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Biomedical applications of inkjet printing at the UoN



Engineering and Physical Sciences Research Council



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Why Material Jetting?

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- Classification of the AM techniques (ASTM, 2012)
 - Binder Jetting
 - Powder Bed Fusion
 - Directed Energy Deposition
 - Sheet Lamination
 - Vat Photopolymerization
 - Material Extrusion
 - Material Jetting





Why Material Jetting?



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 Material Jetting: Droplets of build material are selectively deposited





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Printability requirement

 $Z = \frac{\sqrt{\gamma \rho a}}{2}$



Fluid printability is defined by parameter Z





- Characteristic length а
- Dynamic viscosity n



Stable droplet formation







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Stratasys Objet Connex 260

Dimatix DMP-2800 and DMP-2831





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PiXDRO LP50





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Nordson PICO





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Material Jetting at UoN

- Direct Printing
 - Jetting of Electronic Tracks
 - Inkjet printing of polyimide

Ag PI

Ag

- Reactive Printing
 - Reactive Jetting



Head 2

Substrate

Monomer B

Polymer

Head 1

000

Monomer A



Resistance < 1Ω



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- Reactive Printing
 - Photopolymers
 - Polycaprolactone: inkjet printing of bioresorbable material
 - 3D printing of materials with resistant to bacterial attachment
 - 3D printable biodegradable materials
 - Polymer particle formation using inkjet printing
 - 3D printing of human tissues



Polycaprolactone: inkjet printing of bioresorbable material

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- Polycaprolactone is a bicompatible, biodegradable and bioresorbable material
- Mostly 3D printed by extrusion or energy deposition technique
- Lack of inks based on bioresorbable materials for inkjet printing
- Improvement of printing resolution
- Possibility of multimaterials

[1] Seyednejad, H. et al. Preparation and Characterization of a 3D-printed Scaffold Based on a Functionalized Polyester for Bone Tissue Engineering Application; Functionalized Polyesters 7, 2012, 5, 87.

[2] Williams, J. et al. Bone tissue engineering using polycaprolactone scaffolds fabricated visa selective laser sintering; Biomaterials, 2005, vol26, p.p: 4817-4827



PCL printed by material extrusion technique [1]



PCL printed by direct energy deposition technique [2]



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Polycaprolactone: inkjet printing of bioresorbable material



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 Synthesis: Preparation of UV curable Polycaprolactone-derivative (PCLDMA)



Printability assessment and modification



Polycaprolactone: inkjet printing of bioresorbable material

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- Printability assessment and modification
 - Poly(ethylene glycol) diacrylate was used as diluent.
 - The viscosity of PCLDMA with different diluent proportion was measured
 - Based on the viscosity measurement,
 PCLDMA:PEGDA =70:30 was chosen as the

printing composition



Temperature	PCLDMA:PEGDA Proportion					
	50:50	60:40	70:30	80:20	90:10	100:0
30 °C	29.22±1.00	34.97±1.00	40.07±0.97	50.45±0.98	69.68±0.95	96.14±0.92
35 °C	24.44±1.01	29.00±0.98	33.00±0.99	41.25±0.96	56.47±1.00	76.96±0.93
40 °C	20.90±1.00	24.44±1.00	27.58±1.00	34.21±0.99	46.45±0.99	62.64±0.94
45 °C	18.24±1.02	20.99±1.00	23.41±1.01	28.81±0.98	39.15±0.98	51.78±0.97
50 °C	16.18±1.02	18.33±1.03	20.15±1.03	24.51±1.00	33.05±0.99	43.28±1.00
55 °C	14.56±1.02	16.25±1.02	17.60±1.03	21.14±0.99	28.24±1.01	36.62±1.00
60 °C	13.30±1.02	14.65±1.02	15.63±1.03	18.84±0.99	24.48±1.00	31.48±1.00

Polycaprolactone: inkjet printing of bioresorbable material



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Surface profile of printed mesh samples

SEM pictures of printed mesh structure with different wall thickness: (a) 150µm, (b) 300µm, (c) 500µm, (d) winkle found at sample surface



- Most healthcare-associated infections are associated with biofilms which form on the surfaces of medical devices.
- These microbial colonies develop up to 1000 times higher tolerances to antibiotic treatment and the host immune system compared with their planktonic counterparts
- The discovery of new materials with resistance to bacterial attachment opens a door to the fabrication of bacteria-free implants, with a direct reduction in the incidence rate of infections and therefore an increment in the cost saving.







