



*The use of AI for health  
outcomes predictalytics as  
a foundation for Value  
Based Healthcare (VBHC)*

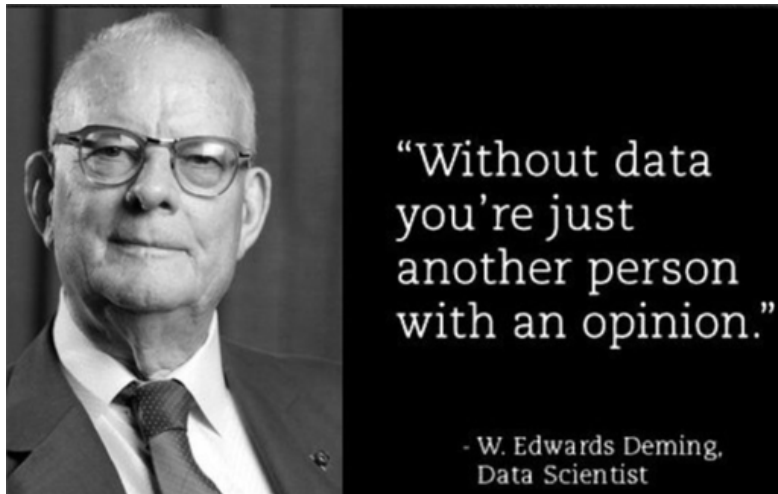
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Senior Radiotherapist

- CEO - Intacare International Ltd
- CEO - The Health Value Partnership



## Scene setting



### Different perspectives on 'benefit and 'value'

The benefit of any medical innovation can only truly be measured by the benefit to the patient, as they see it!

Dr Don Berwick – Advisor, World Health Organisation

When assessing the value of any intervention we must not look solely at whether the patient lived as a result but rather whether they lived (or died) 'well'.

Public Health England

At the beginning of any decision process one must always consider the perspectives of the different stakeholders, and then balance where the value lies for each.

P. Just: PharmD, Senior Principal, Global Health Economics. ICON

- 
- **Payers:** €, QALY, DALY, ICER
  - **Hospitals:** Admissions, length of stay, volume of interventions, AEs
  - **Clinicians:** OS, DFS, PSF,
  - **Patients:** Quality of life, functional ability, emotional well-being, time

**Limited Understanding:** About how decisions are made— what data?

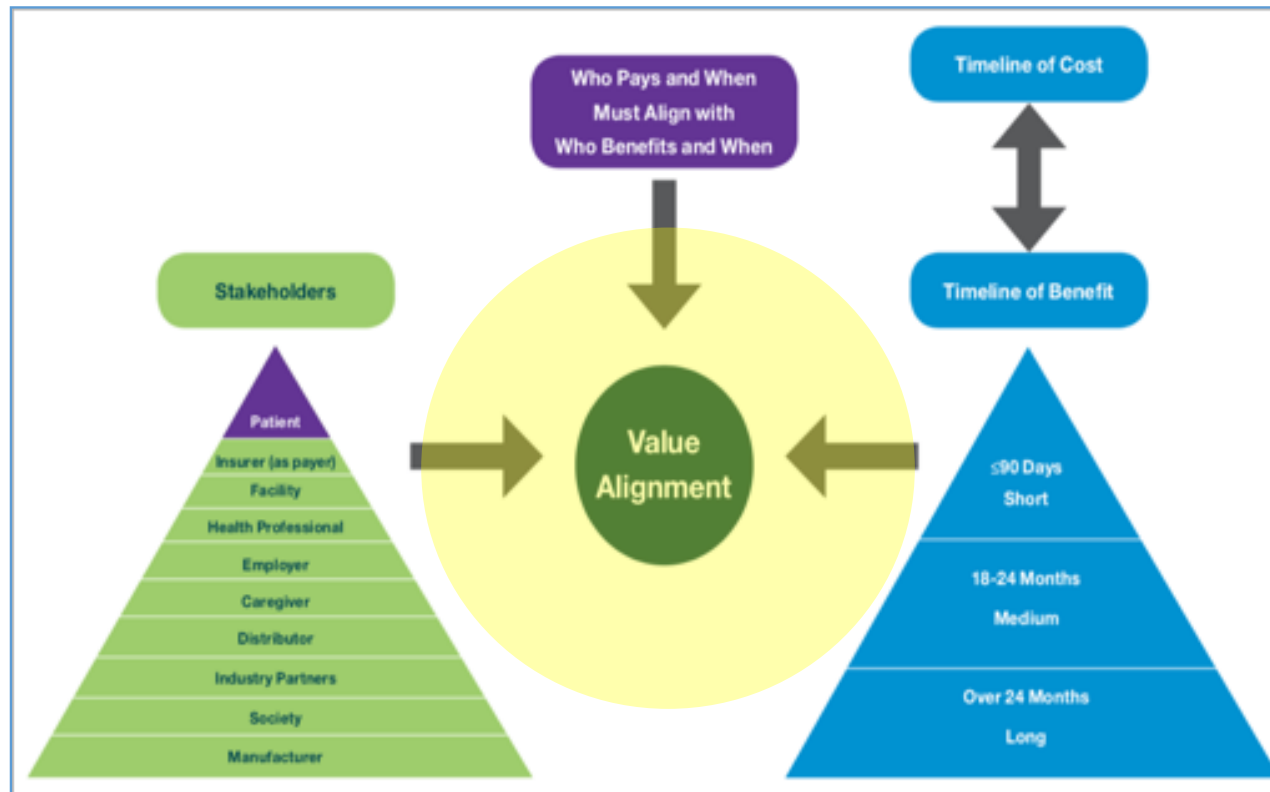
**The result:** Instability and poor 'value exchange', a cycle of diminishing returns

