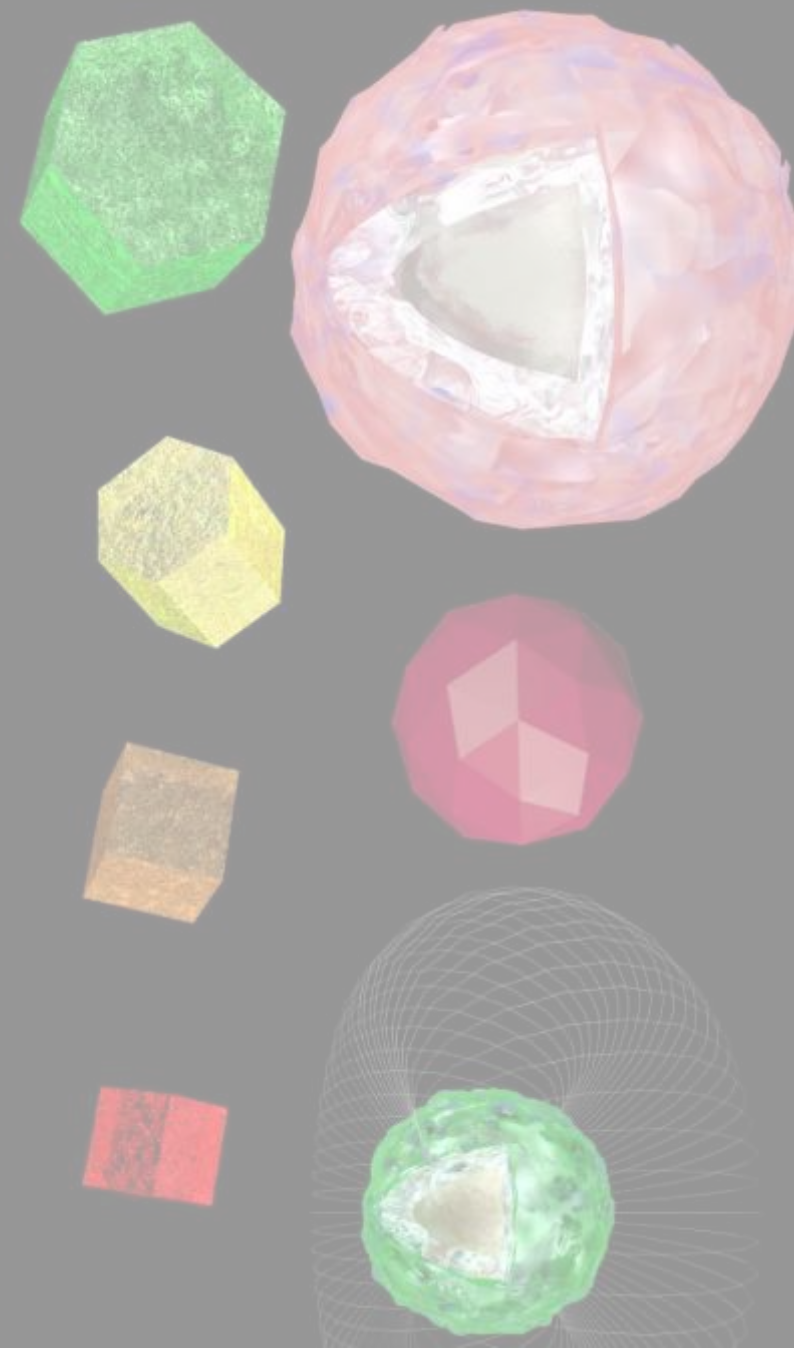
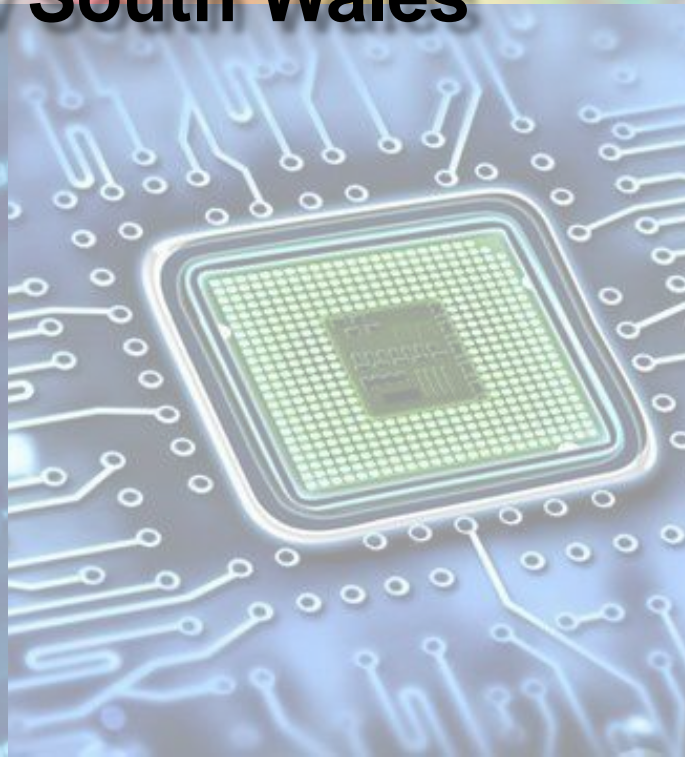


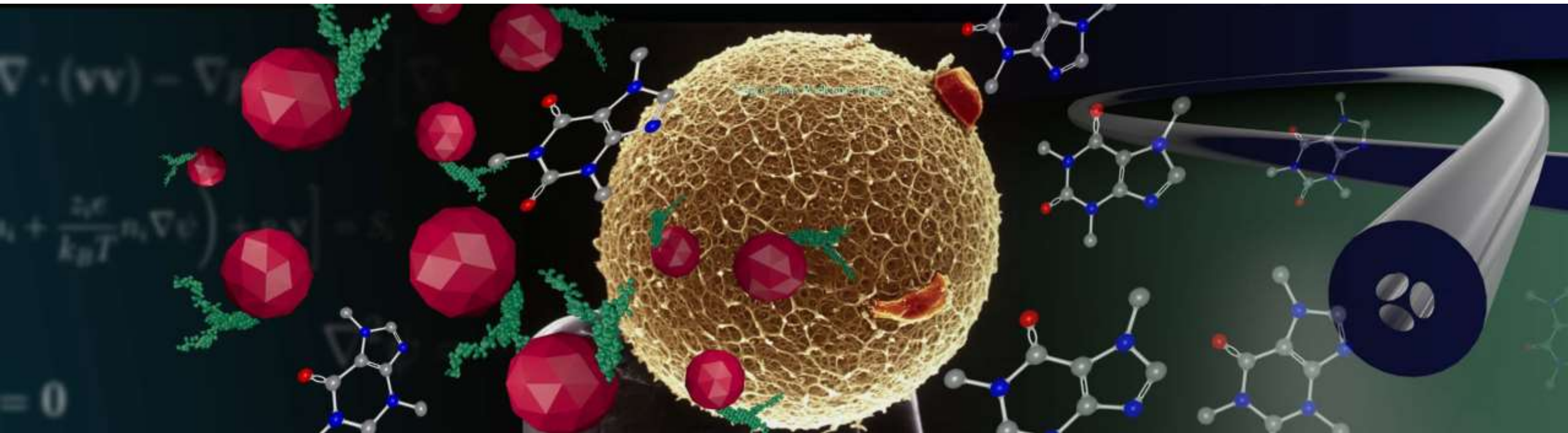
**Prof Ewa Goldys,
Deputy Director,
ARC Centre of Excellence
for Nanoscale Biophotonics
University of New South Wales**

```
parameters.contains("age")  
+= " and p.age = :age";  
typedQuery<Person> query = em.createQuery(hql, Person.class);  
if(parameters.contains("name")){  
    query.setParameter("name", values[0].toString());  
}  
if(parameters.contains("age")){  
    query.setParameter("age", Integer.valueOf(values[1]));  
}  
query.list();
```



