

The JRC-ENCR Quality Check Software (QCS) for the validation of cancer registry data: user compendium

JRC-ENCR QCS 2.x JRC CSV Data layout converter

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Update May 2024, QCS version 2.2

Based on JRC Technical report (May 2024, update) and ECIS protocol 2022 (February 2024)

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Acknowledgments

The authors acknowledge the following institutions and entities for the invaluable feedback provided running the previous versions of the JRC-ENCR Quality Check Software.

- East Switzerland Cancer Registry
- OmniSoftory Engineering S.A., Switzerland
- National Institute for Cancer Epidemiology and Registration (NICER), Switzerland
- National Agency for Cancer Registration (NACR), Switzerland
- Swiss Childhood Cancer Registry
- Basel Cancer Registry, Switzerland
- Bern Cancer Registry, Switzerland
- Samara Cancer Registry, Russian Federation
- Trapani Cancer Registry, Italy
- Cancer Registry of Puglia Section of Health Local Unit Barletta Andria Trani, Italy
- Veneto Cancer Registry, Italy
- Castellon Cancer Registry, Spain
- Basque Country Cancer Registry, Spain
- Girona Cancer Registry, Spain
- Tarragona Cancer Registry, Spain
- Greater Poland Cancer Registry
- Polish National Cancer Registry
- Cancer Registry of Republic of Slovenia
- Morphological Tumour Registry, Luxembourg
- Cancer Registry of Norway
- Isère Cancer Registry, France
- Association of the French Cancer Registries (FRANCIM)
- Cancer Registry of Baden-Württemberg, Germany
- National Cancer Registry Ireland

Introduction

The European Network of Cancer Registries (ENCR) and the European Commission's Joint Research Centre (JRC) have launched in 2022 a data call for updating cancer burden indicators available in the European Cancer Information System (ECIS) (<u>https://ecis.jrc.ec.europa.eu/</u>), following an updated <u>ECIS data call protocol</u>.

In order to enable cancer registries to perform data quality checks and to test the adherence of their data to the protocol required format, the JRC starting from 2015 has been developing the <u>JRC-ENCR Data Quality Check Software</u> (QCS). Complementary to the QCS, the CSV Data Layout Converter has been released in 2021, as an additional tool to facilitate the preparation of the incidence file.

This version of the QCS is updated to the requirements of the 2022 ECIS data call protocol and the <u>2024 JRC Technical report</u> "A common data quality check procedure for European cancer registries". This new version benefits from the experience gathered in validation of previously submitted data from European registries, and from registries' feedback.

The development of the QCS is an evolving process, with regular updates that ensures alignment with European and International recommendations and classifications.

The present document provides technical guidance to the software, and helps to understand and interpret its output.

The main changes and improvements included in this version compared to previous one, are detailed in **Annex 1 – Changes and improvements from the previous versions**. A list of known bugs and issues to be addressed in a later release is available in **Annex 2 – Known** *issues and future improvements.*

1 Software overview

The JRC-ENCR Quality Check Software (QCS) is an open-access software developed to facilitate standardisation and validation of population-based cancer registries (PBCRs) data.

The QCS has been designed as a stand-alone desktop application that can run locally by cancer registries, without the need of internet connection.

The majority of the checks performed by this version are based on the 2024 JRC Technical report, which includes recommendations for checking internal consistency of the cancer incidence data and provides a comprehensive and standardised list of data quality checks to be adopted by European PBCRs.

The software checks the four different data files (incidence, mortality, population and life tables) required by the ECIS protocol. The QCS output consists of four files (in PDF, TXT and CSV format) containing the detailed list of warnings and errors detected in the validation run for each record of the input data file.

Population, life tables and mortality files have a simple structure, which require univariate checks validation (checks on single variables). Incidence files are more complex, and in addition to univariate validation the QCS performs checks between variables of the same record (multivariate checks) and between variables of different records (duplicates and multiple primary tumours checks).

2 System requirements and installation

The QCS has been initially developed for Windows operating systems that support Java (Windows 10 and above).