# The Data Challenge

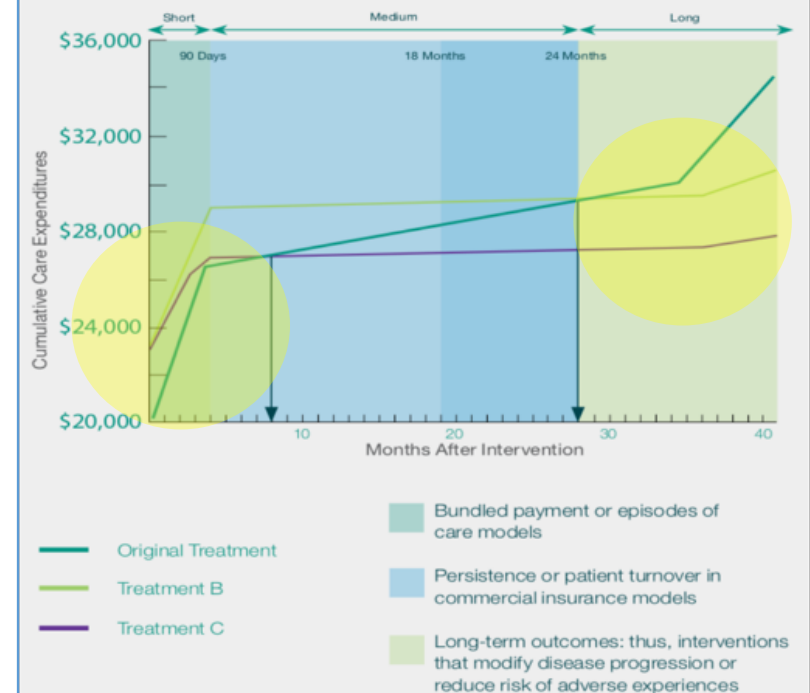
<b>Core clinical parameters</b>	
<ul style="list-style-type: none"> <li>Demographics</li> <li>Co-morbidities</li> <li>Diagnosis dataset – time, methodology, accuracy</li> <li>Regime(s) / protocol(s) incl. Trial e.g. FEC, CHHiP</li> <li>Treatment Dataset - Sx, CT, RT, HRT, IT etc</li> <li>Acute and longitudinal toxicity profiles (RTOG / CTCAE)</li> <li>Adverse event reporting</li> </ul>	<ul style="list-style-type: none"> <li>Personal History</li> <li>Presenting symptoms profile</li> <li>Treatment Dataset - Sx, CT, RT, HRT, IT etc</li> <li>Dose, timing, duration, OAR dose, multi-modalities</li> <li>Regimen(s) / protocol(s) incl. Trial e.g. FEC, CHHiP</li> <li>Survivorship data: OS, PSF, DFS , date of death etc</li> <li>Longitudinal health data</li> </ul>
<b>Clinical Coding</b>	<b>Ontologies</b>
<ul style="list-style-type: none"> <li>ICD-10</li> </ul>	<ul style="list-style-type: none"> <li>SNOMED-CT</li> </ul>
<b>Toxicity Scoring</b>	<b>Clinical Data Systems</b>
<ul style="list-style-type: none"> <li>RTOG, UKONS, CTCAE</li> </ul>	<ul style="list-style-type: none"> <li>EMR, OIS + AE reporting</li> </ul>
<b>Patient Reported Outcomes</b>	<b>Patient Reported Experience</b>
<ul style="list-style-type: none"> <li>380 different questionnaire suites</li> </ul>	<ul style="list-style-type: none"> <li>Numerous</li> </ul>
<b>Claims / Cost datasets / models</b>	<b>Claims Systems</b>
<ul style="list-style-type: none"> <li>UK CCSD, ICS, HRG, QALY, ICER, DALY, QOCRv</li> </ul>	<ul style="list-style-type: none"> <li>CREST, CIARA, Lorica</li> </ul>
<b>Benchmarks - national</b>	<b>Benchmarks - International</b>
<ul style="list-style-type: none"> <li>PHE (COSD, RTDS, PCDS, NatPatSatQ) + ICHOMs, CODE, SAMS, PIE, others</li> </ul>	<ul style="list-style-type: none"> <li>OECD – others</li> </ul>

