



European Network
of Cancer Registries



European
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Joint Research Centre



ENCR Workshop on Software and AI tools for cancer registries

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Registro de Cáncer de Murcia

VALIDATION OF DIGITAL SOURCES OF INFORMATION AND DECISIONAL ALGORITHMS IN CANCER REGISTRATION



Región de Murcia
Consejería de Salud

Dirección General de Salud
Pública y Adicciones

Dra. María Dolores Chirlaque.
Murcia Cancer Registry Director.
Preventive Medicine Specialist. PhD.



Cofinanciado por
la Unión Europea



Instituto Murciano de
Investigación Biosanitaria
Pascual Parrilla

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Epidemiología y Salud Pública

UNIVERSIDAD DE
MURCIA



VALIDATION OF DIGITAL SOURCES OF INFORMATION AND DECISIONAL ALGORITHMS IN CANCER REGISTRATION



Mª Dolores Chirlique



Mónica Ballesta



Ricardo Vaamonde



Mª José Sánchez



Mercedes Expósito



Antonia Sánchez



Sandra Garrido



Juan Francisco Molina

Manuel Lodeiro

Milagros Montesinos

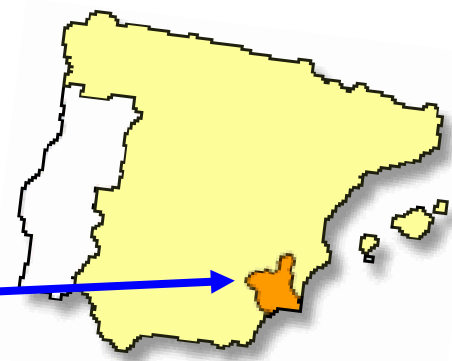
VALIDATION OF DIGITAL SOURCES OF INFORMATION AND DECISIONAL ALGORITHMS IN CANCER REGISTRATION

- Population Murcia Region
2023: 1 556 568 inhabitants.

[WWW.INE.ES](http://www.ine.es)

- Number of incident cancer
cases in 2022-2023: 14599.

<https://sms.carm.es/ricsmur/bitstream/handle/123456789/13104/bem.2024.44.883.pdf?sequence=1&isAllowed=y>