Starting from version 1.7, all versions of the QCS can also run on MacOS and Linux operating systems (see sections 2.5 and 2.6). Sections 2.1-2.4 below refer to Windows operating systems.

2.1 In case Java software is not installed on your computer

Java software is needed in order to run the QCS. In case Java is not installed on your computer, please follow the steps below, otherwise go to **section 2.3**.

- Go to <u>Java.com</u> and click on the **Download Java. On the next landing page click the Download Java** button to start the download;
- If a Filedialog box appears, choose the folder where you wish to save the executable file (*.exe) and click on the **Save** button;
- Double click on the executable file to start the software installation;
- Depending on your security settings, you may be prompted dialog boxes asking for permission to continue. Confirm that you want to proceed with the installation;
- The installation process starts. Click the **Install** button to accept the license terms and to continue.

Following screenshots shows an example of Java installation, which could change in future updates.





After having completed the installation of Java, you can install the QCS, as explained in section 2.3.

2.2 Further information and troubleshooting related to Java

If you need help in installing Java Runtime Environment on your machine, kindly ask to your System Administrator or local IT support to install it for you.

You should also setup the JAVA_HOME environment variable correctly configured. Usually, this is done automatically. Please check with your System Administrator.

If you wish to manually set the JAVA_HOME variable in Windows (for other systems the procedure may vary) please click on *Start* \rightarrow *Control Panel* \rightarrow *search for the "variable" keyword* \rightarrow *Edit environment variables for your account* \rightarrow *Environment variables* to configure the Java environment (see screenshot below):

The official requirements for Java can be found at: <u>https://www.java.com/en/download/win_sysreq-sm.jsp</u>

Remember to choose the correct version for your operating system (Windows 32 bit or Windows 64 bit).

Please note: there are two versions of Java environments, Java Developer Kit (JDK) and Java Runtime Environment (JRE). **Please install the JRE**.

stem Properties	
Computer Name Hardv	ware Advanced Remote
Environment Variabl	es 🗙
User variables for m	entafa
Variable	Value 🔺
ForcePSTPath	\\net1.cec.eu.int\jrc-services\JRC-PST\
StartMenuDir	%USERPROFILE%\Appdata\Roaming\M
TEMP	%USERPROFILE%\AppData\Local\Temp
TMP	%USERPROFILE%\AppData\Local\Temp 🥃
	New Edit Delete
System variables	
Variable	Value
DG_COMPUTER	IPR 🔟
DG_USER	IPR
FP_NO_HOST_C	, NO
JAVA_HOME	C:\Program Files (x86)\Java\jre6 🚽 🚽
1	
	New Edit Delete
	OK Cancel

Further information on how to install Java without third party sponsor offers is available at: <u>https://www.java.com/en/download/faq/disable_offers.xml</u>.

2.3 How to install the QCS

Download the latest version of the software from the <u>JRC-ENCR website</u>, and extract the files on your computer in a dedicated folder. Please follow the instructions below according to your PCs operating system.

2.4 Running on Windows

- 1. Open the folder containing the QCS
- Double click on the "JRC-ENCR-QCS-2GB.bat" file (suggested option). If the QCS does not start or does not work properly, double click on the "JRC-ENCR-QCS.bat"

2.5 Running on macOS

- 1. Double click the Terminal icon (or label, depending by your view settings) to open a Terminal window
- Enter the Terminal window and move into the folder created at step 1. For example, if the target QCS file was named "JRC-ENCR-QCS-V2.x.zip", then you should execute the command like

cd Desktop/JRC-ENCR-QCS-V2.x

3. Execute the file having the extension ".sh". For example if the file is named "start-jrc-encrqcs.sh", then type the command:

source jrc-encr-qcs.sh

2.6 Running the QCS on Linux operating systems

- 1. Move into the directory containing the QCS
- 2. Execute the QCS by running the ".sh" file:

source start-jrc-encr-qcs.sh

2.7 Verify the correct installation

Navigate to the folder where you extracted the software and before running it (as explained in section 5.1), check if it is correctly installed.