**In the above, how well does patients voice REALLY feature?**

# The Time Challenge



**Figure 1: Time Dependency of Value Return**



Compared to the standard treatment, both alternatives (treatment B and C) are more costly in the short-term. But, when evaluated as total treatment cost over time, treatment B is cost saving after eight months - but cost more in total. Treatment C is cost saving beyond two years and gives most return in the long term.

# Introducing Intacare



## The company:

- A clinical AI and data science company dedicated to developing clinical risk prediction solutions that optimise outcomes for multiple stakeholders
- **The patients voice remains central to everything we do**

## Aim:

To socialise the science of predictalytics in VBHC

## Vision:

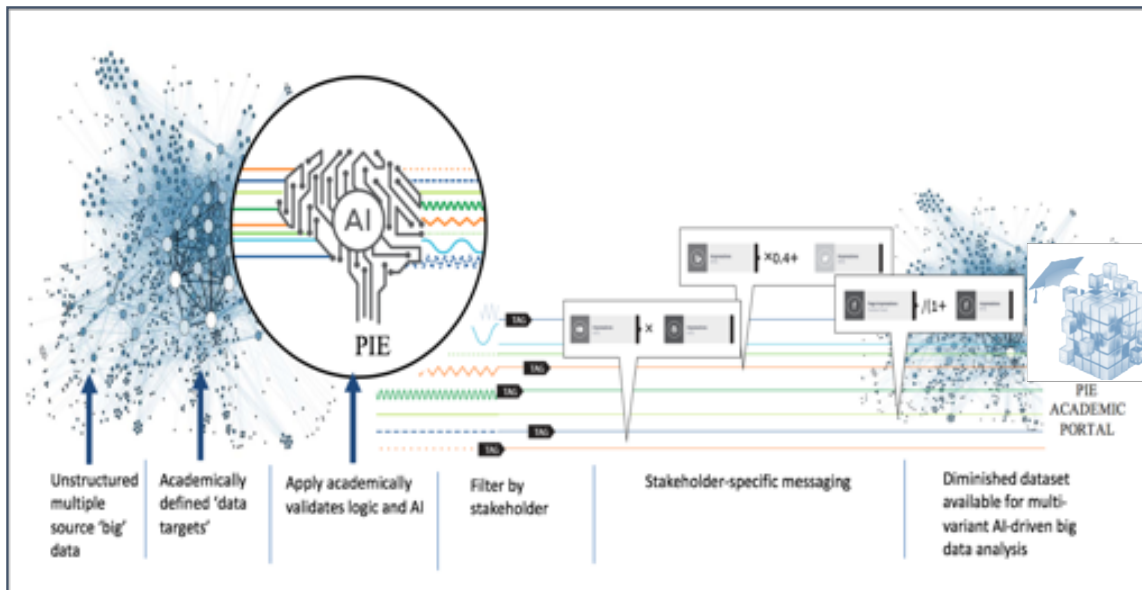
- To be the 'google maps' for clinical outcomes – enabling better value navigation by providing a 'value map' that transcends the healthcare ecosystem