Core aim of ARC Centre for Nanoscale Biophotonics

New approaches to measuring nano-scale dynamic phenomena
in living systems



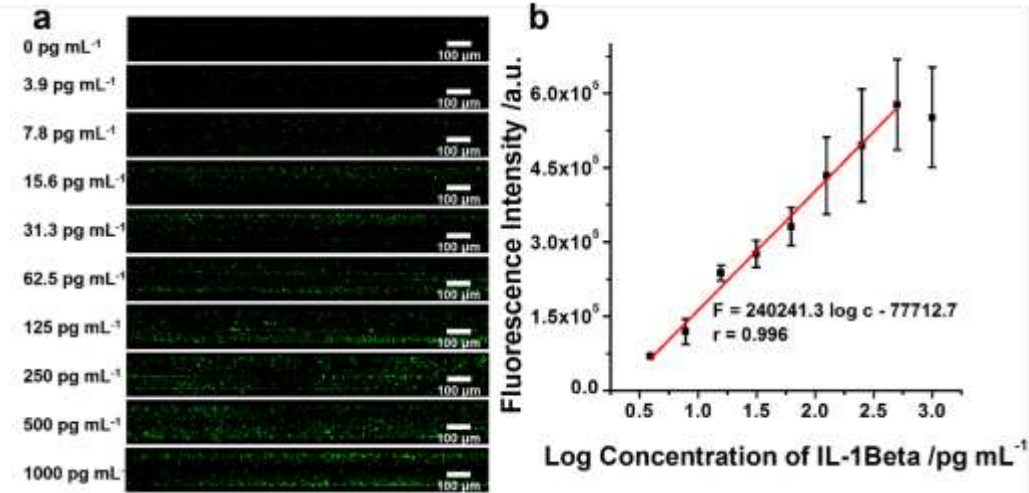
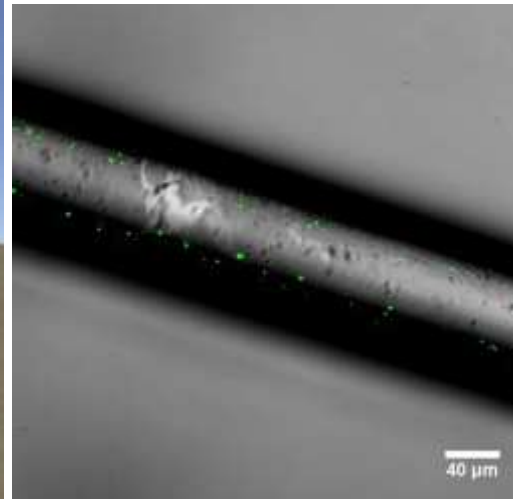
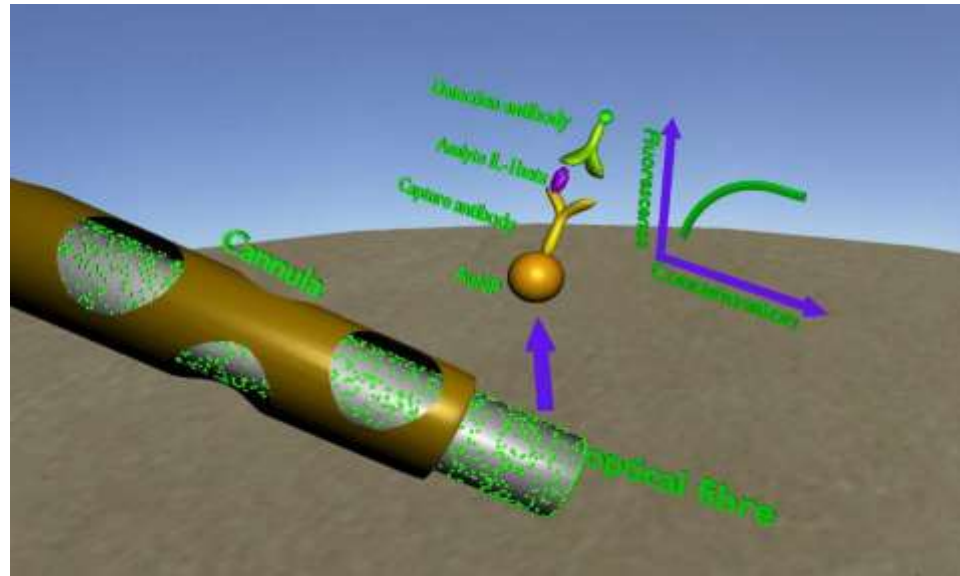
in-vivo

real-time

portable? point-of-care?

BIOMOLECULAR ANALYSIS

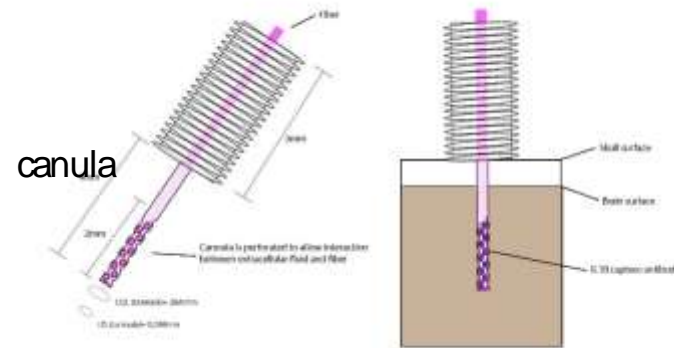
How to detect cytokines in specific locations