The expected directory structure is shown below:

← → ~ ↑ 📙 > T	'his PC →	(U:) JRC.F.1 > 1b-Cancer Information	→ QCS → jrc-qcs-2.x	✓ O Search jrc-qo	cs-2.x
	^	Name	Date modified	Туре	Size
📌 Quick access			25 (01 (2024 15:20	File Colden	
📃 Desktop	*	config	25/01/2024 15:29	File folder	
Documents	*	docs	25/01/2024 15:30	File folder	
		lib	25/01/2024 15:31	File folder	
QCS	*	logs	25/01/2024 15:31	File folder	
Main_docs	*	📙 output	25/01/2024 15:31	File folder	
AIRTUM_XXIV	*		25/01/2024 15:34	File folder	
Presentation	*	JRC-ENCR-QCS.bat	15/01/2024 16:25	Windows Batch File	3 KB
Spese per Piksel	*	📄 jrc-encr-qcs.sh	15/01/2024 16:25	SH File	1 KB
Validation_doc	*	JRC-ENCR-QCS-2GB.bat	15/01/2024 16:25	Windows Batch File	3 KB
		JRC-NET1-QCS.bat	15/01/2024 16:25	Windows Batch File	2 KB
CNS recommandat	tior 🖈	JRC-NET1-QCS-1GB.bat	15/01/2024 16:25	Windows Batch File	2 KB
Piksel	*	🛃 jrc-qcs-2.x.jar	15/01/2024 16:25	JAR File	1290 KB
jrc-qcs-2.1.5	*	LICENSE.txt	05/12/2023 10:14	Text Document	10 KB
Final 20231110		test-suite.sh	15/01/2024 16:25	SH File	14 KB
JRC QC report		RUN-QCS.bat	17/04/2024 17:38	Windows Batch File	3 KB
Otto		🚳 qcs.db	17/04/2024 11:40	Data Base File	396 KB

The folder JRC-ENCR-QCS-V2.x includes the following files and folders:

- The LICENSE.txt file containing the usage license
- The executable files JRC-ENCR-QCS.bat and JRC-ENCR-QCS-2GB.bat; The executable files JRC-NET1-QCS.bat, JRC-NET1-QCS-1GB.bat, RUN-QCS.bat;
- Files *jrc-encr-qcs.sh* and *test-suite.sh*;
- The application's file *jrc-qcs-2.x.jar*;
- Installation folders: config, docs, lib (installed with the QCS);
- Runtime folders: *logs, output, temp* (created at first usage).

The file *JRC-ENCR-QCS.bat* will run the QCS with 1GB of RAM memory allocated by Java, whereas *JRC-ENCR-QCS-2GB.bat* will use 2GB of RAM memory.

The file *jrc-encr-qcs.sh* is the standard ".sh" script for running the application on the macOS and Linux systems (see sections 2.5 and 2.6 above).

config: this folder contains configuration files, such as those for ranges of variable values, general application settings (with the possibility to disable some functionalities) etc.

docs: this folder contains relevant documentation files, i.e. the QCS compendium (this document), the JRC Technical Report "A common data quality check procedure for European cancer registries" and the ECIS call for data protocol 2022 in PDF format. The file change_log.txt is also included with the list of the most recent QCS versions and updates. The subfolder examples includes some sample scripts for advanced users (see Annex 4 – Running the JRC-ENCR QCS in background).

<u>lib</u>: this folder includes the jar library files used by the software at run time.

logs: this folder stores the log file of all the QCS activities.

<u>output</u>: this folder is created after the QCS is run for the first time. It includes four different subfolders, including the output reports produced by the QCS for *Incidence, Mortality, Population* and *LifeTable* files respectively.

temp: this folder is created after the QCS is run for the first time. It contains all the raw files (working files) used to generate the final output reports.

3 How to prepare an input file for the QCS

In this section an example of input files accepted by the software is given. Input files should be formatted as follows:

- as text files (.csv or .txt) with semicolon (;) separators only;
- with first line as header.

3.1 Incidence File

The input file must follow the format of the <u>ECIS data call protocol</u> (section 3.1.1). It can be either created following the instructions below (section 3.1.2), or using the JRC CSV Data layout converter (section 3.1.3).