## History:

- Seven years of academic research (Imperial College London, University of Surrey)
- Three clinical pilots (Royal Marsden, Royal Surrey, Southampton NHS Trusts – partnered with Varian and Macmillan)
- Commercial pilots (GenesisCare UK (10 Sites) + Varian)
- Now linking to the largest insurer in the world and the largest oncology-tech solution in the world



# Intacare AI – Predictive Risk and Outcomes Insights Engine (PIE)



## Liberate academic knowledge

Identify the data that is both necessary and currently available to enable the application of real world insights for complex decision support, risk assessment, tracking and mitigation and cost/benefit identification.

## Target data from multiple datasets

Use AI and machine learning-based system to apply pre-defined academic logic to defined datasets to present risk reports that are tailored to the subscribers needs.

## Federated learning

Provide a secure cloud-based SaaS risk engine to interact with third-party data collection suites and analysed data to elicit dynamic decision aids to put these at the finger-tips of the decision makers

### Patient-Centricity

Simplifying patient reported outcomes and making them diagnostic and dynamic

### Precision

Superior clinical decision support and insights built from the latest evidence

### Prediction

Predicting events, eliciting better value across the ecosystem for value-based reimbursement and performance incentives

### Value

Defined value outputs optimized for Value-Based Healthcare and cost transparency

## Intacare AI – Academic Programme 1: Cancer Dynamic (diagnostic and predictive) PROMs



**Question: What is the most comprehensive holistic PROM that correlates with late effects**

**Identify:** Most comprehensive symptoms toolkit that covers key holistic domains:  
Symptoms, psychosocial, vocational, spiritual / Quality of Life  
(Rotterdam)  
What symptoms are missing

**Process:** Combined 380 PROMs tools.  
Correlated PROM with patient characteristics + clinical data + toxicity data  
Identified what symptoms are most indicative of a clinical event  
Academically and clinically validated = 19

**Develop:** A new, hybrid and all encompassing toolkit with a new, more meaningful presentation structure (Intacare SAMS):

- Correlates with late effects prediction
- Validated for clinical and academic use.

Report	Date
Insurance Reports	15 May 2015
Insurance Reports	20 May 2015

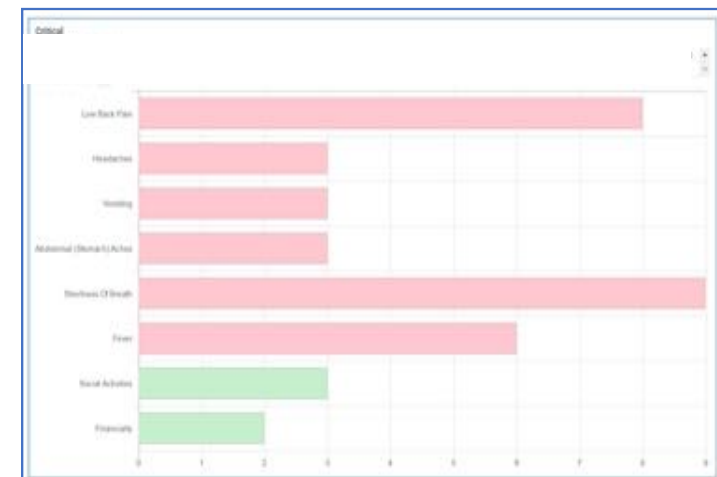
  

Report	Date
Customer Service	15 May 2015

## Intacare AI – Academic Programme 1: Cancer Dynamic (diagnostic and predictive) PROMs

**Presents:**

- Specific symptom severity – Heatmaps
- +/- trends
- Cancer-specific symptom clusters , breaches
- Identifies early sign of recurrence
- Identifies risk and early signs of critical events





