Trustworthy AI and Zinpection@IRyCIS

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About me

- Prof. in Computer Science, University of Alcalá
- Since 2022, coordinator of the Data Science Unit of the Research Institute at the "Ramón y Cajal" Hospital in Madrid.
- Co-founder In2AI
- Established the Z-Inspection Lab@IRyCIS early 2024.



• Universidad







Initial focus @ IRyCIS of using Z-inspection

- Given that AI is considered a crosscutting priority in the organization, the focus is on planning in advance.
- Trustworthy-first design of clinical decision making interfaces.
 - Preparing for transition to actual practice.
 - Avoiding late findings of pitfalls or barriers.
- Current emphasis on key enablers of AI based projects:
 - Adapting socio-technical scenario modeling.
 - Collecting best practices on traceability, especially and starting from research data.



Large case study: AI and pancreatic cancer

- Focus: risk, early detection and diagnostic tools for PC
- Z inspection considered as the basis for the case across institutes in Spain first, then in EATRIS overall.
- Digital capabilities with emphasis on predictive AI.
 - Includes Trustworthy AI as key element.
 - Materials with best practice in a Digital HUB







DEATHS ESTIMATED IN EUROPE 2022- 2045 IN EUROPE



Current treatments



Risk Factors



AVAILABLE INFORMATION AT HOSPITALS

Electronic Health Records (EHR) (standards as <u>enablers</u> for accountability)



а KRAS CDKN2A SMAD TD5 * KDM6A MLL3 MLL2 * TGFBR2 SV subtypes BRCA1 BRCA2 * Histopathology Moffitt turnour class Moffitt stromal class Collisson class ADEX Pancreatic progenitor Souamous Immunogenia -2 Scattered Focal Stable Unstable Basal-Ike Classical Activated stroma Normal stroma Adenosouamous carcino Non-slient SNV or indel Quasi-mesenchymal Acinar cell carcinoma SV Exocrine-like PMN with invasion/mucinous Deletion Classical Amplification (copy number >8) BRCA2 germline mutations Liquid biopsy EVs CON CIDNA

and made

MOOR

Point mutatio

Genomic

Copy Number Variation

M

Molecular Markers



Imaging: CT scan, MRI



Histopathology



IRYCIS/ES TASK FORCE – multidisciplinary team

> M Laura García Bermejo, WP6 coordinator, Biomarkers & Therapeutic Targets Leader

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- Miguel Ángel Sicilia, Data Science Unit Director
- > Elena García Barriocanal, **Data Science Unit member**
- > Ana García-García de Paredes, Gastroenterologist (pancreatic diseases section leader)
- Mercedes Rodriguez Garrote, Medical Oncologist (GI tumor section)
- > Carolina de la Pinta, Radiation Oncologist (GI tumor section) /Radiomics
- Elisa Conde, Biomarkers & Therapeutic Targets Unit
- Julie Earl, Pancreas Cancer Patient registry responsible at IRYCIS, BIOPAC Group
- > Val Fernández, Bioinformatics (Trans-BIONET, ELIXIR)
- > Enrique Aracil, Hospital Deputy Director, Responsible of EHR at Ramon y Cajal
- > Sonia Camaño, HRYC Biobank coordinator, (BBMRI), Responsible IRYCIS Quality
- > Itziar de Pablo, Ethical Committee of Clinical Investigation (SCREEN/ ECRIN), Al expert
- Richard Buck, Patient representative
- Fuensanta Bellvis, Quibim Development Support

<u>eatris</u>onnect **Clinical variables Definition and sources: initial Proposal**

	Structured?	Where to find it?	Surgical histo
Patient characteristics			gastrectomy
sex	Yes	HCIS	D, aspirin, N
age at diagnosis	No	HCIS-	complication
		Gastroenterology	fasting blood
race	No	HCIS	glycosylated
smoker	No	HCIS	diagnosis
alcohol	No	HCIS	weight loss i
cannabis	No	HCIS	abdominal p
exposure to chemical compounds	No	HCIS	meteorism i
Obesity: BMI <30	No	HCIS	diarrhea in y
blood group	No	HCIS	nutritional s
sedentary lifestyle	No	HCIS	exocrine par
history of helicobacter pylori infection	No	HCIS-	Tumor chara
		Gastroenterology	ECOG
history of chronic HBV	No	HCIS-	diagnosis: in
		Gastroenterology	date of diag
history of chronic HCV infection	No	HCIS-	stage
		Gastroenterology	histology
history of other tumor	No	HOIS	grade
DM 1 or 2	No	HCIS	location
date of DM diagnosis	No		size
history of acute paperoatitic	No		Presence of
history of acute pancreatitis	No		metastases:
	No		PET/CT at di
type of pancreatitis	NO	HUIS-	
familiar bistory of sansar	No		liver MRI at
familiar history of cancer	NO	HCIS-Oncology	
familiar pancreatic cancer	NO	HCIS-Oncology	exploratory
nereditary syndromes: familial breast and	NO	HCIS-Oncology	
ovarian cancer (BRCA1, BRCA2, PALB2), Peutz-			resectability
Jegners synarome (STK11), Lynch synarome			committee
(CDKN2A) stavia telengiastasia sundrama			type of tumo
(CDKNZA), ataxia-telanglectasia synurome			unresectable
(ATIVI)	Ne	LICIC	
precursor lesion: cystic mucinous neoplasia,	NO	HUIS	Ca 19.9 at di
Intraductal papillary mucinous neoplasia and			fasting blood
high visit stigments	Ne		glycosylated
nign-risk sugmata	NO	Costroontorology	CEA at diagn
	Ne	Gastroenterology	hemoglobin
worrisome teatures	NO	HUIS-	PCR at diagn
		Gastroenterology	Total Bilirub
Date of diagnosis of the precursor lesion	NO	HCIS	alkaline pho
Suspicious findings according to Fukuoka	No	HCIS	GGT at diagr
criteria.			GOT at diagr