▶ I. SOURCES OF INFORMATION

▶ II. VALIDATION OF DIGITAL SOURCES OF INFORMATION

▶ III. AUTOMATED PROCESS OF DATA LOAD IN THE OFICIAL HEALTH CARE DATABASE

Fist step: normalization

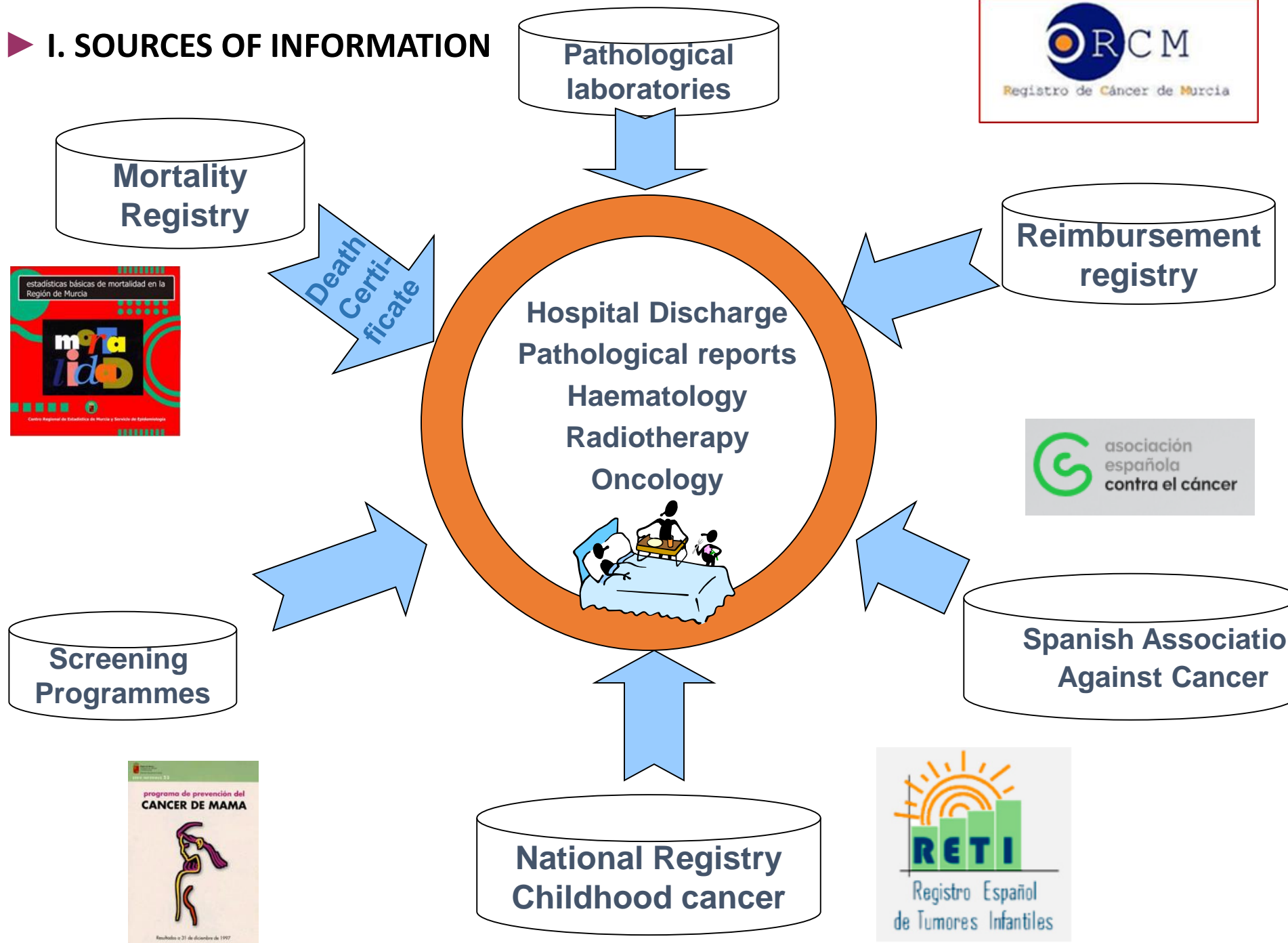
Second step: codification

Third step: linkage with person in oficial Health Care Database

▶ IV. AUTOMATED PROCESS OF CANCER CONSOLIDATION IN CANCER REGISTRY PROGRAM

Decisional algorithms

I. SOURCES OF INFORMATION



FUENTES DE INFORMACIÓN HOSPITALARIAS	
HOSPITAL	Servicio
H. Virgen de la Arrixaca	Anatomía Patológica. CMBD. Oncología,Radioterapia,Hematología
H. R. Sofía	Anatomía Patológica. CMBD
H. Virgen del Rosell	Anatomía Patológica. CMBD. Hematología.
H. Morales Meseguer	Anatomía Patológica. CMBD. Onco-Hematología
H. R.Méndez	Anatomía Patológica. CMBD.
H. Noroeste	Anatomía Patológica CMBD.
Hospital Altiplano	Anatomía Patológica CMBD. Hematología.
H. los Arcos	Anatomía Patológica. CMBD.
H. de Cieza	Anatomía Patológica. CMBD
H. de Molina	CMBD
H. la Vega	CMBD
H.San Carlos	CMBD
H.PerpetuoS	CMBD
H. Caridad	CMBD
H. Cehegín	CMBD
H. Mesa C.	CMBD
H. Alcantar.	CMBD
H.V. Alcázar	CMBD
H. Cli. Bernal	CMBD
R.Villademar	CMBD
R. Alarcón	CMBD



I. SOURCES OF INFORMATION

FUENTES DE INFORMACIÓN NO HOSPITALARIAS
Laboratorio Dr. Sola- Anatomía Patológica
Laboratorio Dra. Arcas- Anatomía Patológica
Laboratorio Dr. Remezal- Anatomía Patológica
Laboratorio Dr. Pérez Guillermo- Anatomía Patológica
Laboratorio Dr. Ortiz- Anatomía Patológica
Reintegro Gastos SMS- Listado reintegro gastos
Registro de SIDA- Casos diagnosticados
Programa Prevención Cáncer Mama- Casos diagnosticados
Programa de Prevención de Cáncer de Colon y Recto- Casos diagnosticados
Certificados de Defunción- Casos diagnosticados

► I. SOURCES OF INFORMATION

1. Available in electronic support?

HOSPITAL DISCHARGE
PATHOLOGICAL REPORTS
DEATH CERTIFICATE



2. Code?

HOSPITAL DISCHARGE ICD-10
PATHOLOGICAL REPORTS SNOMED VS NO CODE
DEATH CERTIFICATE ICD-10



3. Diagnostic codes valid?

STUDY ON VALIDATION OF DIAGNOSTIC CODES:
HOSPITAL DISCHARGE-PATHOLOGICAL REPORTS



▶ I. SOURCES OF INFORMATION

▶ II. VALIDATION OF DIGITAL SOURCES OF INFORMATION

▶ III. AUTOMATED PROCESS OF DATA LOAD IN THE OFICIAL HEALTH CARE DATABASE

Fist step: normalization

Second step: codification

Third step: linkage with person in oficial Health Care Database

▶ IV. AUTOMATED PROCESS OF CANCER CONSOLIDATION IN CANCER REGISTRY PROGRAM

Decisional algorithms

Just to apply decisional algorithms... we need:

VALIDATION OF DIAGNOSTIC CODES FROM HOSPITAL DISCHARGE

- **A. Validation of colorectal cancer diagnostic codes in a hospital administration data set.**
- **B. Validity of Hospital discharge with diagnostic and procedures codes related to female breast cancer in Spain.**
- **C. Is hospital discharge administrative data an appropriate source of information for cancer registries purposes? Some insights from four Spanish registries.**

VALIDATION OF DIAGNOSTIC CODES PATHOLOGICAL REPORTS

- **D. Quality of data on morphology codes from pathology reports in Murcia-Spain.**

- **A. Validation of colorectal cancer diagnostic codes in a hospital administration data set.**

[Validation of colorectal cancer diagnostic codes in a hospital administration data set]

[Article in Spanish]

Mirari Márquez Cid ¹, Isabel Valera Niñirola, María Dolores Chirlaque López, Jacinta Tortosa Martínez, Encarnación Párraga Sánchez, Carmen Navarro Sánchez

Affiliations + expand

PMID: 16942712 DOI: [10.1157/13091140](https://doi.org/10.1157/13091140)

Full text at
GACETA SANITARIA

Objectives: To validate the ability of a hospital administration data set (minimum data set [MDS]) to detect incident cases of colorectal cancer using the Murcia Cancer Registry (MCR) as the gold standard and to measure agreement between the MDS and registration of colorectal cancer.

Conclusions: Because of its high sensitivity, the MDS is a good source for detecting incident cases of cancer. The high agreement found in the site of colorectal cancer helps to consolidate the MDS as a data source for cancer registration.