# Intacare AI – Academic Programme 2:

## Cancer late effects prediction models (predictalytics)



**Inception:** What are the data that are associated with a late effect risk / presentation. How to target these data to predict future events

**Assess:** Academic literature (100's of evidence level I and II papers) relating to toxicity and late effects

**Analyses:** Disparate datasets and targets defined data (precision data)

**Filters:** Data into international ontologies and coding structures (ICD etc)

**Correlates:** Patient characteristics + clinical data + utility data + toxicity data + Patient Reported Data (PROMs) + Patient Satisfaction Data (PREMs)

**Identify:** The symptoms assessed in the report, importance use  
Patient characteristics + clinical data + toxicity data  
Determine what symptoms are most indicative of a clinical event

**Develop:** A new, dynamic, late effects prediction and state change assessment model

A screenshot of a clinical report titled "Syndrome, Causation, Probability Analysis Reports". The report is structured as a table with two columns: "Item" and "Ref".

Item	Ref
Adverse Events	11/10/2018
Adverse Events	21/04/2019
Holistic Quality of Life Questionnaire Alerts	
Ref	Ref
Customer data	11/10/2018

# Intacare AI – Academic Programme 2: Cancer late effects prediction models (predictalytics)

**Risk management logic** Validate Submit Cancel

Treatment mode	Acute effects	Late effects	Monitoring	PROMS	Remove?
Radiology	<input type="checkbox"/> Bowel dysfunction - diarrhoea	<input type="checkbox"/> Rectal bleeding	<input checked="" type="checkbox"/> 6-monthly PSA	<input type="checkbox"/> Urinary symptoms	<a href="#">Remove</a>
	<input type="checkbox"/> Bowel dysfunction - tenesmus	<input type="checkbox"/> Irregular bowel habit	<input checked="" type="checkbox"/> 12-monthly consult	<input type="checkbox"/> Nocturia	
	<input type="checkbox"/> Dysuria	<input type="checkbox"/> Stricture of the urethra	<input type="checkbox"/> Advise patient to report new symptoms	<input type="checkbox"/> Fatigue	
	<input type="checkbox"/> Rectal bleeding	<input checked="" type="checkbox"/> Erectile dysfunction		<input type="checkbox"/> Weakness	
	<input type="checkbox"/> Urinary incontinence	<input checked="" type="checkbox"/> Haematuria		<input type="checkbox"/> Breathlessness	
	<input checked="" type="checkbox"/> Urinary frequency	<input checked="" type="checkbox"/> Secondary malignancies		<input type="checkbox"/> General disability	
	<input type="checkbox"/> Haematuria			<input type="checkbox"/> Skeletal pain	
	<input type="checkbox"/> Pelvic pain			<input type="checkbox"/> Instability, change in mood, or confusion	
				<input type="checkbox"/> Depression	
				<input type="checkbox"/> Bowel dysfunction - rectal bleeding	
Surgery	<input type="checkbox"/> Bowel dysfunction - diarrhoea	<input type="checkbox"/> Rectal bleeding	<input checked="" type="checkbox"/> 6-monthly PSA	<input type="checkbox"/> Urinary symptoms	<a href="#">Remove</a>
	<input type="checkbox"/> Bowel dysfunction - tenesmus	<input type="checkbox"/> Irregular bowel habit	<input type="checkbox"/> 12-monthly consult	<input type="checkbox"/> Nocturia	
	<input checked="" type="checkbox"/> Dysuria	<input type="checkbox"/> Stricture of the urethra	<input type="checkbox"/> Advise patient to report new symptoms	<input type="checkbox"/> Fatigue	
	<input type="checkbox"/> Rectal bleeding	<input checked="" type="checkbox"/> Erectile dysfunction		<input type="checkbox"/> Weakness	
	<input type="checkbox"/> Urinary incontinence	<input checked="" type="checkbox"/> Haematuria		<input type="checkbox"/> Breathlessness	
	<input type="checkbox"/> Urinary frequency	<input checked="" type="checkbox"/> Secondary malignancies		<input type="checkbox"/> General disability	
	<input type="checkbox"/> Haematuria			<input type="checkbox"/> Skeletal pain	
	<input type="checkbox"/> Pelvic pain			<input type="checkbox"/> Instability, change in mood, or confusion	
				<input type="checkbox"/> Depression	
				<input type="checkbox"/> Bowel dysfunction - rectal bleeding	