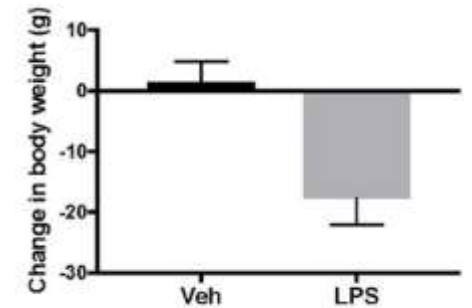
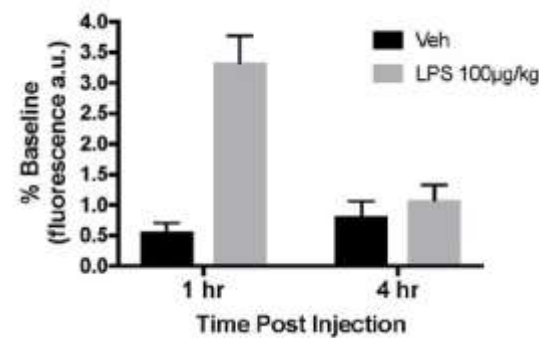
“sandwich immunoassay”,



K. Zhang, Guozhen Liu, Mike Baratta,
Macquarie U U Colorado

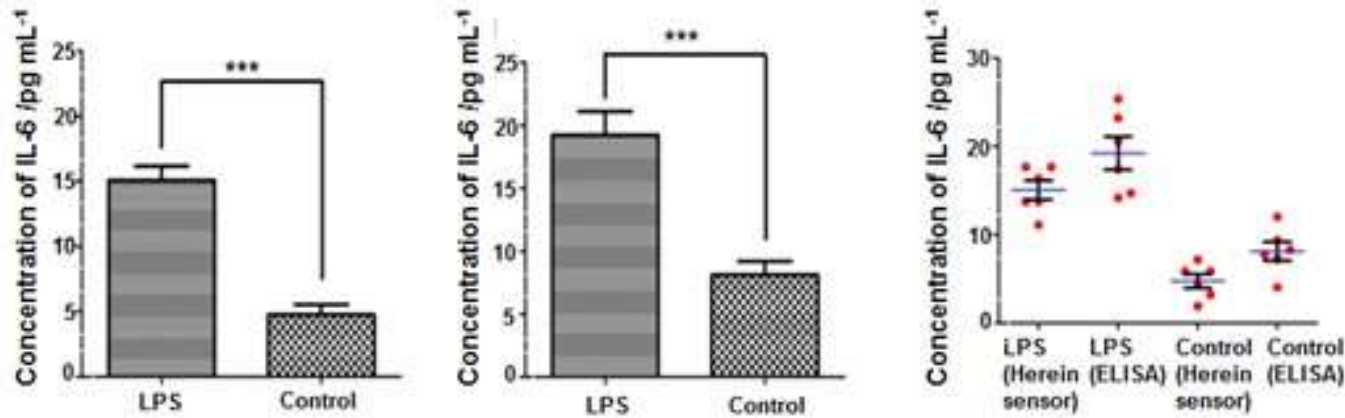
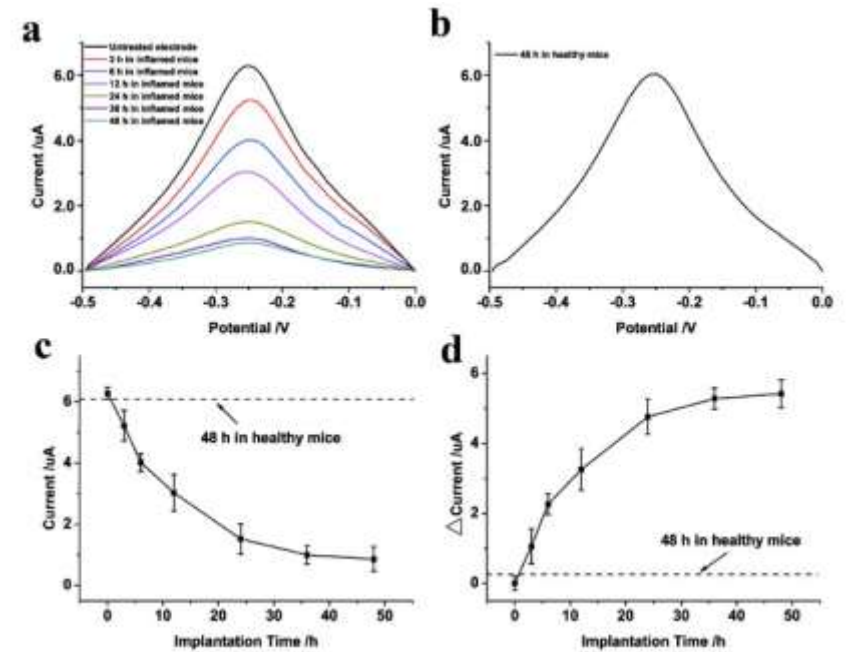
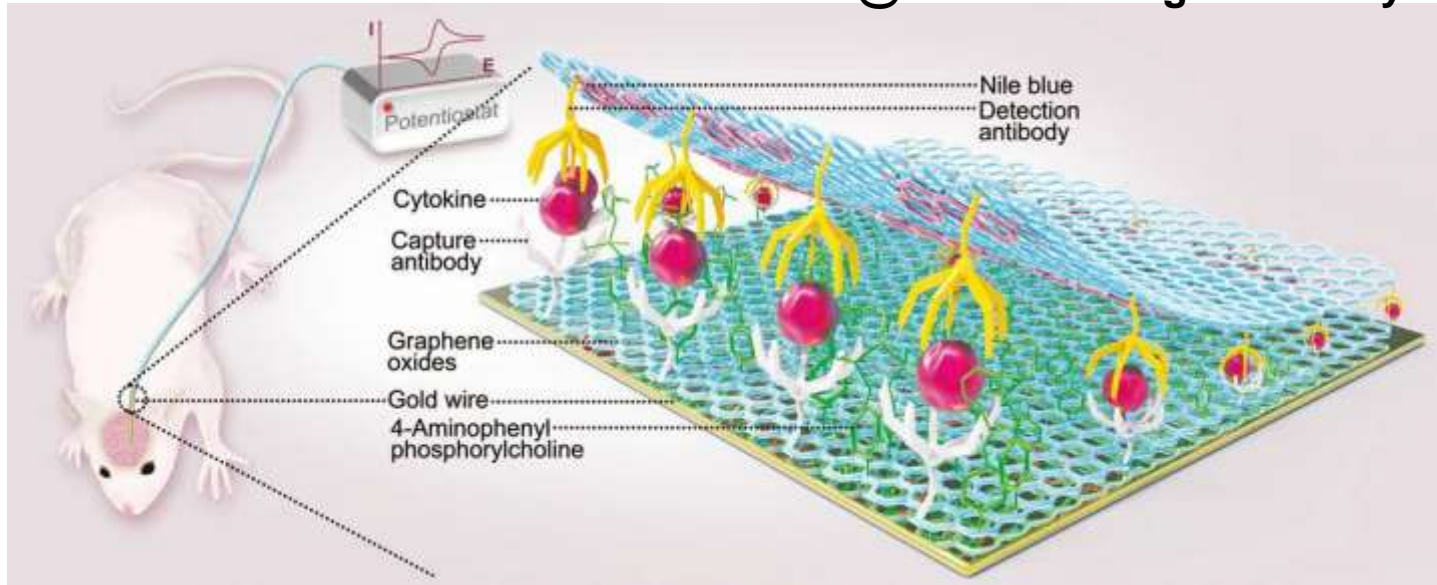