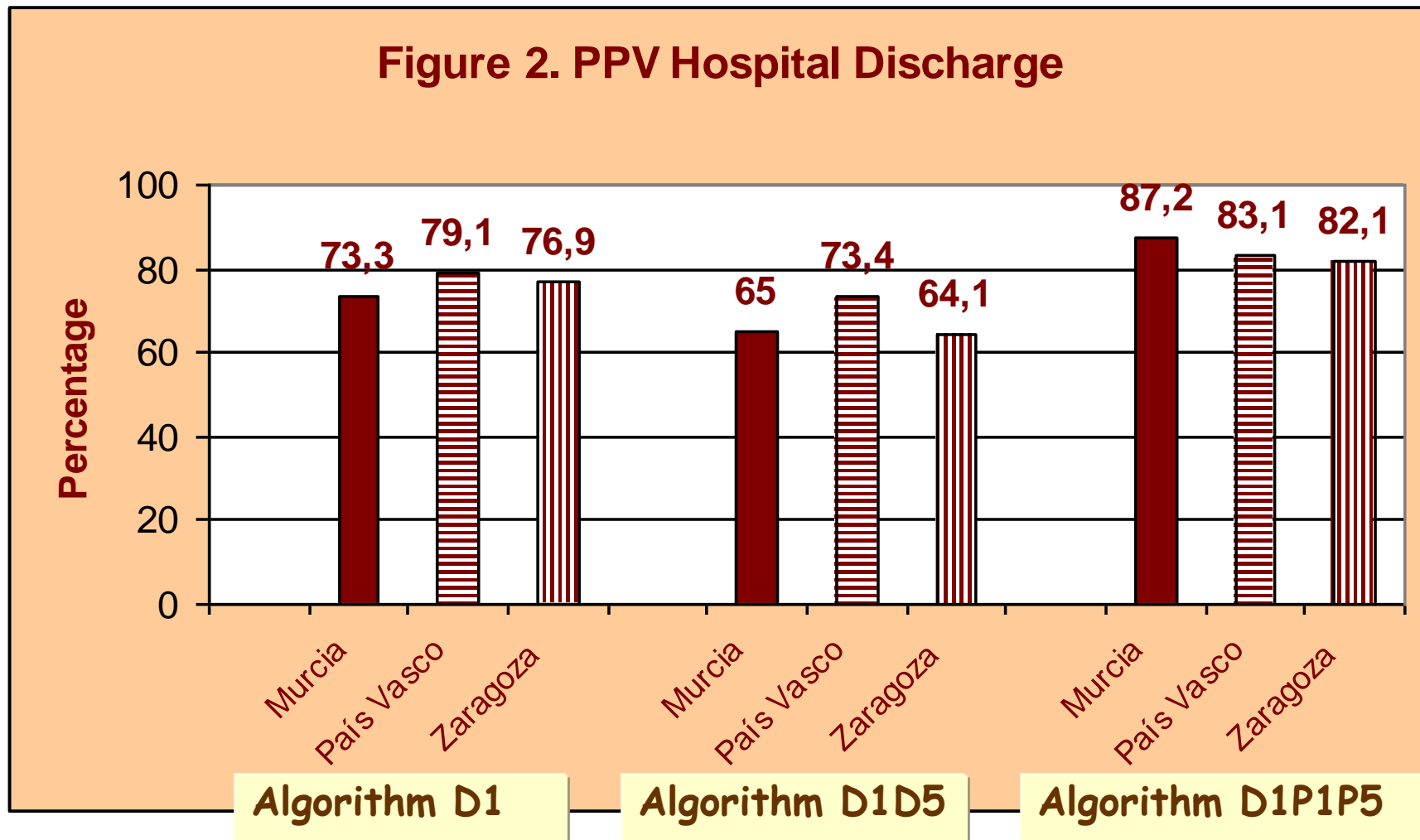
- **B. Validity of Hospital discharge with diagnostic and procedures codes related to female breast cancer in Spain.**

➤ Four regions in Spain with population based cancer registries participate in the study, covering a total of 5 million inhabitants (11% of total Spanish population).

➤ Case definition is episode of hospitalization in women residing in one of the study areas during caused by breast cancer included in the CMBD.



- **B. Validity of Hospital discharge with diagnostic and procedures codes related to female breast cancer in Spain.**



- **B. Validity of Hospital discharge with diagnostic and procedures codes related to female breast cancer in Spain.**

- A high percentage of cases from minimum data set in hospital discharge are confirmed as true positive mainly when the selection of cases is made taking into account diagnoses codes and procedures.
- Hospital discharge records can be used for identifying incident cases of female breast cancer.
- Further analysis will be necessary continuously because of the completeness of CMBD has been increasing in the last years and we would expect to improve these results.

- **C. Is hospital discharge administrative data an appropriate source of information for cancer registries purposes? Some insights from four Spanish registries.**

Bernal-Delgado E *et al.* *BMC Health Services Research* 2010, **10**:9
<http://www.biomedcentral.com/1472-6963/10/9>



RESEARCH ARTICLE

Open Access

Is hospital discharge administrative data an appropriate source of information for cancer registries purposes? Some insights from four Spanish registries

Enrique Bernal-Delgado E^{1*}, Carmen Martos², Natalia Martínez¹, María Dolores Chirlaque^{4,7}, Mirari Márquez^{4,7}, Carmen Navarro^{4,7}, Lauro Hernando⁵, Joaquín Palomar⁵, Isabel Izarzugaza^{3,7}, Nerea Larrañaga³, Olatz Mokoroa³, M Cres Tobalina³, Joseba Bidaurrezaga³, María José Sánchez^{6,7}, Carmen Martínez^{6,7}, Miguel Rodríguez^{6,7}, Esther Pérez^{6,7}, Yoe Ling Chang^{6,7}

- **C. Is hospital discharge administrative data an appropriate source of information for cancer registries purposes? Some insights from four Spanish registries.**

Abstract

Background: The use of hospital discharge administrative data (HDAD) has been recommended for automating, improving, even substituting, population-based cancer registries. The frequency of false positive and false negative cases recommends local validation.

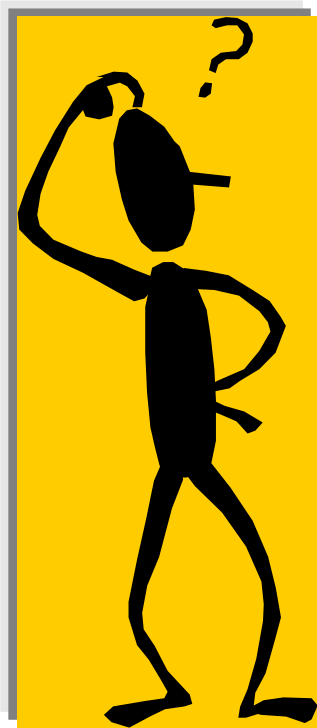
Methods: The aim of this study was to detect newly diagnosed, false positive and false negative cases of cancer from hospital discharge claims, using four Spanish population-based cancer registries as the gold standard. Prostate cancer was used as a case study.

Results: A total of 2286 incident cases of prostate cancer registered in 2000 were used for validation. In the most sensitive algorithm (that using five diagnostic codes), estimates for Sensitivity ranged from 14.5% (CI95% 10.3-19.6) to 45.7% (CI95% 41.4-50.1). In the most predictive algorithm (that using five diagnostic and five surgical codes) Positive Predictive Value estimates ranged from 55.9% (CI95% 42.4-68.8) to 74.3% (CI95% 67.0-80.6). The most frequent reason for false positive cases was the number of prevalent cases inadequately considered as newly diagnosed cancers, ranging from 61.1% to 82.3% of false positive cases. The most frequent reason for false negative cases was related to the number of cases not attended in hospital settings. In this case, figures ranged from 34.4% to 69.7% of false negative cases, in the most predictive algorithm.

