyser. The typical mass resolution (at  $m/z$  41) was just over 6000.

## Results

The formation of a first generation array was achieved by printing 279 unique solutions for polymerisation onto a pHEMA coated glass slide with subsequent UV initiated curing.<sup>[21]</sup> The polymers were formed from 23 amphiphilic monomers (Figure S11). Automated pico litre sessile drop WCA measurements were made for all 279 materials, initially at 8  $^{\circ}\text{C}$  and then at 40  $^{\circ}\text{C}$ , as a screen to identify thermally responsive polymers. This temperature range was chosen due to its biological relevance and the ease at which these temperatures can be achieved in many laboratories. The resultant  $\Delta\text{WCA}$  ( $\text{WCA}^{40} - \text{WCA}^8$ ) for each polymer is shown in Figure 1A. The  $\Delta\text{WCA}$  was assumed to be 0 for polymers where the measured  $\Delta\text{WCA}$  was below the limit of detection (3 times the standard deviation of a measurement). From this initial screen, the top 11 ‘hit’ compositions producing either a positive or negative  $\Delta\text{WCA}$  were selected for a second generation array where the two monomers from each composition were varied systematically from 0 - 100% in increments of 10%. The second generation array contained a total of 121 polymers and three replicate arrays were produced on the

same slide. The resultant  $\Delta$ WCA when the temperature was increased from 8 °C to 40 °C is shown in Figure 1B.

The 16 polymer compositions that produced the largest absolute  $\Delta$ WCA were selected for scale up to 10 mm diameter polymer coupons. The monomer composition of selected 'hit' formulations was chosen such that monomer content varied by at least 15% to maximise the compositional variation. The WCA for each of these materials was measured from 8 °C to 40 °C in increments of 8 °C. The WCA is plotted as a function of temperature in Figure SI2 and for the 4 materials with the largest overall  $\Delta$ WCA in Figure 2D. A significant difference in the  $\Delta$ WCA between the measurement on the microarray samples and the polymer coupons was noted for 5 of the 16 compositions (Figure SI3). The different thermo-responsive properties of the polymer coupons could be a result of the decreased surface area:volume ratio, resulting in an altered surface energy. This could cause the material to no longer undergo a temperature induced change in WCA in the temperature range studied. In summary, the largest negative  $\Delta$ WCA of  $-18.5^\circ \pm 1.8^\circ$  was measured for the homopolymer of monomer L (Figure 2A) and the largest positive  $\Delta$ WCA of  $17.1^\circ \pm 4.0^\circ$  was measured for the copolymer V(70%)L(30%). These values are of a similar magnitude to the  $\Delta$ WCA of 12-23° reported for pNIPAM.<sup>[22-24]</sup> The inclusion of 2-(2-methoxyethoxy)ethyl methacrylate (J) (Figure 2C) with monomer L did not significantly alter the WCA of the material nor the absolute change in the WCA with temperature, but rather increased the temperature at which the WCA of the polymer decreased (Figure 2D).

The 4 polymer coupons with the largest measured  $\Delta$ WCA with a change in temperature were analysed by ToF-SIMS at 2 temperatures to see if any molecular structural changes could be detected at the surface that cause the temperature induced  $\Delta$ WCA. It is important to note that these measurements are obtained in ultra high vacuum conditions and relating them to other environments, for example in aqueous conditions, should be done with caution. Nevertheless, any surface enrichment of ions is likely to be indicative of changes that occur at the surface at ambient conditions. A subset of characteristic ions with the largest relative change in intensity when the temperature of the materials was changed between 5 °C and 37 °C is shown in Table 1 (the corresponding full list of ions is shown in Table SI1). For copolymers L(85%)J(15%) and L(70%)J(30%) and the homopolymer of monomer L, an increase in intensity was observed at low temperature for ions originating from monomer L, such as ions  $\text{CHNO}^-$  and  $\text{C}_8\text{H}_{13}\text{NO}_3^-$ , and from the acrylate/methacrylates backbone, such as  $\text{CHO}_2^-$ ,  $\text{C}_2\text{H}_2^+$  and  $\text{CH}_3^+$ . These results suggest at 5 °C the whole monomer L side-group is surface enriched. At high temperature an increase in intensity was observed for ions originating from ethylene glycol moieties on both monomers L and J, such as  $\text{C}_2\text{H}_5\text{O}_2^-$ ,  $\text{C}_3\text{H}_7\text{O}^+$  and  $\text{C}_4\text{H}_5\text{O}_2^-$ , and for ions from the terminus of monomer L, such as  $\text{C}_3\text{H}_7^+$  and  $\text{C}_2\text{H}_4\text{N}^+$ . As these polymers also show a decreased contact angle at 37 °C (Figure 2D) it is likely that with an increased temperature leads to the surface enrichment of hydrophilic groups such as ethylene glycol and di(ethylene glycol). Taken together, these results suggest that the polymer pendant groups are surface-enriched at higher temperature, likely due to reduced intramolecular interactions. However at reduced temperature the monomer backbone is surface enriched as intramolecular interactions dominate and cause rotational movement of the side groups towards the polymer bulk. This is likely caused by the temperature increasing above the polymer's upper critical solution temperature (UCST). This conformational change is depicted schematically in Figure 2E, and is similar to conformational changes observed on pNIPAM.<sup>[25]</sup>