Living brain
(hippocampus)



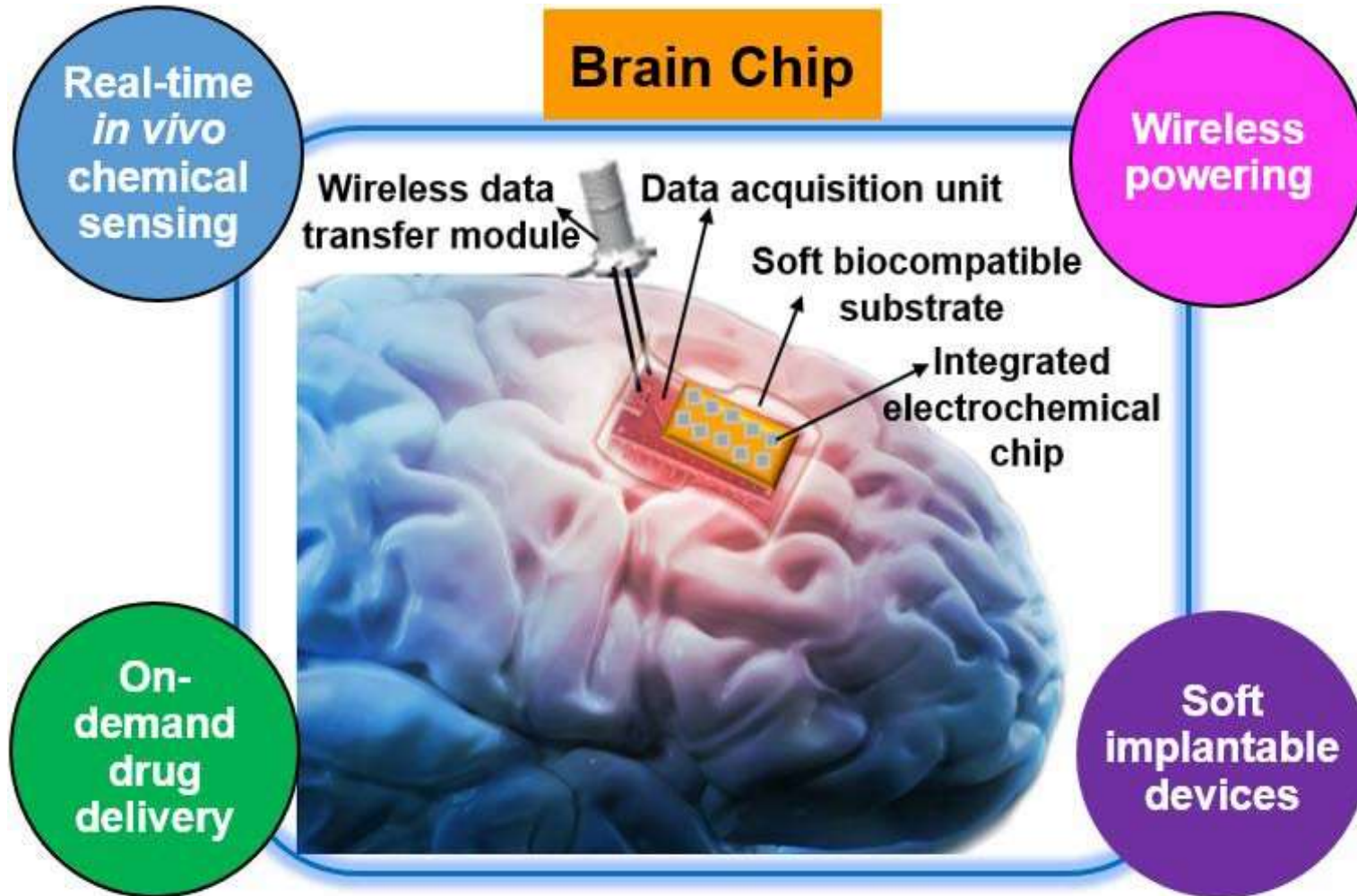
How to detect cytokines in specific locations

ARC Future Fellow, Dr Guozhen Liu, UNSW, A/Prof Rui Li@Centra China Normal University,
Prof Xin Chen@Xi'an Jiaotong University



Real-time sensing of cytokines on a relevant time scale has been demonstrated

VISION: WIRELESS IMPLANTABLE REAL-TIME SENSORS

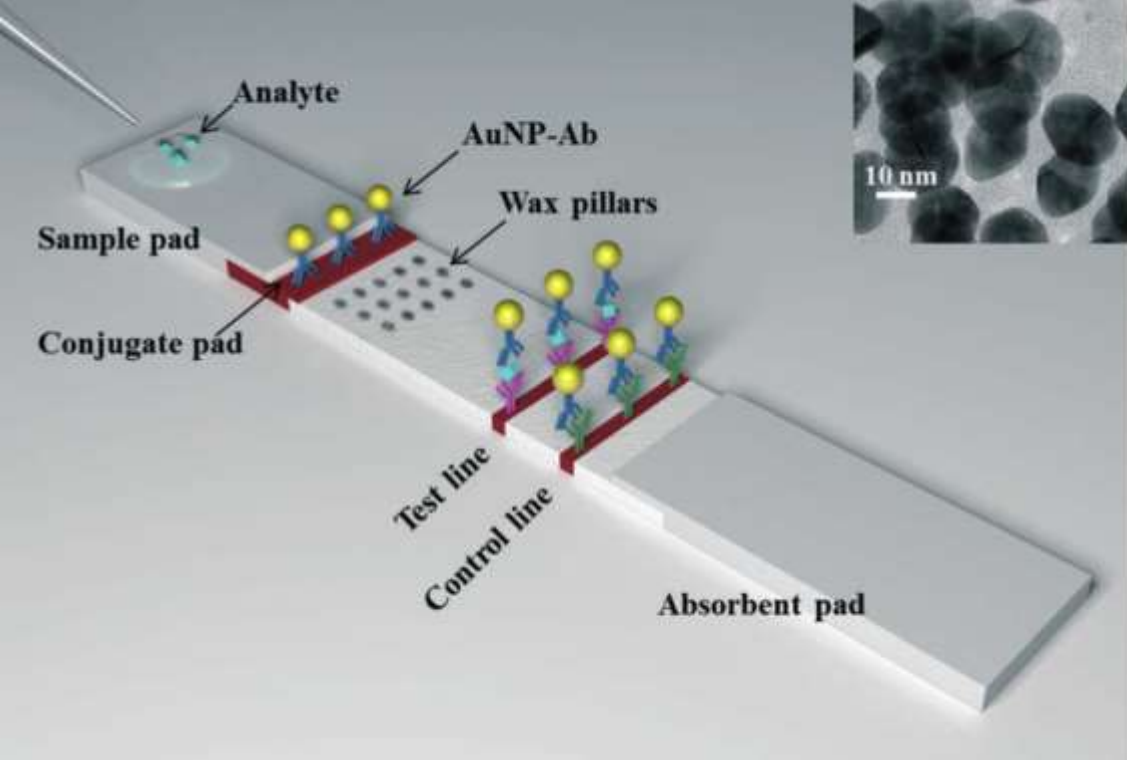


NEEDED:

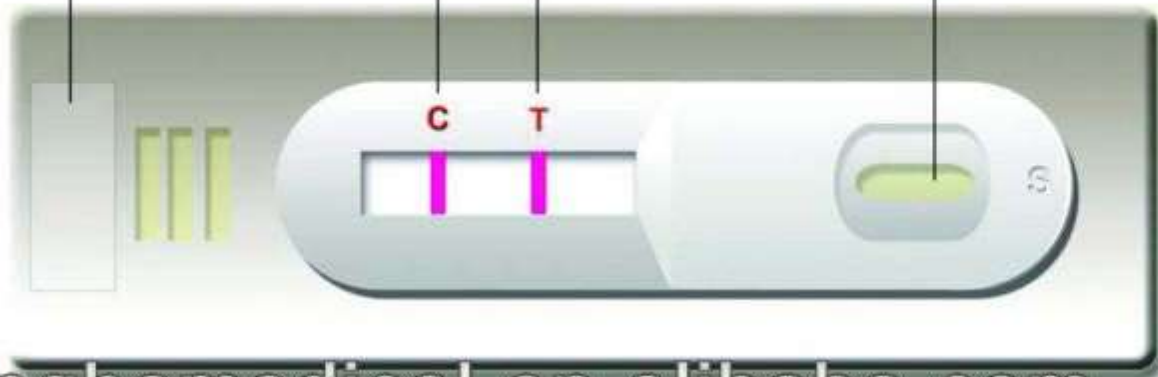
- Switchable, shape changing capture molecules to dynamically respond to analyte variations
- Capture molecules for specific targets, and they must be working well on surfaces
- Non-degradable capture molecules
- Amplification strategies (FETs?)
- Technology integration
- Protect against fibrosis over the long term
- Usual biocompatibility, degradability,

TARGETS: “Neuro” analytes – cytokines, neurotransmitters, metabolites

VISION: LATERAL FLOW ASSAYS



Specimen ID Control Line Test Line Sample Well



NEEDED:

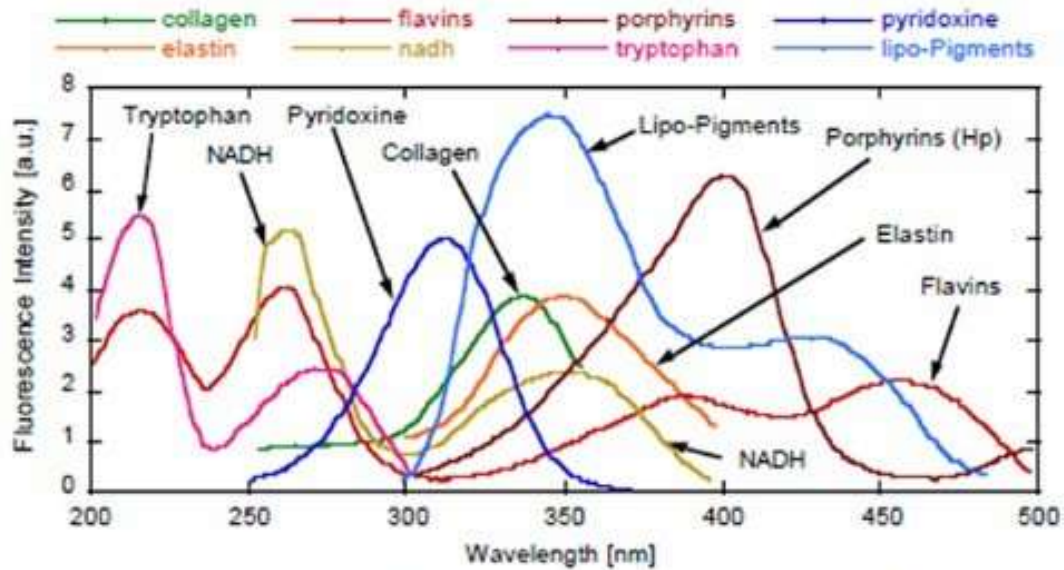
- Switchable, shape changing capture molecules to dynamically respond to analyte variations – may enable repetitive use of the same device
- Capture molecules for specific targets, and they must be working well on surfaces
- Non-degradable capture molecules
- **Amplification strategies** maybe based on nanomaterials
- Mobile phone readout (almost there)

TARGETS:

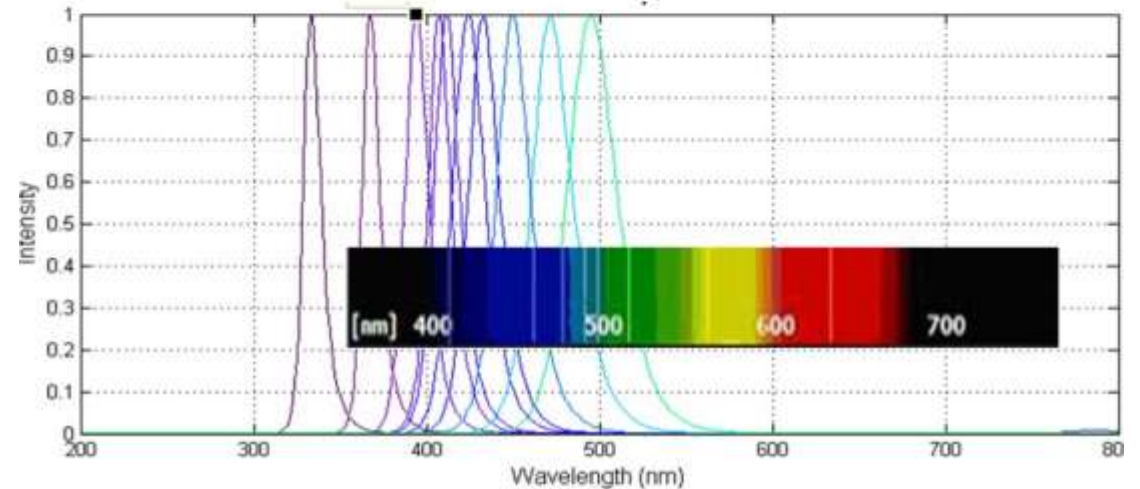
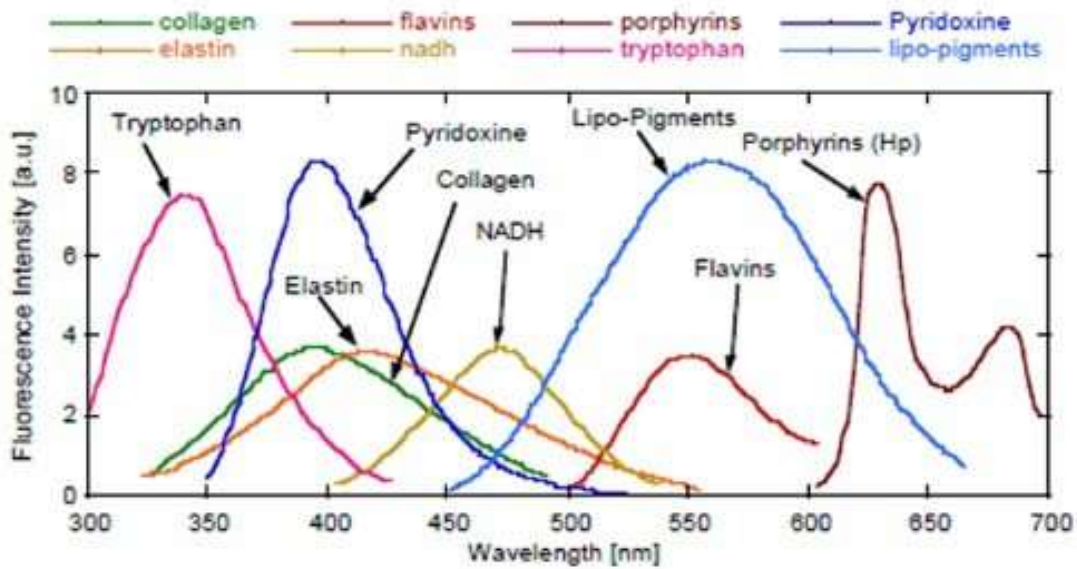
- “Neuro” analytes – cytokines, neurotransmitters, metabolites
- Bacteria and viruses
- DNA sequences
- Hormones
- Standard medical analytes, also those relevant to injury and acute conditions
- Warfare agents
- Toxins and contaminants
- Etc etc.