## Presents:

- The probability of late-effect (syndrome) e.g. cardiac, respiratory syndrome
- Identifies the associated / contributing risk factors
- Predicts the % probability of and time to occurrence
- Predicts the presentation e.g. myocardial infraction
- Proposes a monitoring plan e.g. monthly BP + weight +, 6-monthly MUGA LVEF
- Provides a link to the supporting evidence that the rule(s) was derived from

Syndrome Causation, Probability Analysis Reports

[Effects](#) [Syndromes](#) [Monitor](#)

**Syndromes Message:**

- The Insight Engine has detected a risk for cardiac syndrome.
- The Insight Engine has detected a risk for neuro-cognitive syndrome.
- The Insight Engine has not detected risk for musculoskeletal syndrome.
- The Insight Engine has not detected risk for metabolic syndrome.
- The Insight Engine has not detected risk for respiratory syndrome.
- The Insight Engine has not detected risk for intra-pelvic syndrome.
- The Insight Engine has not detected risk for gastroenteric syndrome.

[Effects](#) [Syndromes](#) [Monitor](#)

Thank you, your inputs have been received and triggered 19 patient insights.

88.0% risk of effect cardiac abnormality, asymptomatic changes related to the cardiomyopathy system according to rule 59 from the cardio-metabolic survey report

88.0% risk of symptom pre-existence (abnormalities of the cardiac muscle related to the cardiac toxicity system according to rule 108 from the cardio-metabolic survey report

There is a 3.0% probability of developing the effect abnormality of breast related to the cardiomyopathy system according to rule 303 from the musculoskeletal survey report

26.0% risk of symptom pain in the upper body related to the localized pleuropathy system during the next 3 to 12 months according to rule 323 from the neuro-cognitive survey report

12.0% risk of effect related to the peripheral neuropathy system during the next 60 months according to rule 330 from the neuro-cognitive survey report

11.0% to 30.0% risk of effect cardiac abnormality, cardiac disease related to the cardiomyopathy system in 60 to 120 months according to rule 57 from the cardio-metabolic survey report

11.0% to 30.0% risk of effect related to the cardiac toxicity system during the next 60 to 120 months according to rule 103 from the cardio-metabolic survey report

present risk of developing the effect pericarditis related to the cardiac toxicity system according to rule 64 from the cardio-metabolic survey report

present risk of developing the effect thoracic dysfunction related to the cardiac toxicity system according to rule 69 from the cardio-metabolic survey report

Syndrome Causation, Probability Analysis Reports

[Effects](#) [Syndromes](#) [Monitor](#)

**Monitor Message:**

Monitoring for cardiac syndrome may include items 3 tests

Monitoring for neuro-cognitive syndrome may include items 47 tests

# Case Study: Research Programmes

## Simplifying Long-form PROMs & Predictalytics



- Royal Marsden NHS Foundation Trust, UK:** - **6-yr Prostate Brachytherapy multiple PROMs analysis (complete)**  
**Royal Surrey NHS Foundation Trust, UK:** - **5-years Colorectal Cancer multi-PROMs analysis (July 2019)**  
- **5-years Prostate multi therapy data analysis (July 2019)**

### Aims:

- Assess the correlation between treatment characteristics (dose etc), patient characteristics (age), patient's own perception of health
- Patterns in symptoms presentation against time, the individual and their treatment characteristics
- Identify symptoms that are typical in this cohort and indicate toxicity and declining Quality of Life
- Map to Predictive Insights Engine for late effects correlation