Conclusions: HDAD might be a helpful tool for cancer registries to reach their goals. The findings suggest that, for automating cancer registries, algorithms combining diagnoses and procedures are the best option. However, for cancer surveillance purposes, in those cancers like prostate cancer in which care is not only hospital-based, combining inpatient and outpatient information will be required.

- D. Quality of data on morphology codes from pathology reports in Murcia-Spain.

Some insights



To evaluate the agreement on ICD-O-3 topography and morphology codes in real cases of a population based cancer registry doubly coded:

cases coded **manually** (staff) and

cases coded **automatically** (developed software).

Chirlaque MD¹, Márquez M², Navarro C², Bermejo J^{1*}, Rodríguez-Barranco M², Tortosa J², Valera I², Garrido S², Párraga E².

¹Epidemiology Department, Murcia Health Council.
²University Hospital Virgen de la Arrixaca, Murcia.
 mdolores.chirlaque@car.m.es

INTRODUCTION:

The automation of cancer registration allows to obtain a high benefit of the codified information and reduce manual time to complete cancer cases. As a previous step to an automatic process is necessary to know the validity of codified and stored information in magnetic support because the automatic process does not have to decrease the quality of data, quite the opposite, it will have to improve.

OBJETIVE: to measure the agreement in morphology codes of malignant neoplasms between pathological reports and those registered at the population-based cancer registry.

METHODS:

The information of pathological reports come from the Pathological Department of the main hospital in Murcia Region, south-east Spain, whereas incident tumors have been obtained from Murcia Cancer Registry (RCM). Both have been selected from year 1998. The pathological reports included in the study have been those with morphology codes malignant primary site (5th digit /3), in one or several reports but with the same morphology codes. So, reports that show more than one morphology code of different invasive tumor (except 800/3) have been excluded.

We analyse the agreement in SNOMED (Systematized Nomenclature of Medicine) morphology codes at first three digits level that for neoplasms range from 800 to 998.

Agreement has been measured for all morphological groups and by specific morphological groups:

- Epithelial, squamous cell and basal cell neoplasms (801-811)
- Transitional cell papillomas and carcinomas (812-813)
- Adenocarcinomas (814-855)
- Complex epithelial and specialized gonadal neoplasms (856-867)
- Paragangliomas and glomus tumors, nevi and melanomas (868-879)
- Soft tissue tumors and sarcomas (880-958)
- Lymphomas and leukemias (959-998)

by means of the Kappa index and a 95% confidence interval. The statistical software has been Stata V7.0.

		Tabla 1. PATHOLOGICAL REPORTS								Total
		801-811	812-813	814-855	856-867	868-879	880-958	959-998		
RCM	801-811	716	35	34	1	0	1	0	787	
	812-813	0	20	0	0	0	1	0	21	
	814-855	14	0	417	1	1	0	1	434	
	856-867	0	0	0	1	0	0	0	1	
	868-879	0	0	1	0	25	0	0	30	
	880-958	2	0	2	0	0	66	0	70	
	959-998	2	0	1	0	0	0	44	47	
Total	734	55	485	3	30	68	45	1390		

RESULTS:

1405 incident cases have been included in the analysis, but 15 of them present a pathological code of 800 (not include in table 1 and figure 2). These cases will have to be codified always manually in cancer registration.

The percentage in overall agreement between the two sources of information has been 78.7%, Kappa index 0.75 (IC 0.73-0.77).

By morphology groups:

- Epithelial, squamous cell and basal cell neoplasms, which represents 51% of total cases, show the highest agreement, Kappa index 0.87 (CI 0.81-0.92).
- Lymphomas and leukemias, which represents 3% of total cases, offer the lowest agreement with a percentage of 25%, Kappa 0.20 (CI 0.14-0.27).
- Transitional cell papillomas and carcinomas stand out from the rest by disagreement, with a negative Kappa index.

The lack of agreement normally takes place within next morphology groups.

Figure 1. NUMER OF CASES BY MORPHOLOGY GROUPS

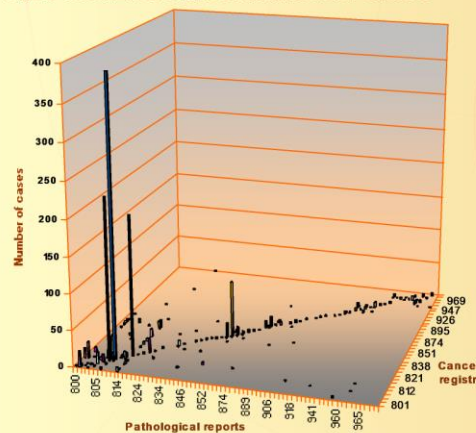
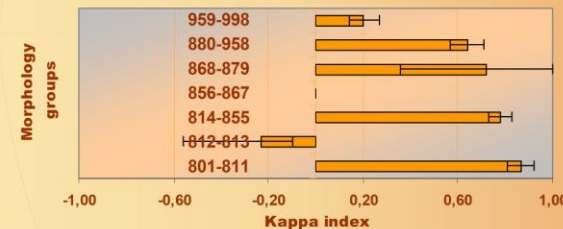


Figure 2. KAPPA INDEX BY MORPHOLOGY GROUPS



CONCLUSIONS:

A high overall agreement is observed. For some morphology groups the future automated process could be implemented maintaining the current quality indicators in the cancer registry. Other groups will require a manual examination before being stored.

- D. Quality of data on morphology codes from pathology reports in Murcia-Spain.

Conclusion:

- A high overall agreement is observed.
- For some morphology groups the future automated process could be implemented maintaining the current quality indicators in the cancer registry.
- Other groups will require a manual examination before being stored.

▶ I. SOURCES OF INFORMATION

▶ II. VALIDATION OF DIGITAL SOURCES OF INFORMATION

▶ III. AUTOMATED PROCESS OF DATA LOAD IN THE OFICIAL HEALTH CARE DATABASE

Fist step: normalization

Second step: codification

Third step: linkage with person in oficial Health Care Database

▶ IV. AUTOMATED PROCESS OF CANCER CONSOLIDATION IN CANCER REGISTRY PROGRAM

Decisional algorithms

▶ III. AUTOMATED PROCESS OF DATA LOAD IN THE OFICIAL HEALTH CARE DATABASE

Fist step: normalization: automatic with commands

Second step: automatic codification in no codificated variables

Third step: linkage with person in oficial Health Care Database

Process of data load:

- **CMBD**
- **DEATH CERTIFICATE**
- **PATHOLOGICAL REPORTS**

Importación	Ventana	?
Carga Fichero Fuente		
Carga Datos DEFUN		
Informe Fuente		
Procesar Informes Fuente		
Informes sin Persona		
Eliminación de Informes sin Persona		
Informes con Codificación Errónea		
Captación de Tumores		

▶ AUTOMATED PROCESS OF DATA LOAD ▶▶ CMBD AND DEATH CERTIFICATE

Automated selection of cases that meet the criteria: ICD-10 codes in any
CMBD/ DEATH CERTIFICATE
diagnosis.