3.1.1 The ECIS data call protocol

Below is the summary table of the variables and related coding, as required by the ECIS data call protocol.

Variable name	Variable description	Format	Maximum length	Missing/ unknown	Coding
PAT ²	Patient identification code	А	50	Not allowed	According to registry coding
MoB	Month of birth	F	2	99	Range of allowed values: 1 - 1
YoB	Year of birth	F	4	9999	Range of allowed values: > 1842 and ≤ the current year
Age	Age at diagnosis (incidence date) in years	F	3	999	Range of allowed values ≥ 0 and < 12
Sex	Sex at birth	F	1	9	1 → Male 2 → Female 3 → Other

² *PAT* should be a code assigned by the registry that is <u>not</u> to be used elsewhere (e.g. in a hospital). So, it cannot be an official personal number. It may be an encrypted personal number as long as this specific encryption is not used by any other organisation. The JRC will provide the tool to the CRs to do it.

Surv_time	Duration of survival in days	F	5	99999	≥0
YoF	Year of last known vital status	F	4	9999	Range of allowed values: > 1941 and ≤ the current year
MoF	Month of last known vital status	F	2	99	Range of allowed values: From 1 to 12
Vit_stat	The last known vital status	F	1	9	1→ Alive 2→ Dead
Autopsy ⁵	Incidental finding of cancer at autopsy	F	1	9	0→No 1→Yes
Variable name	Variable description	Format	Maximum length	Missing/ unknown	Coding
	ated to follow-up				1
Variables rela					
					7→ Null cell; Non T-non B 8→ NK cell (natural killer cell) 0→ Not applicable
	пппппорпенотуре				5→ 1-cell; 1-precursor 6→ B-Cell; Pre-B; B-precursor
Grade ⁴	ICD-O-3 grade of differentiation / immunophenotype	F	1	9	differentiated 3→ Grade III, Poorly differentiated 4→Grade IV, Undifferentiated, anaplastic 5→ T-cell; T-precursor
					1→Grade I, Well differentiated 2→ Grade II, Moderately
Beh	ICD-O-3 behaviour	F	1	Not allowed	 0→ Benign neoplasm 1→ Neoplasm of uncertain and unknown behaviour 2→ In situ neoplasm 3→ Malignant neoplasm
Morpho	ICD-O-3 morphology code	F	4	Not allowed	Valid code in any ICD-O-3 version
Торо	ICD-O-3 topography code	A	4	Not allowed	Valid code in ICD-O-3
BoD	Basis of diagnosis	F	1	9	0→Death certificate only (DCO) 1→Clinical 2→Clinical investigation 4→Specific tumour markers 5→Cytology 7→Histology 8→Cytogenetic and/or molecular testing 9→ Unknown
Yol	Year of incidence	F	4	Not allowed	Range of allowed values: From 1941 to present
Mol	Month of incidence	F	2	99	Range of allowed values: 1 - 12
TUM	Tumour identification	A	50	Not allowed	According to registry coding
Tumour variab					
Geo_label	Name of the geographical area of residence at Diagnosis	А	50	9	Blank \rightarrow not applicable
Geo_code	Code for the geographical area of residence at Diagnosis	A	10	XX99	highest level of administrative sub divison that can be provided ³ . Blank → not applicable

³ NUTS 3 codes should be provided for regional registries covering NUTS 3 areas such as French *Départements*, Italian *Province* and Spanish *Provincias*.

⁴ The grade of tumours of the central nervous system should be coded according to table 27 of ICD-O-3.

⁵ In autopsy cases, incidentally found at autopsy, the *vital status* is always 2 (dead) and the *survival* time is 0 days.