### Royal Marsden Review:

<b>Final Cohort analysed:</b>	94 men who had been treatment with prostate brachytherapy between 2006 – 2013 (mean age: 64-yrs)
<b>PROMs data:</b>	EPIC, FACT-P(v4), IPSS, Fulham questionnaires (full or partial) at baseline, 6-months, 1, 5, 6 years
<b>Treatment system data:</b>	Age at treatment, Total Target Dose D100 (Gy), Target dose D90 (Gy), Rectal Volume (cm3), Rectum dose D100 (Gy), Urethra volume (cm3), Urethra dose D100 (Gy), Number of needles, Number of seeds
<b>Key Missing data:</b>	Patient stage and grade

## Analysis:

1. PROM response timeline
2. Features importance
3. Cluster longitudinal analysis of correlating factors and response
4. Effect on other parameters

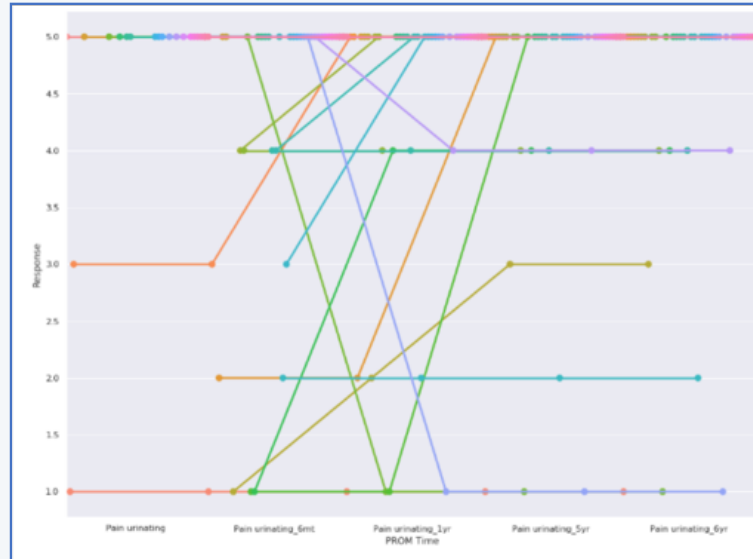
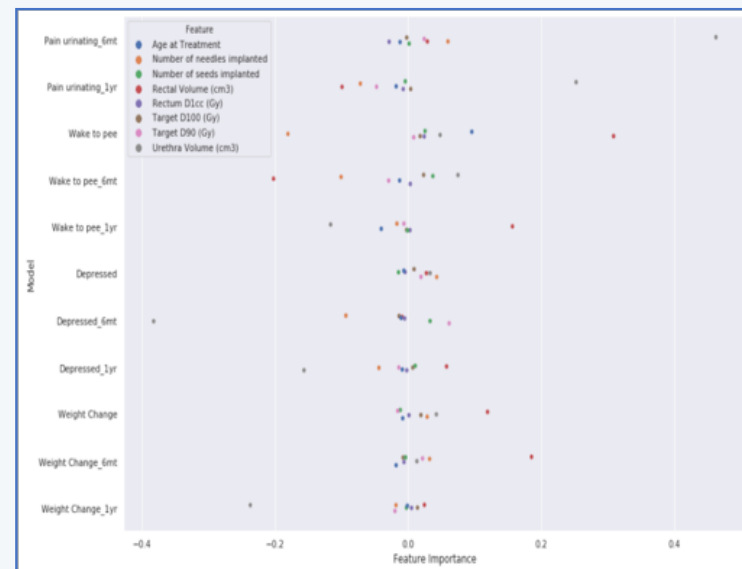
Symptoms	Correlation
↑ depression at 1 year Persistent depression	Number of needles, rectal vol, urethra vol, pain
Potency decline	Age, number of needles, urethra vol
Weight change at 6 months, 1 year	Age, potency decline, pain, depression
Pain (general)	Rectal vol, depression
Pain urinating	Urethra vol, depression
Nocturia at 1-year	Number of needles, urethra vol, depression