An increase in WCA was measured for the copolymer of V and L with increasing temperature (Figure 2C), which differs from the other 3 polymers studied. Analysis by ToF-SIMS revealed an increase in the intensity of characteristic ions  $\text{C}_7\text{H}_{15}\text{NO}_2^+$  and  $\text{C}_3\text{H}_7\text{NO}^+$  from monomer L and ions  $\text{C}_4\text{H}_9\text{O}^+$  and  $\text{C}_5\text{H}_9\text{O}_2^+$  from propylene glycols with increased temperature. With a decreased temperature the  $\text{C}_4\text{H}_7^+$  and  $\text{C}_5\text{H}_9^+$  ions from aliphatic carbon and ions  $\text{C}_5\text{H}_{10}\text{N}^+$  and  $\text{C}_4\text{H}_7\text{N}^+$  from the terminus of monomer L were found to increase. Monomer V (Figure 2C) is a diacrylate and is thus less mobile than monomer

L, thus any conformational changes within this polymer likely result from a rearrangement of monomer L. This suggests that the pendant group of monomer L is surface enriched and possibly upright at lower temperatures whilst at higher temperature monomer V is exposed at the surface as monomer L is rotated towards the bulk.

## Conclusion

A high throughput methodology has been demonstrated to identify thermally responsive materials based upon altered hydrophilicity. This approach has been applied to polymer microarrays, resulting in the discovery of novel switchable materials L(100%), L(85%)J(15%), L(70%)J(30%) and V(70%)L(30%) that were scaled up to polymer coupons whilst preserving their stimuli responsive nature. ToF-SIMS analysis provided insight into the molecular conformation changes that cause the temperature responsive  $\Delta$ WCA. Specifically, the copolymers of monomer J and L alter between an extended and collapsed surface conformation when the temperature is varied from 5 °C to 40 °C. This study, which utilised ToF-SIMS with a temperature-controlled stage, represents a novel way to investigate the surface wettability changes of thermo-responsive materials and thus understand their interactions with cells and proteins.

## Acknowledgements

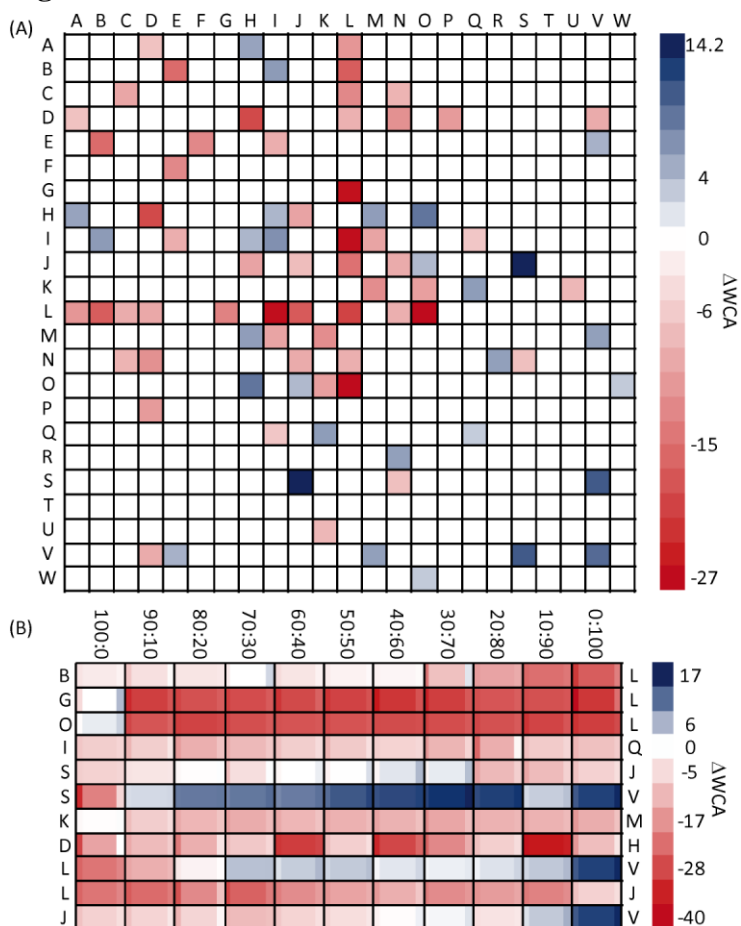
Experimental assistance with material scale up by Josephine Wong Yunn Shyuan and Ka To Fung is kindly acknowledged. Funding from the Wellcome Trust (Grant number 085245/Z/08/Z) is kindly acknowledged.