ICD ^{6,7}	ICD edition for coding cause of death	F	2	99	Range of allowed values: <12 Blank → Not applicable
CoD ^{6,7}	Official underlying cause of death	А	4	R99 (ICD-10) 7999 (ICD-9)	According to ICD Blank → Not applicable
Stage variab	les				
TNM_ed	TNM edition	F	2	99	Allowed values: ≤ 8
cT ⁸	Clinical T-category	А	12	9	
cN ⁸	Clinical N-category	А	12	9	According to the TNM
сМ ⁸	Clinical M-category	А	12	9	Classification of Malignant
рТ ^{8,9}	Pathological T-category	А	12	9	Tumours
pN ^{8,9}	Pathological N-category	А	12	9	Blank → not applicable
рМ ^{8,9}	Pathological M-category	А	12	9	
ToS	Staging system	A	3	9	D → Dukes' stage E → Extent of disease F → FIGO stage S → TNM stage, unknown whether clinical or pathological clS → clinical TNM stage paS → pathological TNM stage paS → pathological TNM stage ypS -> pathological TNM stage after neoadjuvant therapy cpS → combination of clinical & pathological TNM stage coS → condensed TNM stage esS → essential TNM stage Ti1 → Tier 1 stage for paediatric tumours Ti2 → Tier 2 stage for paediatric tumours, findings at surgery when NO chemotherapy prior to surgery SIO → SIOP Tier 2 stage for Wilms tumours: findings at surgery when chemotherapy prior to surgery 8 → Other staging system
Stage variab	les				2 2 .

Stage variables

Variable name	Variable description	Format	Maximum length	Missing/ unknown	Coding
Stage	Stage	A	3	9	$0 \rightarrow Stage 0, stage 0a, stage 0is, carcinoma in situ, non-invasive 1 \rightarrow Stage I, FIGO I, localized, localized limited (L), limited, Dukes A 1A -> Stage IA, FIGO IA, Ann Arbor 1B -> Stage IB, FIGO IB 1B1 -> FIGO IB1 2 \rightarrow Stage II, FIGO II, localized advanced (A), locally advanced, advanced, direct extension, Dukes B 2A -> Sage IIA, FIGO IIA 2B -> Stage IIB, FIGO IIB 3 \rightarrow Stage III, FIGO III, regional (with$

					or without direct extension), R+, N+, Dukes C 4 → Stage IV, FIGO IV, metastatic, distant, M+, Dukes D See also Annex 4 for
Treatment	variables	1	1	1	1
Surgery ^{10,11}	Resection of the primary tumour	F	1	9	 0 → No 1 → Yes, without specification 2 → Yes, local surgery only¹² 3 → Yes, 'operative' surgery¹³
Rt	Radiotherapy	F	1	9	 0 → No 1 → Yes, without specification 2 → Yes, neoadjuvant (pre-operative) radiotherapy 3 → Yes, adjuvant (post-operative) radiotherapy
Cht	Chemotherapy	F	1	9	 0 → No 1 → Yes, without other specification 2 → Yes, neoadjuvant (preoperative) 3 → Yes, adjuvant (postoperative) 4 → Yes, both neoadjuvant and adjuvant
Tt ¹⁴	Targeted therapy (including monoclonal antibodies)	F	1	9	$\begin{array}{c} 0 \rightarrow No \\ 1 \rightarrow Yes \end{array}$
lt	Immunotherapy (excl. monoclonal antibodies)	F	1	9	$\begin{array}{c} 0 \rightarrow No \\ 1 \rightarrow Yes \end{array}$
Ht	Hormone therapy	F	1	9	$\begin{array}{c} 0 \rightarrow No \\ 1 \rightarrow Yes \end{array}$
Ot	Other or unspecified systemic therapy	F	1	9	 0 → No 1 → Yes, without other specification 2 → Yes, neoadjuvant (preoperative) 3 → Yes, adjuvant (postoperative)
SCT	Stem cell transplantation	F	1	9	0 → No 1 → Yes

⁶ If the vital status is 1 (alive) the *CoD* and *ICD* should be left blank.

⁷ if the vital status is 2 (dead) and the cause of death is unknown, CoD should be coded as R99 (ICD-10)/7999 (ICD-9) or 9999 and ICD should be coded as 99.

⁸ If TNM is not available or not applicable, cTNM (cT, cN, cM) and pTNM (cT, cN, cM) should be coded as 9 and be left blank respectively

and (if possible) *Staging system (ToS)* and *stage* should be coded.

