Turning Your Smart Phone into a Tricorder

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1. Background

- 2. Turning Your Smart Phone into a Tricorder
- 3. Tricoder Applications
- 4. Conclusion



Turning Your Smart Phone into a Tricorder





Disease Burden in Canada

Disease	Number of new	Number (%) of	Cost of
	cases per year	Canadian living	treatment
		with disease	(million \$)
Alzheimer	60,000	280,000 (0.74%)	\$431
Parkinson	5,500	100,000 (0.31%)	\$201
Stroke	50,000	300,000 (1%)	\$665
Heart Disease	1,200,000	16,000,000 (5%)	\$6,818
Asthma	138,000	2,744,000 (8%)	\$705
Atherosclerosis	112,000	544,000 (1.7%)	\$19,800
Schizophrenia	10,240	259,200 (0.81%)	\$1,637
Lung Cancer	25,600	39,884 (0.12%)	\$163
Prostate Cancer	25,500	150,135 (0.47%)	\$1,033
Breast Cancer	23,590	153,777 (0.48%)	\$1,161

Healthcare Mandates

• Diagnostic

Do you have a given disease/condition?

• **Prognostic**

How well will you do with this disease/condition?

• Predictive

Odds of getting a given disease/condition.

Environmental Influence



Protein

List of Metabolites

Glycerol	Indoxyl sulfate	Carnosine
Vitamin D	Choline	Vitamin B12
Phenylalanine	Lactate	Folate
Tyrosine	Glutamate	Taurine
Leucine	Glutamine	Bilirubin
Aminoadipic < acid	Glucose	ADMA
ТМАО	Formate	Pyruvate
Betaine	Aldosterone	Cortisol
Homocysteine	Testosterone	Dopamine
Uric acid	Estradiol	НРНРА

Utilizing Biomarkers

• Historically we have stuck with the idea one biomarker = one disease





This limits accuracy, precision and sensitivity/specificity Glucose (> 6.1 mM – diabetes)

Fatty Acid



Nat Med. 2011 Apr;17(4):448-53. Epub 2011 Mar 20.

Metabolite profiles and the risk of developing diabetes.

Wang TJ, Larson MG, Vasan RS, Cheng S, Rhee EP, McCabe E, Lewis GD, Fox CS, Jacques PF, Fernandez C, O'Donnell CJ, Carr SA, Mootha VK, Florez JC, Souza A, Melander O, Clish CB, Gerszten RE.

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Discovering Biomarkers

- Omics technologies now suggest many biomarkers = one disease
- This improves accuracy, precision and sensitivity/specificity
- The problem is the math (or biomarker model) gets much more complicated



Cholesterol (> 5.2 mM – CVD risk) Triglycerides (> 1.8 mM – CVD risk)

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TMAO	Formate	Pyruvate
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Discovering Biomarkers

• Historically we have stuck with the idea



hCG (human chorionic gonadotrophin) is a hormone secreted in pregnancy that is made by the developing embryo soon after conception. It is used for pregnancy test.

The Test Cost

Metabolite	Cost of Test
Betaine	\$113.70
Choline	\$100.00
Estradiol	\$125.00
Formate	\$84.00
Glucose	\$45.00
Glutamate	\$98.94
Glutamine	\$98.94
Glycerol	\$48.50
Taurine	\$98.94
Vitamin D	\$210.00

Total: \$1023.02 for 10 tests



What Are the (Standard) Metabolomic Technologies?







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- UPLC, HPLC
- CE/microfluidics
- LC-MS
- FT-MS
- QqQ-MS
- NMR spectroscopy
- X-ray crystallography
- GC-MS
- LIF detection
- Instruments are large (500 kg) and not portable
- Expensive (>\$1 million), hard to use (requires PhD)

The Problem...

- Metabolomic analysis is expensive, non-portable and difficult to do
- Need to make small (palm size), inexpensive (<\$1,000 for an instrument, \$1 for a test), easy-to-use (Smartphone application), portable (farm/industry friendly) devices

The Solution...

- Easy-to-use, platform specific kits
- Nanoelectromechanical devices (NEMS)
- Microfluidics
- Engineered biosensors
- Integration of extraction, separation & detection on a single platform



StarTrek & Tricorders



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	Name: John Smith Age: 26 Gender: Female Male
	Race: Mongoloid / Asian
the application will dis	play either a red exclamation mark

Analyzing



present in biofluids by using electrical impedance-based detection.



signifying that the patient's health is at risk and that they should seek medical attention



Proof-of-concept Design

- Synthesize streptavidin-poly-T
- Synthesize GNPs-poly-A
- Detect the signal on bio-molecule sensor





Gold Nanoparticle

Test Sequence: The structure of Poly A and Poly T

Experimental results



The amount of GNP-Poly A is 0.24, 0.48, 0.96, 1.44 nM

Electrode Sensor Chips (Nanotechnology)







nanofab.ualberta.ca

Analytical and Numerical Simulations



Sensitivity of 2D vs 3D simulations





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Conclusion

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