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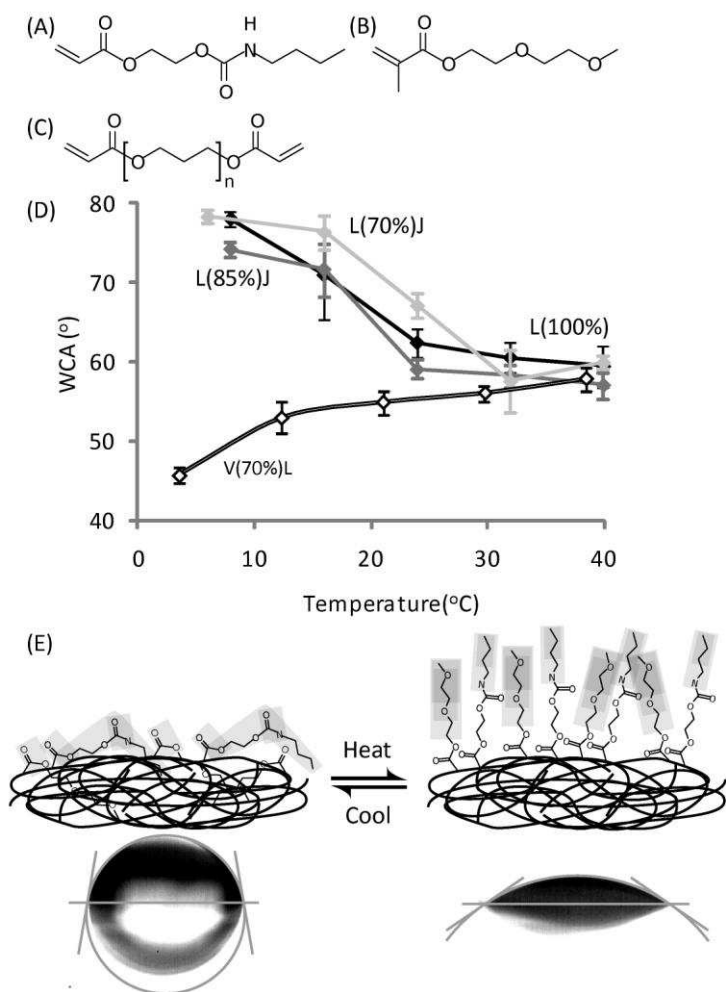
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## Figures



**Figure 1.** Intensity map showing the change in WCA when temperature is switched from 8 to 40 °C. On (A) the first generation array and (B) the second generation array. Blue indicates a positive shift whilst red indicates a negative shift as indicated by the intensity scale. Materials shown in white had a shift in WCA below the LOD (3 times the standard deviation of repeated measurements on the pHEMA background). Monomers are indicated by a letter. For (A) monomers were mixed at a 50:50 ratio. For (B) the monomer composition ratios are indicated across the top of the figure and denote the ratio between the two monomers indicated as a letter on the left and right of the figure. The large block indicates the value of the change in WCA whilst the small blocks to the left and right of the large block indicate the mean  $\pm$  the standard deviation,  $n = 3$ .