Surgical history of cholecystectomy or	No	HCIS
gastrectomy, treatment with statins, vitamin		
D, aspirin, NSAIDs, other diseases or		
complications		
fasting blood glucose in year prior to diagnosis	Yes	Laboratory- HCIS
glycosylated hemoglobin in year prior to	Yes	Laboratory- HCIS
diagnosis		
weight loss in year prior to diagnosis	No	HCIS
abdominal pain in year prior to diagnosis	No	HCIS
meteorism in year prior to diagnosis	No	HCIS
diarrhea in year prior to diagnosis	No	HCIS
nutritional status at diagnosis	No	HCIS
exocrine pancreatic insufficiency at diagnosis	No	HCIS
Tumor characteristics		
ECOG	No	HCIS
diagnosis: incidental, symptomatic	No	HCIS
date of diagnosis	No	HCIS
stage	No	HCIS
histology	No	GESPAD reports
grade	No	GESPAD reports
location	No	HCIS
size	No	HCIS
Presence of metastases at diagnosis, type of	No	Radiology reports-
metastases: hepatic, pulmonary, bone		HCIS
PET/CT at diagnosis	No	Nuclear medicine
		reports- HCIS
liver MRI at diagnosis	No	Radiology reports -
		HCIS
exploratory laparotomy at diagnosis	No	HCIS- General
		surgery
resectability assessed by a multidisciplinary	No	HCIS -
committee		Multidisciplinary
		committee reports
type of tumor (resectable, bordeline,	No	HCIS -
unresectable)		Multidisciplinary
		committee reports
Ca 19.9 at diagnosis	Yes	Laboratory- HCIS
fasting blood glucose at diagnosis	Yes	Laboratory- HCIS
glycosylated hemoglobin at diagnosis	Yes	Laboratory- HCIS
CEA at diagnosis	Yes	Laboratory- HCIS
hemoglobin at diagnosis	Yes	Laboratory- HCIS
PCR at diagnosis	Yes	Laboratory- HCIS
Total Bilirubin at diagnosis	Yes	Laboratory- HCIS
alkaline phosphatase at diagnosis	Yes	Laboratory- HCIS
GGT at diagnosis	Yes	Laboratory- HCIS
GOT at diagnosis	Yes	Laboratory- HCIS

GPT at diagnosis	Yes	Laboratory- HCIS
Treatment characteristics		
endoscopic biliary drainage		
type of surgery	Yes	HCIS- General
		surgery
pTNM	Yes	Pathology reports
date of surgery	Yes	Pathology reports –
		General surgery
neoadjuvant chemotherapy and type	No	HCIS -Medical
		oncology
date of start of chemotherapy	Yes	HCIS -Medical
		oncology- Pharmacy
number of cycles	Yes	HCIS -Medical
		oncology- Pharmacy
response to treatment according to RECIST	No	Radiology reports
response to treatment according to Ca 19.9		
ypTNM in surgical specimen	Yes	Pathology reports
adjuvant chemotherapy	No	HCIS -Medical
		oncology
radiation therapy (neoadjuvant, radical,	No	HCIS -Radiation
adjuvant)		Oncology
type of radiotherapy (SBRT or conventional	No	HCIS -Radiation
radiotherapy)		Oncology
doses of radiotherapy	No	HCIS -Radiation
		Oncology
unresectable or metastatic patient	No	Radiology reports
local progression	No	Radiology reports
distant progression	No	Radiology reports
first line of chemotherapy	Yes	HCIS -Medical
		oncology- Pharmacy
second line of chemotherapy	Yes	HCIS -Medical
		oncology
status	No	HCIS
Exploratory variables related to analytical		
parameters, other medical history and		
treatments prior to diagnosis will be included		
Radiomics: GLSZM, NGLDM, energy, entropy,	NA	DICOM

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Traceability – from DMPs to Trustworthy Al

- Translational medicine:
 - Often described as an effort to carry scientific knowledge "from bench to bedside," builds on basic research advances and uses them to develop new therapies or medical procedures.
- Research planned in advance and data processes and products documented first in Data Management Plans (DMPs)
- This sets the need to advance the data infrastructure and data engineering practices to trace from DMPs to all downstream data artifacts.

Processes and trustworthy Al



CRISes, DMPs and Data Lakes

• The Data Management Plan (DMP) is the specification that mediates between the Project and the artifacts being generated in the Data Lake.



Bridging the DMP, the metadata catalogue and actual data