	Rango CIE-9	Rango CIE-10	Incluidos	CIE-9	CIE-10
Malignos	140-209	C00-C97	Se incluyen todos	140-209	C00-C97
Benignos	210-229	D10-D36	Sólo SNC	225	D32-D33
In situ	230-234	D00-D09	Se incluyen todos		
Inciertos	235-239	D37-D48	Vejiga	236.7	D41.4
			SNC	237.5-237.9	D42-D43
Inciertos en CIE9 y CIE10 pero malignos en CIEO-3			Trastornos mieloproliferativos	238.4	D45
				238.5	D46
			crónicos	238.6	D47
				238.7	

Automatic linkage with the people
Health Care System database.

Calidad	Gestión Proyectos	Auxiliares	Listas
Fusión de Expedientes			
Reasignación Expediente de Desconocido			
Informes de CMBD sin id de Person			

Correction of error (less than 1%).

New people no previously registered (less than 1%).

▶▶ AUTOMATED PROCESS OF DATA LOAD ▶▶ PATHOLOGICAL REPORTS (NO CODES):

An automated process has been implemented to select cases and to generate ICEO-3 location and morphology codes from the pathological anatomy diagnoses

--Table of terms with their corresponding codes.

--Automatic algorithm to classify reports:



Cases selected as registered:

high sensitive algorithm that includes most of the registered cases (true positives) but also includes non-registered (false positives).



Cases selected as no registered:

high specificity: automatically discarded (>90% of reports) without manual review.

SNOMED Mapping – ICD-10 – ICD-3
Topography and Morphology
Conversion tables

Palabra	Código	Localización	Código	Histología	Prioridad	Palabra completa	Fecha	Profesional
ADAMANTINOMA	92513		ADAMANTINOMA DE HUESOS LAROS (C40...)				24/11	MDN24R
ADENOCARCINOMA	81403		ADENOCARCINOMA, SAI				28/11	MCL59Z
ALTO GRADO	80772		NEOPLASIA INTRAEPITELIAL ESCAMOSA, ALTO GRADO				24/11	MDN24R
ANEMIA REFRACTA...	99803		ANEMIA REFRACTARIA				24/11	MDN24R
ANGIOENDOTELIOM...	97123		LINFOMA DE CELULAS B GRANDES INTRAVASCULAR (...)				16/11	MDN24R
AREB	99803		ANEMIA REFRACTARIA				16/11	MDN24R
ASKN	93653		TUMOR DE ASKN				16/11	MDN24R
ASTROCITOMA	93841		ASTROCITOMA SUBEPENDIMARIO DE CELULAS GIGAN...				16/11	MDN24R
ATIPICO	80013		CELULAS TUMORALES MALIGNAS				24/11	MCL59Z
BLASTOMA	80003		NEOPLASIA MALIGNA				16/11	MDN24R
CADENAS PESADAS	97623		ENFERMEDAD DE LA CADENA PESADA SAI				16/11	MDN24R
CARCINO	80103		CARCINOMA SAI				16/11	MDN24R
CELULAS NEOPLASI...	80001		NEOPLASIA INCERTO SI ES BENIGNO O MALIGNO				22/12	MCL59Z
CN II	80772		NEOPLASIA INTRAEPITELIAL ESCAMOSA, ALTO GRADO				16/11	MDN24R
CN-III	80772		NEOPLASIA INTRAEPITELIAL ESCAMOSA, ALTO GRADO				16/11	MDN24R
CN-III	80772		NEOPLASIA INTRAEPITELIAL ESCAMOSA, ALTO GRADO				16/11	MDN24R
CITOFENIA REFRAC...	99853		CITOFENIA REFRACTARIA CON DISPLASIA MULTINUCLE...				16/11	MDN24R
CLOROMA	99303		SARCOMA MELOIDE (VER TAMBIEN M- 98617G)				16/11	MDN24R
CORDOMA	93723		CORDOMA DESDIFERENCIADO				16/11	MDN24R

▶ ▶ AUTOMATED PROCESS OF DATA LOAD ▶ ▶ PATHOLOGICAL REPORTS (NO CODES):