**Figure 2.** (A-B) The chemical structure of monomers (A) L, (B) J and (C) V. (D) The WCA measured for each of the polymer compositions for temperatures of 8-40 °C. Error bars equal  $\pm$  one standard deviation;  $n = 9$ . The monomer compositions studied were L(100%) (◆), L(85%)J(15%) (◆), L(70%)J(30%) (◆), V(70%)L(30%) (◇). (E) Schematic depiction of the molecular conformation of a copolymer of monomers L and J upon heating or cooling and the corresponding WCA measurements, which used a circle fit. The molecular fragments that ions enriched at each temperature are likely to have originated from are highlighted in grey.

**Table 1.** Summary of ions characteristic to each monomer with the highest relative change at the surface of polymer coupons at temperatures of 5 °C and 37 °C as detected by ToF-SIMS. The normalised (total ion count) ion intensities at both temperatures are shown. The top half shows the top ions that decreased with an increase in the temperature and the bottom half of the table shows ions that increased with increasing temperature.

| L(70%)J(30%)   |         |         | L(85%)J(15%)   |         |         | L(100%)   |         |         | V(70%)L(30%)  |         |         |
|--|---------|---------|--|---------|---------|---|---------|---------|---|---------|---------|
| Ion  | 5 °C    | 37 °C   | Ion  | 5 °C    | 37 °C   | Ion   | 5 °C    | 37 °C   | Ion   | 5 °C    | 37 °C   |
| CHNO <sup>-</sup>  | 0.00285 | 0.00194 | C <sub>2</sub> H <sub>2</sub> <sup>+</sup>                 | 0.00782 | 0.00234 | C <sub>8</sub> H <sub>13</sub> NO <sub>3</sub> <sup>-</sup> | 0.01177 | 0.00628 | C <sub>5</sub> H <sub>10</sub> N <sup>+</sup>               | 0.00151 | 0.00083 |
| C <sub>2</sub> H <sub>5</sub> NO <sub>2</sub> <sup>+</sup> | 0.00102 | 0.00070 | C <sub>3</sub> H <sub>2</sub> <sup>+</sup>                 | 0.00308 | 0.00126 | C <sub>8</sub> H <sub>11</sub> NO <sub>3</sub> <sup>-</sup> | 0.00092 | 0.00051 | C <sub>5</sub> H <sub>9</sub> <sup>+</sup>                  | 0.00281 | 0.00188 |
| CHO <sub>2</sub> <sup>-</sup>                              | 0.02198 | 0.01577 | C <sub>4</sub> H <sub>2</sub> <sup>+</sup>                 | 0.00216 | 0.00110 | CHO <sub>2</sub> <sup>-</sup>                               | 0.03330 | 0.02163 | C <sub>4</sub> H <sub>7</sub> N <sup>+</sup>                | 0.00104 | 0.00071 |
|  |         |         |  |         |         |   |         |         | C <sub>4</sub> H <sub>7</sub> <sup>+</sup>                  | 0.01251 | 0.00874 |
| C <sub>2</sub> H <sub>5</sub> NO <sup>+</sup>              | 0.00145 | 0.00184 | C <sub>4</sub> H <sub>3</sub> NO <sub>2</sub> <sup>-</sup> | 0.00005 | 0.00017 | C <sub>2</sub> H <sub>5</sub> NO <sup>+</sup>               | 0.00122 | 0.00257 | C <sub>7</sub> H <sub>15</sub> NO <sub>2</sub> <sup>+</sup> | 0.01063 | 0.01728 |
| C <sub>2</sub> H <sub>5</sub> O <sub>2</sub> <sup>-</sup>  | 0.01802 | 0.02247 | C <sub>3</sub> H <sub>7</sub> O <sup>+</sup>               | 0.00444 | 0.01457 | C <sub>3</sub> H <sub>7</sub> <sup>+</sup>                  | 0.00716 | 0.01250 | C <sub>4</sub> H <sub>5</sub> O <sup>+</sup>                | 0.00165 | 0.00259 |
|  |         |         | C <sub>2</sub> H <sub>5</sub> O <sub>2</sub> <sup>-</sup>  | 0.01120 | 0.02794 | C <sub>2</sub> H <sub>3</sub> O <sub>2</sub> <sup>+</sup>   | 0.00369 | 0.00442 | C <sub>3</sub> H <sub>7</sub> NO <sup>+</sup>               | 0.00345 | 0.00522 |
|  |         |         |  |         |         |   |         |         | C <sub>5</sub> H <sub>9</sub> O <sub>2</sub> <sup>+</sup>   | 0.00187 | 0.00269 |

Table 1