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 Alexander, M et al.; Discovery of novel materials with broad resistance to bacterial attachment using combinatorial polymer microarrays, *Advanced Materials* 2013, 25, 2542-47







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 The aim of this work is to develop a methodology to inkjet print customised parts for biomedical applications with graded compliance and such antibacterial property, using a combination of inks with widely separated moduli.









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- Printability of both inks tested.
- First trial for gradation of properties using comercial inks

Image 2: Gradation of color using comercial inks

100% Blue Ink

0% TGME



Image 1: Aprox. 120 µm wall thicknes cylinder printed with the rigid monomer



3D printable biodegradable materials



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- 3D printing technologies allows a precise shape and morphology for personalised structures.
- In order to completely optimize the material to the application, new families of safe biomaterials for personalized medicine need to be developed

Synthetic Biodegradable and Photocurable polymer-

based Materials







3D printable biodegradable materials



- Synthesis of macromonomers
 - D., G. Anderson, C. A. Tweedie, N. Hossain, S. M. Navarro, D. M. Brey, K. J. Van Vliet, R. Langer, J. A. Burdick, *Adv. Mater.* **2006**, *18*, 2614–2618.



- Biodegradable (24h- > 3 months)Very high viscosity
- Solution: Ink formulation by mixing the macromers with photocurable degradable monomers of low viscosity.
- A combinatorial approach is going to be used for screening





3D printable biodegradable materials



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2D printing of spots





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3D printable biodegradable materials



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Polymer particle formation using inkjet printing



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 Design of new materials is hindered by the lack of knowledge on the physiochemical parameters controlling the range of cellular responses required of modern devices



Polymer particle formation using inkjet printing



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Polymer particle formation using 3D printer Dimatix



3D printing of human tissues



- Aim of the project: Build a tissue constructs by directed cell deposition
- Vasculature, cells, and extra-cellular matrix (ECM) are co-printed to yield engineered tissue constructs composed of heterogeneous subunits
- The vascular tissue is later dissolved or evacuated to let nutritious perfused through the empty fluidic channels.



D. B. Kolesky , R. L. Truby , A. S. Gladman , T. A. Busbee , K. A. Homan , J. A. Lewis , Adv. Mater. 2014 , 26 , 3124 .

3D printing of human tissues



 Our Group proposal: 3D printing of gelatin derivatives as structural and vascular tissues, using high viscosity jetting valve system (PicoDot Nordson)



Photron	FASTCAM-APX RS model 1/500000 sec frame : -115 Time : 14:36	Partition : 003 128 x 432 -5.111 ms	22500 fps Center Date : 2015/5/1
		J	



3D printing of human tissues



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Gelatine pillars, 10 layers, 15x15 mm²

Gelatine cube, 10 layers





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3D printed smart limbs



- 3D-printed pressure sensitive finger
 - UV curable polymeric ink forms the finger structure.
 - 3 silver tracks were inkjet printed to power up a pressure sensor in the finger tip and also collect pressure data.

Pressure sensor was inserted and a rubbery cap was printed to seal the pressure around the sensor



3D printed smart limbs



- All 3D-printed capacitive touch sensitive hand.
 - UV curable polymeric ink forms the hand structure.
 - Touch sensitive silver tracks were printed within the hand structure connecting a data line to a microprocessor at the bottom of the hand.

Silver ink printed inside the Objet VeroClear structure. The conductive track acts as a capacitive touch sensor to activate embedded LEDs.





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Thank you very much

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