WORKSHOP A: PRECISION MEDICINE

	Who must be involved	Who can initiate?
Data standards	clinicians, scientists (IT), legal system	scientists (IT)
Cross-disciplinary collaboration	clinicians, scientists	clinicians, scientists
Prospective clinical trials	clinicians, pharma industry	clinicians, pharma industry
New improved consent handling	clinicians, IT, ethicists, legal system	ethicists, IT, legal system

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	How will this: ...contribute to more efficient cancer care?	... contribute to more equal cancer care?	... improve the individual patient's quality of life?	... influence which health decisions the patient and her kin can make?
Data standards	Common rules, interoperability	Common rules, better informed patients	Improved access to own health data	Improved access to own health data
Cross-disciplinary collaboration	Comprehensive result delivery	More precise understanding	Comprehensive knowledge of own health	Improved understanding of own health
Prospective clinical trials	Evidence-based drugs	Treat more people	Less side-effects	Larger spectrum of informed choice
New improved consent handling	Transparent, data sharing	Power of decision in the hands of the individual	Indirectly benefit future patients	More flexibility to choices made