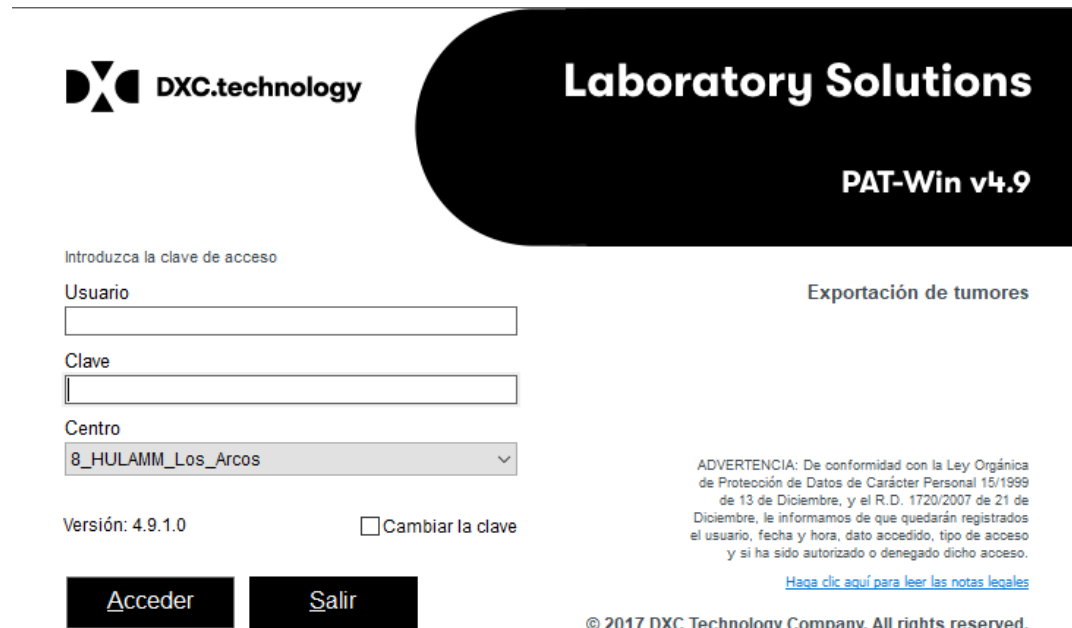
Término codificación									
Palabra	Código	Localización	Código	Histología	Prioridad	Palabra completa	Fecha	P	
ADAMANTINOMA			92613	ADAMANTINOMA DE HUESOS LARGOS (C40...)		<input type="checkbox"/>	24/11...	M	
ADENOCARCINOMA			81403	ADENOCARCINOMA, SAI		<input type="checkbox"/>	29/11...	M	
ALTO GRADO			80772	NEOPLASIA INTRAEPITELIAL ESCAMOSA, ALTO GRADO		<input type="checkbox"/>	24/11...	M	
ANEMIA REFRACTA...			99803	ANEMIA REFRACTARIA		<input type="checkbox"/>	24/11...	M	
ANGIOENDOTELIOM...			97123	LINFOMA DE CELULAS B GRANDES INTRAVASCULAR (...)		<input type="checkbox"/>	16/11...	M	
AREB			99803	ANEMIA REFRACTARIA		<input checked="" type="checkbox"/>	16/11...	M	
ASKIN			93653	TUMOR DE ASKIN		<input type="checkbox"/>	16/11...	M	
ASTROKITOMA			93841	ASTROKITOMA SUBPENDIMARIO DE CELULAS GIGAN...		<input type="checkbox"/>	16/11...	M	
ATIPICO			80013	CELULAS TUMORALES MALIGNAS		<input type="checkbox"/>	24/11...	M	
BLASTOMA			80003	NEOPLASIA MALIGNA		<input type="checkbox"/>	16/11...	M	
CADENAS PESADAS			97623	ENFERMEDAD DE LA CADENA PESADA SAI		<input type="checkbox"/>	16/11...	M	
CARCINO			80103	CARCINOMA SAI		<input type="checkbox"/>	16/11...	M	
CELULAS NEOPLASI...			80001	NEOPLASIA INCIERTO SI ES BENIGNO O MALIGNO		<input type="checkbox"/>	22/12...	M	
CIN III			80772	NEOPLASIA INTRAEPITELIAL ESCAMOSA, ALTO GRADO		<input type="checkbox"/>	16/11...	M	
CIN-III			80772	NEOPLASIA INTRAEPITELIAL ESCAMOSA, ALTO GRADO		<input type="checkbox"/>	16/11...	M	
CINIII			80772	NEOPLASIA INTRAEPITELIAL ESCAMOSA, ALTO GRADO		<input type="checkbox"/>	16/11...	M	
CITOPENIA REFRAC...			99853	CITOPENIA REFRACTARIA CON DISPLASIA MULTILINAJE		<input type="checkbox"/>	16/11...	M	
CLODOMA			99303	SARCOMA MIELOIDE (VER TAMBIEN M- 98617/3)		<input type="checkbox"/>	16/11...	M	
CORDOMA			93723	CORDOMA DESDIFERENCIADO		<input type="checkbox"/>	16/11...	M	

Conversion tables

▶ ▶ AUTOMATED PROCESS OF DATA LOAD ▶ ▶ PATHOLOGICAL REPORTS (NO CODES):

- Automated upload of cases as external sources to RCM: all pathology reports that have met the inclusion criteria are uploaded to RCM.

Pathological anatomy
downloads from PATWIN ▶



DXC.technology **Laboratory Solutions**
PAT-Win v4.9

Introduzca la clave de acceso

Usuario

Clave

Centro

Versión: 4.9.1.0 Cambiar la clave

[Exportación de tumores](#)

ADVERTENCIA: De conformidad con la Ley Orgánica de Protección de Datos de Carácter Personal 15/1999 de 13 de Diciembre, y el R.D. 1720/2007 de 21 de Diciembre, le informamos de que quedarán registrados el usuario, fecha y hora, dato accedido, tipo de acceso y si ha sido autorizado o denegado dicho acceso.

[Haga clic aquí para leer las notas legales](#)

Acceder **Salir**

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Configuración de la Consulta



Conexión Hospitalaria

Hospital desde el que se realiza la consulta:

Opciones de búsqueda

- Exportación Estándar
 Exportación SMS
 Recuperar Estudios Anteriores
- Estudios por literal
 Exportación SAS

Exportación

Salto de línea:
 Tabulación:
 Coma:

Comilla doble:
 Comilla simple:

Exportar en formato XML

Datos del Paciente

NHC	mynhc	14
Apellido1	myApellido1	1
Apellido2	myApellido2	2
Nombre	myNombre	3
Fecha Nac.	myfnac	4
NASS	myNSS	7
DNI	myDNI	6
CIP	mytarjetasanitaria	8
CIP Regional	mytarjetasanitariaregional	9
Sexo	mysexo	5
Domicilio	mydom	10
Localidad de nacimiento	mylocalidad	15

Datos del Estudio

Número de estudio	myestudio	13
Tipo de estudio	Tipo de estudio	0
Fecha de registro	Fecha de registro	0
Fecha de toma	myftoma	17
Servicio solicitante	Servicio solicitante	0
Médico solicitante	Médico solicitante	0
Patólogo responsable	Patólogo responsable	0
Fecha de diagnóstico	myfinforme	16
Descr. diagnóstico	Diagnostico_AP	25
Descr. macroscópica	mymacro	21
Descr. macroscópica	mymicro	22
Datos clínicos	mydiagclinico	20
Muestra	myorgano	26
Técnicas	mytecnica	27

Datos del Tumor

Comentarios	Comentarios	0
Tratamiento sincrónico	Tratamiento sincrónico	0
Recidiva	Recidiva	0
Estudio post-mortem	Estudio post-mortem	0
Metástasis	Metástasis	230
Estadio	Estudio del tumor	240
Tamaño menor (mm.)	Tamaño menor (mm.)	0
Tamaño mayor (mm.)	Tamaño mayor (mm.)	0
Nº de ganglios afectados	Nº de ganglios afectados	270
Nº de ganglios extraídos	Nº de ganglios extraídos	0
Diferenciación tumoral	Diferenciación tumoral	0
Lateralidad	Lateralidad	0
PN	PN	0
PT	PT	0