Autofluorescence analysis

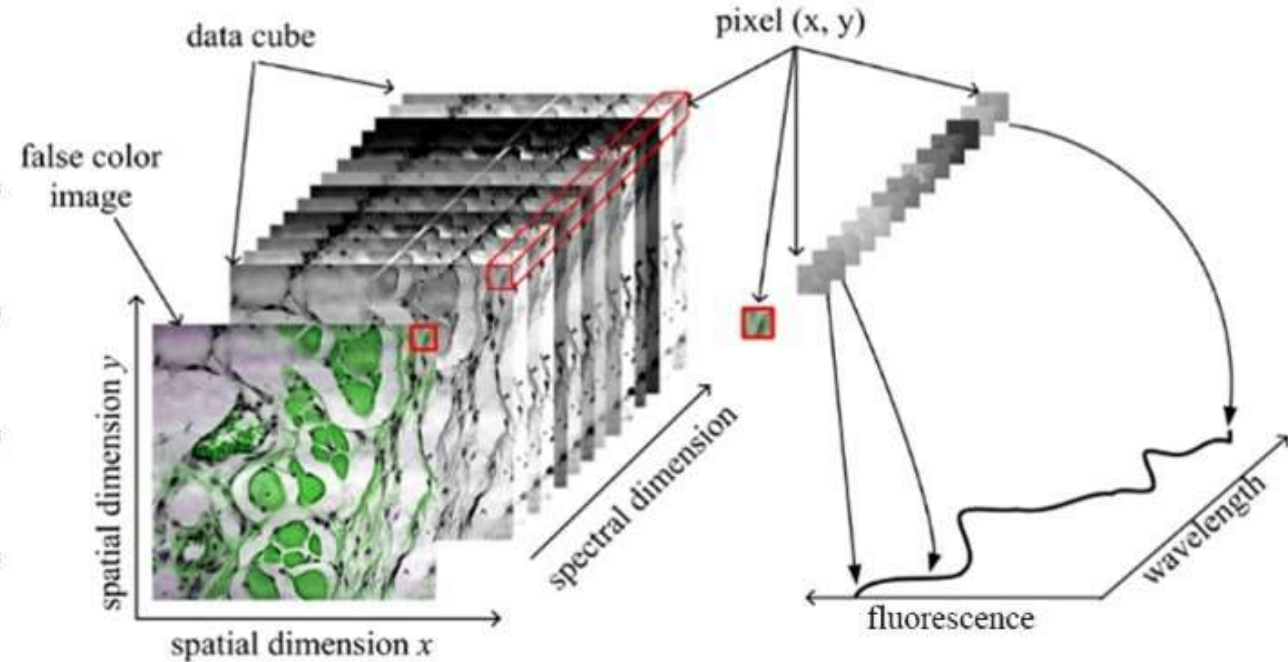
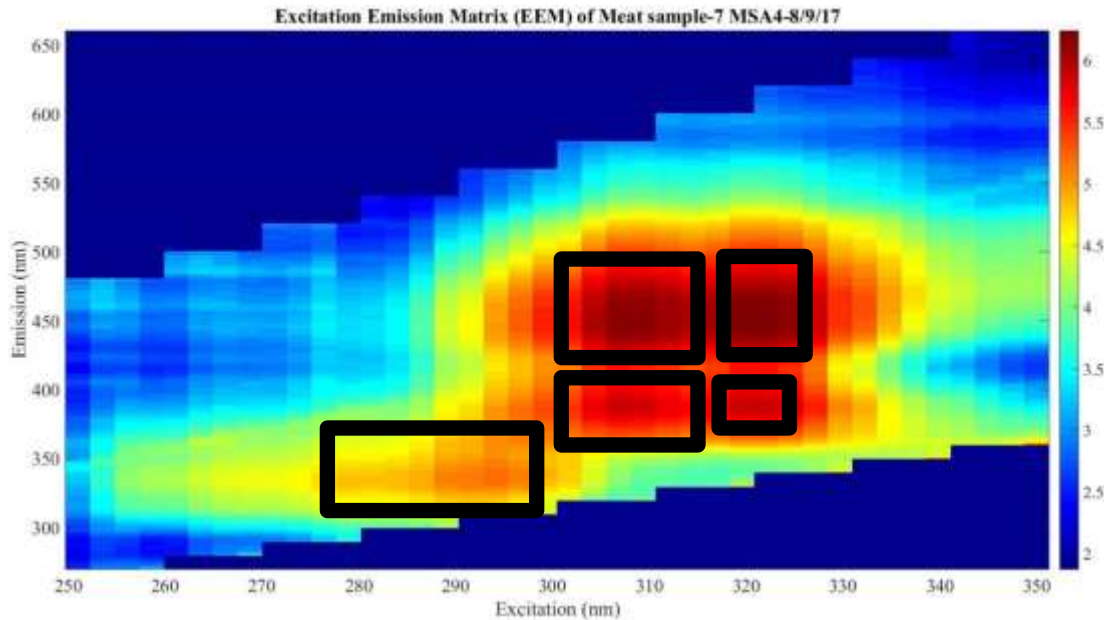
excitation



emission



Hyperspectral image data set



Cellular feature:

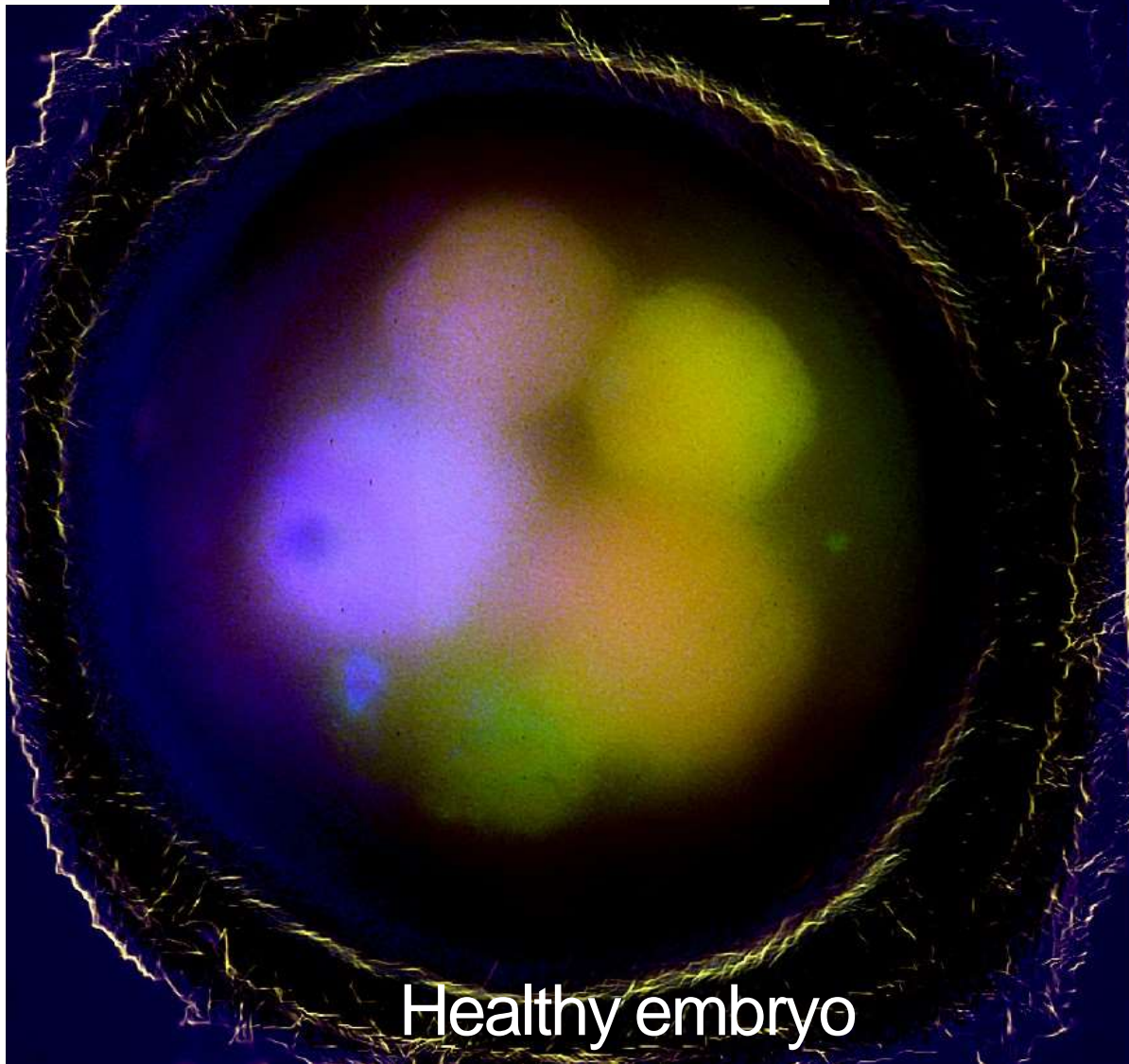
anything that can be calculated for a given cell

- average reading in that cell in a given spectral channel
- ratios of such readings for various channel pairs (channel ratios)
- Haralick textural features etc etc.

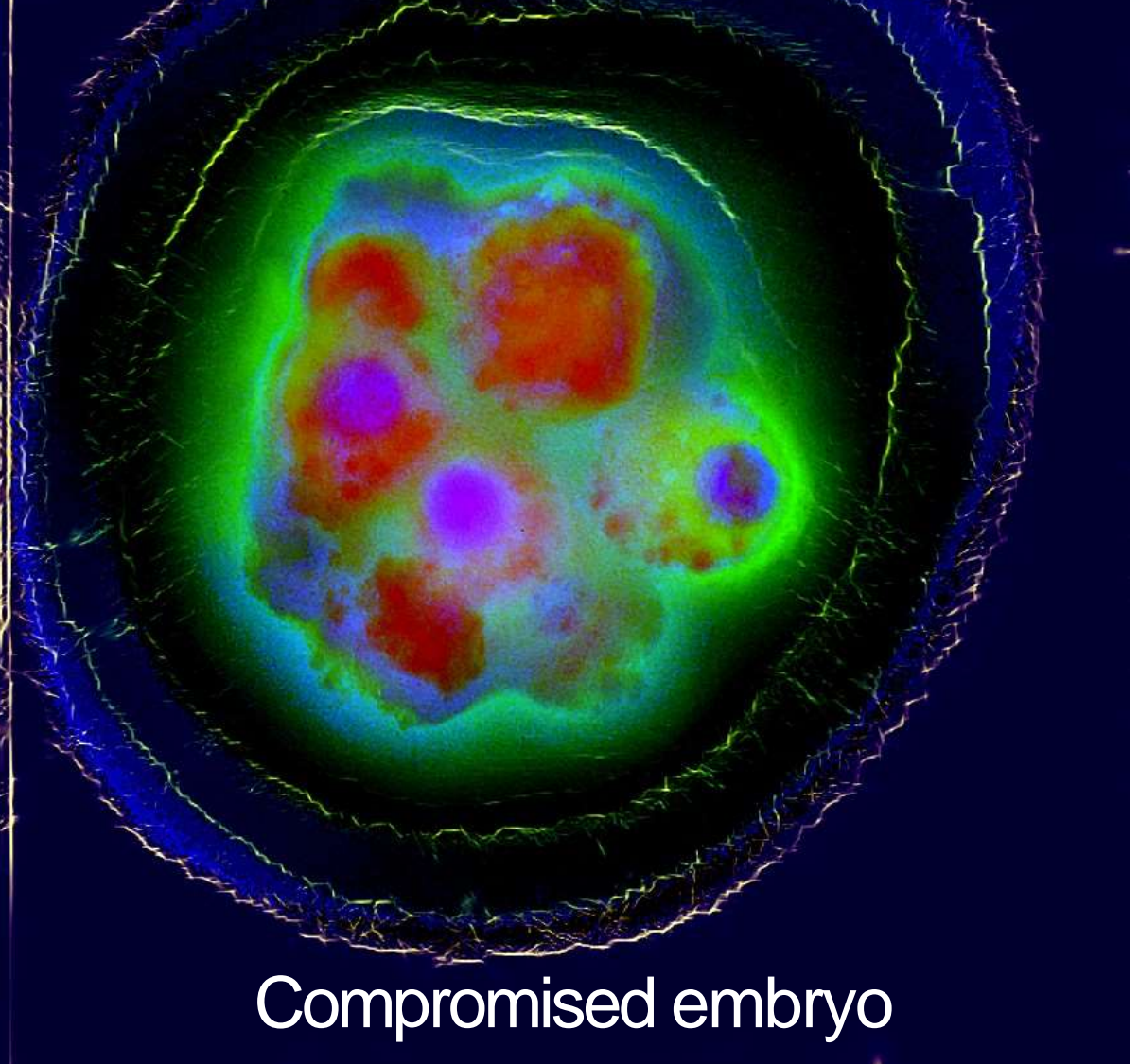
Example features: “squareness” – similarity to a square, length scale, colour



