

“Innovating better health outcomes for a developing world”

The Narrative

- We want to develop new, better products that will meet unmet needs
- We should provide solutions to patient population that is more acutely aware of disease outcome than other populations
- We should provide early detection of pharmacodynamic outcomes
- We should provide clinicians with information to move patients onto alternative care pathways much earlier than we currently achieve
- Enhance the reputation of your company by your participation

The Consortium and its tractability

Stakeholder

- Patient advocates
- Regulators
- Therapy developers (“pharma”)
- Device and technology companies
- CROs

Role, contribution, benefit

- Emotional drive to adoption,
- Early adoption, incorporate regulation into design,
- £/product, philanthropy reputation
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- Service provision, new business

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Opportunity

- Work on high cost therapies, for example monoclonal antibodies
- The therapies are highly targeted, and need a personalized diagnosis
- Provides new business through development of a companion diagnostic
- Oncology is an area with severe patient outcome
- Paediatrics is an area where new sampling modalities are required
- Benefits will accrue from improved precision of the assessment/diagnostic method, that provides enhanced clinical outcome or care pathway choice
- Benefits accrue to companies, by segmenting out patients to trial on new therapies “find the needle in a haystack”
- Benefits accrue to the health economy by selecting responding patients
- Benefits accrue by opening currently closed markets e.g. work WITH China

Risk/Benefit Calculations

- Risk management is a barrier to innovation
- Failure of technology is not failure – carry out failure mode analysis
- Understand probability of success, and assess patient rejection
- Understand protection & ownership of data
- Difficulty in using ‘old’ drugs, may reveal new unknowns about your drug with regulatory implications.