Configuración SMS

Fecha de búsqueda:

Receptores hormonales:

Inmunohistoquímica:

Texto para las fechas:

Separador:

Conversión sexo

Hombre: →

Mujer: →

Otro: →

Aceptar

▶ I. SOURCES OF INFORMATION

▶ II. VALIDATION OF DIGITAL SOURCES OF INFORMATION

▶ III. AUTOMATED PROCESS OF DATA LOAD IN THE OFICIAL HEALTH CARE DATABASE

Fist step: normalization

Second step: codification

Third step: linkage with person in oficial Health Care Database

▶ IV. AUTOMATED PROCESS OF CANCER CONSOLIDATION IN CANCER REGISTRY PROGRAM

Decisional algorithms

▶ **CONSOLIDATION OF TUMOURS:**

▶ ▶ **Joint information of several sources in the same tumor:**

-Tumor already exist :

--- Same tumor: update.

--- Different tumor: multiple: create a new second primary tumor.

-Tumor doesn't exist:

--- Creation of new incident tumors.

▶ **DEVELOPMENT OF ALGORITHMS TO FILL IN THE TUMOUR VARIABLES:**

- Date of incidence: ENCR rules.

- Site / morphology

- behaviour / grade

- Base of diagnosis

- Sources of information

- ICD-10

- Date of death

▶ IV. AUTOMATED PROCESS OF CANCER CONSOLIDATION IN CANCER REGISTRY PROGRAM: Decisional algorithms

Definition of automated case:

- Codes from different sources must be in agreement:
 - - Topography: 3 digits level.
 - - Morphology: morphology group ICDO.

- Number of sources of information:
 - always more than one, except for pathological reports.

Select the more specific code in morphology of the same group, the higher number.

Auxiliares	Listados	Importación	V
Tipos de Asistencia			
Tipos de Grado de Diferenciación			
Modod de Presentación			
Tipo de Extensión			
Tipo de Resultado de Autopsia			
Tipo de Lugar Muerte			
Estado Vital			
Tipo de Documentación			
Tipo de Fuente			
Fuente Departamento			
Localización Grupo			
Histología Grupo			
Base diagnóstico			
Lateralidad			
Tipo de Informe			
Tratamiento			
Localización			
Localización Múltiple			
Histología			
Histología-Localización			
Histología Múltiple			
Grupo Histológico Múltiple			
Estado			
Fase			
Estado-Fase			
Centro - Servicio - Codificación			
Variables Globales			
Término - Codificación			

▶ IV. AUTOMATED PROCESS OF CANCER CONSOLIDATION IN CANCER REGISTRY PROGRAM: Decisional algorithms

A selected sample of cases have been compared:

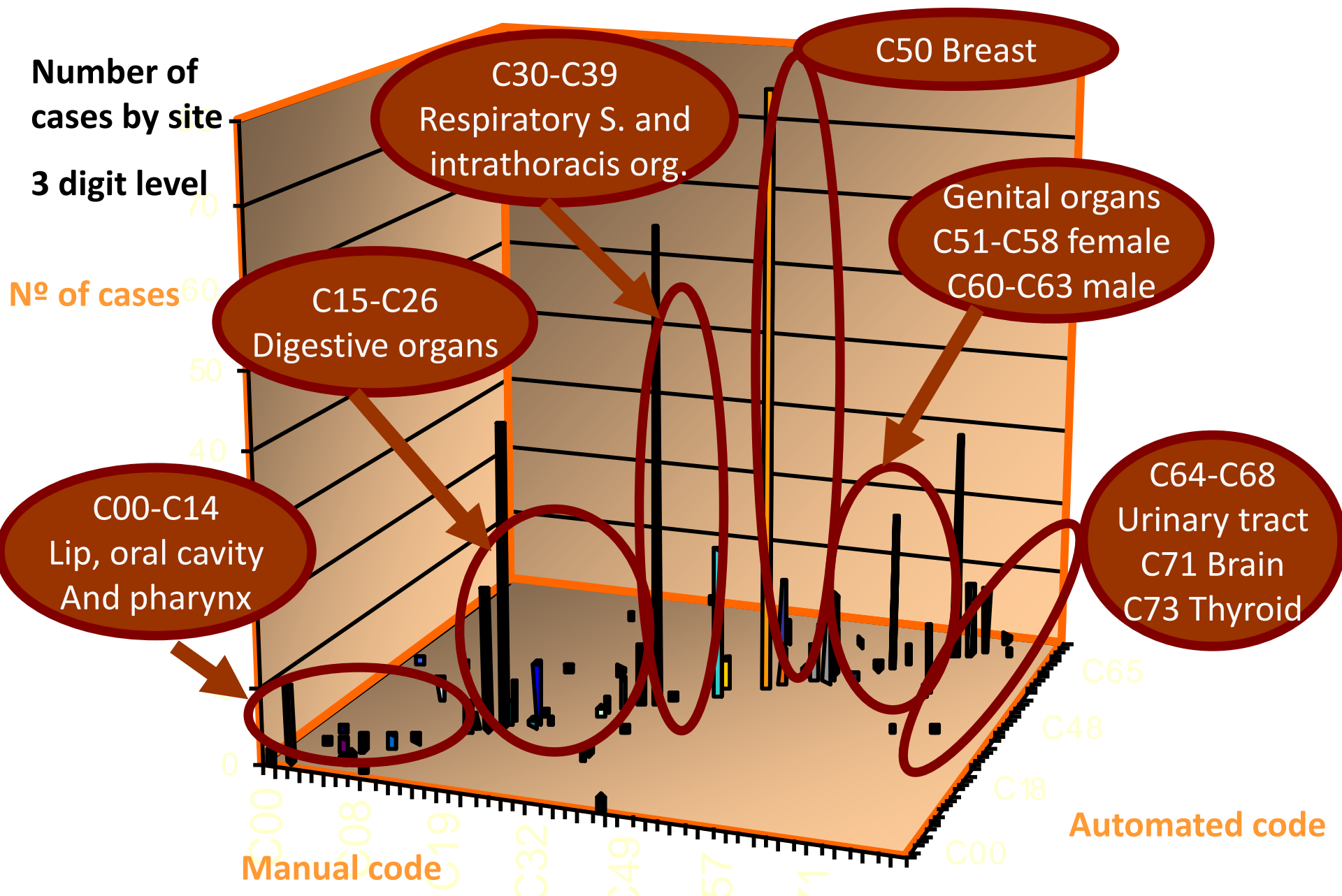
- ▶ 79% of cases are correctly coded automatically at three digit level topography.
- ▶ 71% of cases are correctly coded automatically at three digit level morphology.
- ▶ 84% of cases are correctly coded automatically by morphology groups ICDO.