⁹ If cTNM is available and the primary tumour was not resected the pTNM (*pT, pN, pM*) should be left blank.

¹⁰ If available, type of surgery (*local surgery* [12] versus *operative surgery* [13]) should be coded for solid cancers of the following cancer sites: C01-C06, C16-C20, C30-C34, C53-C55, C61 and C65-C68. For other cancers, code 1 (surgery without specification) suffices.

¹¹ If both *local surgery* and *operative surgery* were performed for the same tumour, *operative surgery* should be coded.

¹² The following procedures should be coded as local surgery: polypectomy (mainly gastro-intestinal tract), transurethral resection (TUR; bladder & other urinary tract), cone biopsy/loop excision (cervix), as well as all other procedures which leave the organ in situ, such as cryosurgery, laser coagulation, thermoablation, radiofrequency ablation (RFA), etc.

- ¹³ This includes all resections of the tumor which require the removal of an organ or a major part of that organ, such as a lobectomy, hemicolectomy, hysterectomy, cystectomy, prostatectomy, etc.
- ¹⁴ Targeted therapy comprises all drugs that block the growth of cancer cells by inhibition of certain pathways in the cancer cell. Traditional chemotherapy also affects other cells in the body that divide quickly. The main categories of targeted therapy are small molecules (mostly tyrosine kinase inhibitors such as imati<u>nib</u> and many other -<u>nib</u>s) and monoclonal antibodies (such as rituxi<u>mab</u> and many other -<u>mabs</u>). Monoclonal antibodies are considered a form of immunotherapy but should be coded as targeted therapy.

3.1.2 Incidence file creation

As a first step, the file header needs to be created with 39 variables, as per ECIS data call protocol. The incidence file has a fixed structure as for names, variables' order and semicolon (;) as column separators. As a consequence, the header line should be the following:

PAT; MoB; YoB; Age; Sex; Geo_Code; Geo_Label; TUM; MoI; YoI; BoD; Topo; Morpho; Beh; Grade; Autopsy; Vit_stat; MoF; YoF; Surv_time; ICD; CoD; TNM_ed; cT; cN; cM; pT; pN; pM; ToS; Stage; Surgery; Rt; Cht; Tt; It; Ht; Ot; SCT

Please note: do NOT put a semicolon at the end of the line. The line ends in "SCT" and <u>not</u> in "SCT;".

After the creation of the header line, you can populate your incidence file with records (rows) corresponding to incidence cases each. When you finish populating, please save it in csv or txt format and this will be your incidence file ready to be run by the QCS.

The paragraph below explains how to use the CSV Data layout converter tool if you need assistance in preparing or checking the incidence file format before running the QCS.

3.1.3 JRC CSV Data layout converter

The JRC CSV Data layout converter assist users (with Windows operating systems) in the creation of incidence files to be run by the QCS.

To use the tool, please select a data file to import and convert into the proper format. The file can be in any text format (CSV, TXT) with allowed supported columns separators as follows:

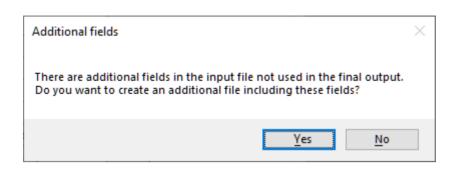
- **TAB** (tabulation)
- | (pipe)
- , (comma)
 - ; (semicolon).If the incidence file is formatted correctly, the converter tool maps the variables as defined in the protocol (Required protocol fields) with the corresponding variables of the input data file (Available fields).To facilitate the data import, the converter tool tries to automatically map fields with the same name: mapped variables are displayed in GREEN, while unmapped variables are marked in RED.

When automatic mapping is not possible, the user can:

- either map manually the variables;
- or leave the field blank (for example, if no mapping is possible, data are not available, etc.).

Only when all the fields defined in the protocol are mapped (or blank), it is possible to Export the content of the original file in the format required by the ECIS protocol.

For variables in the data file not relevant to the ECIS protocol, the tool enables exporting an additional file including one or more of these variables in addition to those required by the ECIS protocol.