**Predictive modelling** - How will patients will answer PROM at each timepoint and what features are important

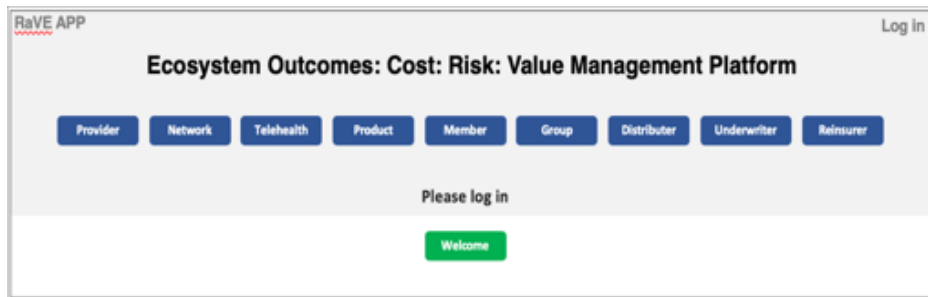
(Support Vector Regression with mean squared error (low = well performing))

## Future Analysis:

1. Stage and Grade correlations
2. Analyse subset of patients who feel their pain gets worse over time
3. Run targeted PROM against new PROM Protocol using EORTC-QOL PR25 to further develop predictive model



# Intacare AI – Dynamic Risk and Value Engine (RaVE)



## Aim:

- Provide a multi-sided digital QOCRv insights platform
- Improve consumer and provider risk profiling
- A secure medium of exchange for healthcare outcomes and value-based payment communication
- A machine learning platform for identifying unwarranted variation, fraud, waste and abuse

**Analyses:** Disparate claims / cost datasets

**Filters:** Data into standardised coding - coding (ICD / HRG / CCSD...)

**Correlates:** Patient characteristics + clinical data + utility data + toxicity data + Patient Reported Data (PROMs) + Patient Satisfaction Data (PREMs) + cost / claims (by code)

## Produces:

- Bespoke targeted 'risk insights' to our subscribers
- Using standard industry health-economic measures e.g. Quality Adjusted Life Years, Daly Adjusted Life Years, ICER, Relative Live Value, Relative Risk, Utility and utility cost.

# Case Study: Cost of Cancer Analysis



**PMI providers (1 x New Zealand, 2 x UK): Typical Spend GBP £300m - 400m, NZD \$17m**

## **Aims:**

1. To determine the 'total cost of cancer', root cause of costs, establish mechanisms to assess, predict and mitigate cost risk
2. Establish Outcomes Based Commissioning Model for Targeted Commissioning from high performers

## **Process**

1. Assess historic claims data (5-ys) – what are patients/ hospitals/ clinicians claiming for
2. How do these claims relate to typical toxicity / co-morbidity profiles
3. What are the characteristics of individuals – treatment-related / patient-specific - correlate with PIE
4. Where were they treated, by whom and with what (appropriateness)
5. Run some value models (QALYs, ICERs, DALYs, TDABC, total cost of claim/ life, time, clustering's, Relative Life Value/Risk

## **Results:**

- 1795 code narrative variations (standard CCSD model = 33)
- Wide variation in claims – areas of significant suspected fraud, waste, abuse and unwarranted variation
- £44-56m in savings (UK): 14% savings (excl. operational efficiencies)

## **Next steps:**

- New 'quality and value assurance' model mandated to all hospitals - includes PROMs and PIE dataset.



# Summary

- **The patients voice is diagnostic and predictive**
- **Captured via PROMs / PREMS and toxicity datasets**
- **Align our perspectives on 'benefit and 'value' – ecosystem value exchange**
- **Standardize our datasets – ensure PROMs / PREMs etc are central**
- **Include correlation between clinical, patient reported and cost/ utility data (transcending views)**
- **Identify root cause of value erosion – evidence based best practice?**
- **Evolve the models for predictalytics (proactive not reactive care) – let AI do the heavy lifting**
- **Socialised e.g. via ICHOMs**
- **Observe, learn, optimise, repeat...**



**Thank you**

**Matt Hickey**

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[www.intacare.com](http://www.intacare.com)