Percentage of agreement by main sites

Manual code

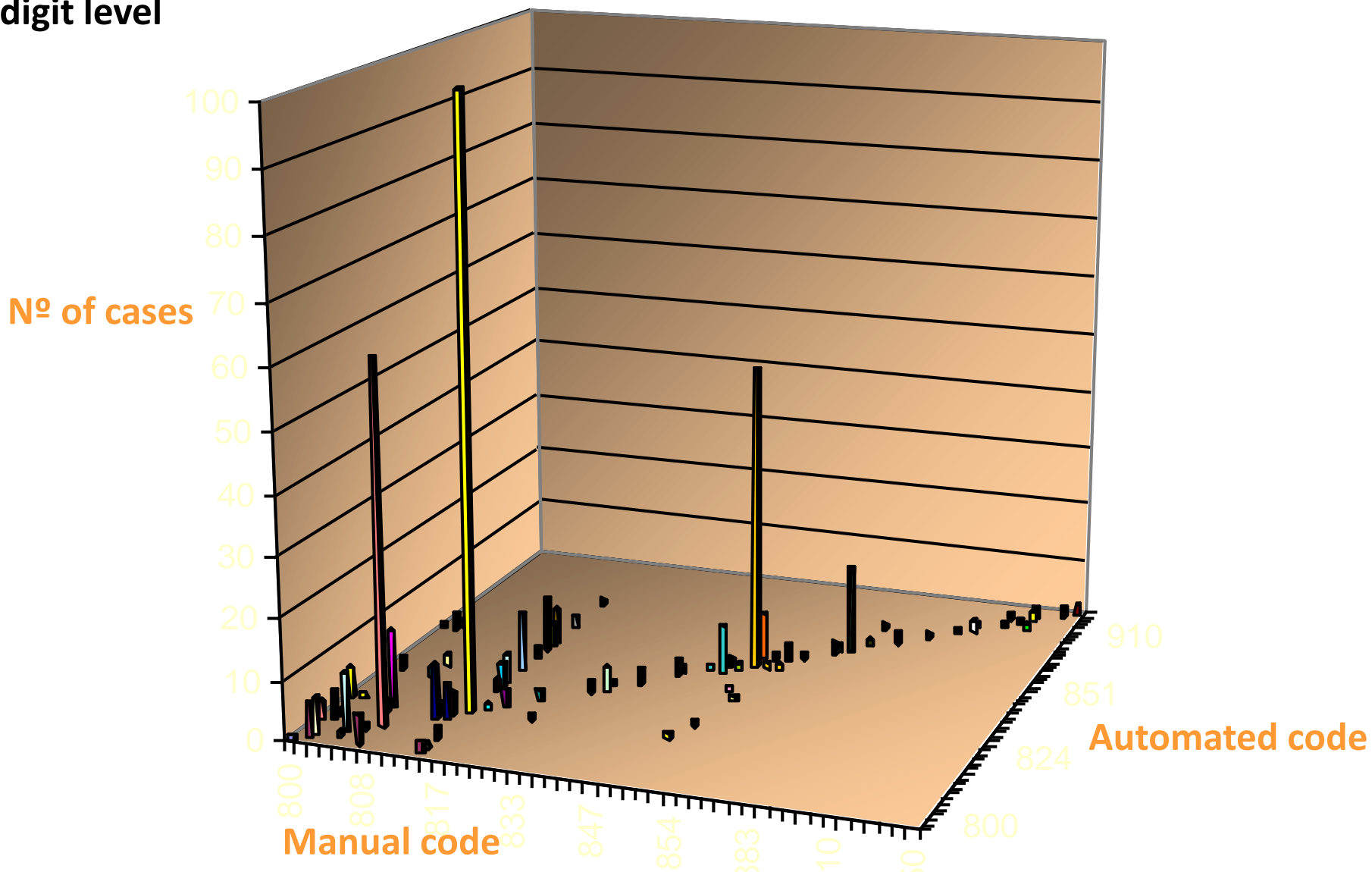
Automated code

% Agreeemnt	C02	C08	C15	C16	C18	C20	C21	C34	C50	C52	C53	C54	C56	C57	C61	C67	C71	C73	C75
C02 Other and un. tongue	91,0	9,0																	
C16 Stomach			10,0	90,0															
C18 Colon					100,0														
C20 Rectum					33,0	53,0	7,0	7,0											
C34 Lung								98,0								2,0			
C50 Breast									100,0										
C53 Cervix uteri											33,0	60,0				7,0			
C54 Corpus uteri										5,0		42,0		5,0		47,0			
C56 Ovary												100,0	0,0						
C61 Prostate															100,0				
C67 Bladder					1,0					6,0		3,0				89,0			
C71 Brain																		100,0	
C73 Thyroid									9,0									82,0	9,0



Number of cases by morphology

3 digit level



By site:

-High agreement in

--BREAST, COLON, PROSTATE, BRAIN and LUNG that represent a high number of incident cases.

-Low agreement in

--FEMALE GENITAL ORGANS, BLADDER

By morphology group:

-High agreement in

- Squamous cell neoplasms**

- Adenomas and adenocarcinomas**

- Nevi and melanoma**

-Low agreement in

- Transitional cell papillomas and carcinomas**

- Cystic, mucinosus and serous neoplasms**

Causes of disagreement:

- Lack of codes accuracy in sources of information.**
- Problems with the conversion between classifications.**
- The algorithms implemented in the software are being reviewed.**
- A high percentage of discordant site have been coded in adjacent topography.**
- A high percentage of discordant morphology have been coded in a less specific morphology group.**

BY THE MOMENT,

‘AUTOMATIZATION HELP BUT IS NOT ENOUGH’

WE HAVE TO IMPROVE:

- **DIGITALIZATON**
- **PRIMARY CODIFICATION**
- **IMPLEMENTATION OF RULES**
- **MAPING**
- **CONVERSION PROGRAMS...**



¡THANKS FOR YOUR ATENTION!