Healthy and compromised early bovine embryos



Healthy embryo



Compromised embryo

Wong-Baker FACES™ Pain Rating Scale Instructions For Usage

Explain to the person that each face is for a person who has no pain (hurt) or some, or a lot of pain.

Face 0 doesn't hurt at all. Face 2 hurts just a little bit. Face 4 hurts a little bit more. Face 6 hurts even more. Face 8 hurts a whole lot. Face 10 hurts as much as you can imagine, although you don't have to be crying to have this worst pain.

<5 = Panadol
>5 = Opiates

Ask the person to choose the face that best describes how much pain he has.



0

No Hurt



2

Hurts Little Bit



4

Hurts Little More



6

Hurts Even More



8

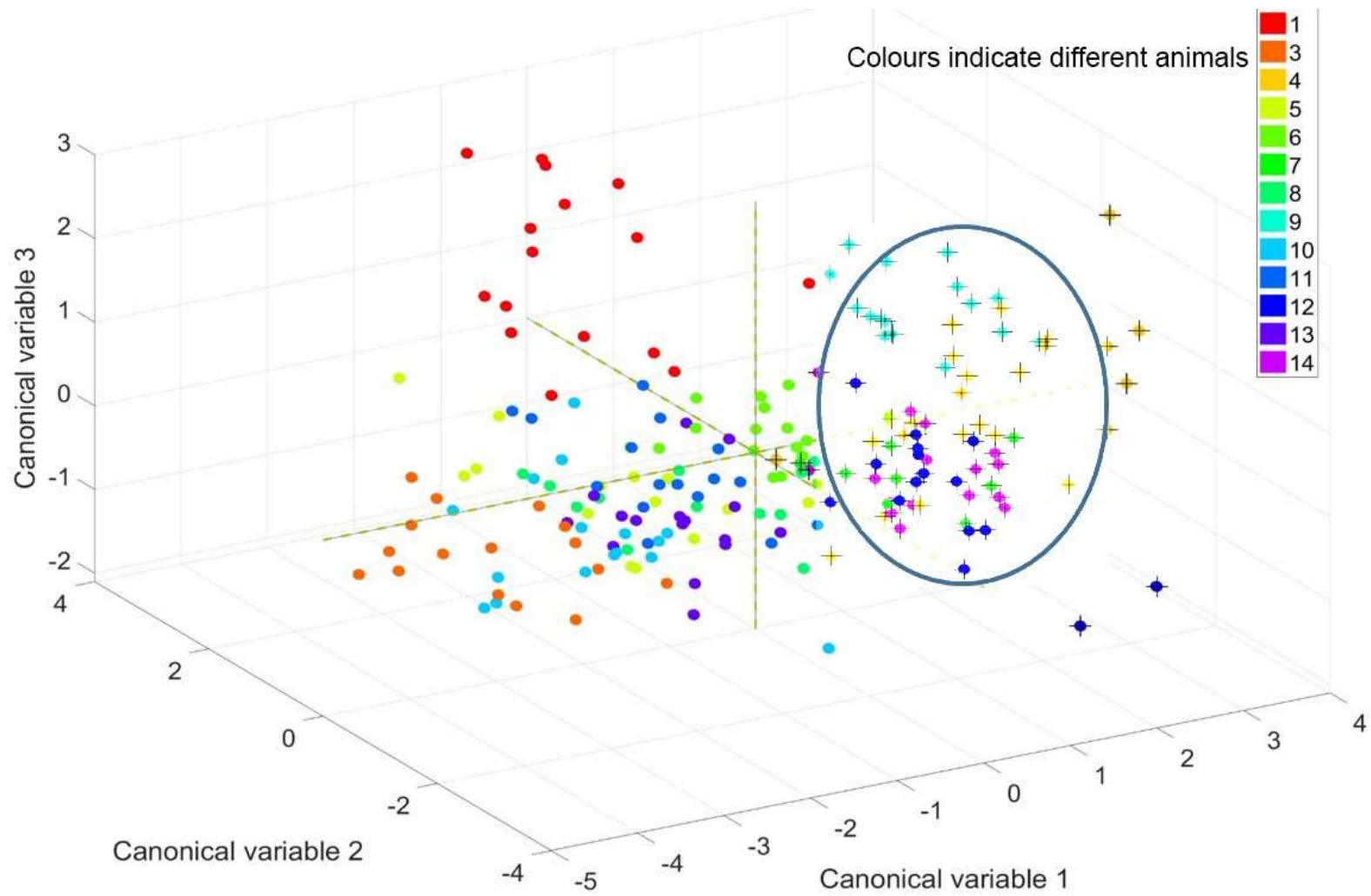
Hurts Whole Lot



10

Hurts Worst

Molecular signatures of pain



Vicky Staikopoulos, Mark Hutchinson, UAdelaide, Martin Gosnell, AyadAnwer, Macquarie U

"Hyperspectral imaging of endogenous fluorescent metabolic molecules to identify pain states in central nervous system tissue",
[Staikopoulos, V., et al., Proceedings Volume 10013, SPIEBioPhotonics Australasia; 1001306 \(2016\); doi: 10.1117/12.2243158](#)

VISION: HIGH CONTENT AUTOFLUORESCENCE IMAGING

NEEDED:

- Rapid multispectral imaging systems
- LED or laser based
- Improved ultrasensitive low noise cameras
- Mobile phone technologies
- Bespoke software (may need to be specific for each problem)
- Validation of methods

TARGETS:

- Characterisation of cells, tissues and underlying conditions (Cancer? Toxic exposure? Infection? Subtle immune effects? Sepsis? Microbial infection of wounds? Response to surgery? Transplantation? Pain?)
- “See right through” – autofluorescence in deep tissue (currently unexplored)
- Preparation-free histology
- Portable phone based devices for some of the above medical conditions

Acknowledgements



SAHMRI

persona EYES



ROYAL NORTH SHORE HOSPITAL



THE VIS ,Q\$PECIALISTS