In the example above, variables "Extra1", "Extra2" and "Extra3" are not relevant for the ECIS protocol. Among these, "Extra3" (marked as to be included) will be kept in the export file.

	Europea Commis	un Ouality Ch	V Data layout cc eck Software - Protocol d				
col: ENCR	Protocol	~	Select the file to process: C:\databas	se2.csv			
	Rec	quired protocol fields		Availa	ble fields		
Position	Name	Description	Map to		Additional field		
24	cT	Clinical T	23 - cT	~		~	
25	cN	Clinical N	24 - cN	~		\sim	
26	сМ	Clinical M	25 - cM	\sim		\sim	
27	рT	Phatological T	26 - pT	~		\sim	
28	рN	Phatological N	27 - pN	~		\sim	
29	рM	Phatological M	28 - pM	~		\sim	
30	ToS	Staging system	29 - ToS	~		\sim	
31	Stage	Stage	30 - Stage	~		~	
32	Surgery	Surgery	31 - Surgery	~		\sim	
33	Rt	Radiotherapy	32 - Rt	~		\sim	
34	Cht	Chemotherapy	33 - Cht	~		~	
35	Tt	Targeted therapy	34 - Tt	~		\sim	
36	lt	Immunotherapy	35 - It	~		~	
37	Ht	Hormone therapy	36 - Ht	~		~	
38	Ot	Other therapy	37 - Ot	~		~	
39	SCT	Stem cell tranplanatation	38 - SCT	~		~	
40	Extra1	Additional field Extra1	39 - Extra 1	~	<< ignore >>	~	
41	Extra2	Additional field Extra2	40 - Extra2	~	<< ignore >>	~	
42	Extra3	Additional field Extra3	41 - Extra3	~		~	

3.2 Mortality file

Similarly to incidence files, the creation of the file header is the first step. For mortality files, the number of required variables is 5.

Depending on the reporting of deaths by single years of age (Age unit) or by age range (Age range), the line header for the mortality file is the following:

Calendar_Year;Sex;Age unit;Cause of death;Number of deaths

Calendar_Year;Sex;Age range;Cause of death;Number of deaths

Please note: make sure that the 5 variables are in the correct order, and separated by semicolons (;). The header line is mandatory. Do NOT put a semicolon at the end of header.

After having created the header, you can populate your mortality file. Once done, please save it in csv or txt format. You are now ready to run the mortality file by the QCS.

3.3 Population file

Please create first the header of the file. For population files, the number of required variables is 6.

Depending on the reporting of deaths by single years of age (Age unit) or by age range (Age range), the line header for the population file is the following:

Calendar Year;Sex;Age unit;Geo_code;Geo_label;Number of residents

Calendar Year;Sex;Age range;Geo_code; Geo_label;Number of residents

Please note: Please make sure that the 6 variables are in the correct order, and separated by semicolons (;). The header line is mandatory. Do NOT put a semicolon at the end of the header.

After having created the header, you can populate your population file. Once done, please save it in csv or txt format. You are now ready to run the population file by the QCS.

3.4 Life Table file

Please create first the header of the file. For life table files, the number of required variables is 6.

The line header for the life table file is the following:

Calendar Year;Sex;Annual age;Geo_code; Geo_label;All causes death probability

Please note: Please make sure that the 6 variables are in the correct order, and are separated by semicolons. The header line is mandatory. Do NOT put a semicolon (;) at the end of each line.

After having created the header, you can populate your life table file. Once done, please save it in csv or txt format. You are now ready to run the population file by the QCS.

4 How does the software work?

The analysis process of an input *incidence* file is described below. Similar processes are performed for the other input data files: *mortality*, *population* and *life table* files.

The incidence file should include 39 variables, semicolon-separated (;), and in the correct format as described in sections 3.1.1 and 3.1.2 above.

The software checks that variable names are correct, and every single record is compliant with the valid format and value for each variable according to the ECIS protocol as for:

- the number of variables;
- the presence of non-missing and non-blank values in the fields affecting incidence calculation (if year of incidence (variable *YoI*) is missing, the software rises the error E-MISS etc.);
- the field content against a list of valid values, when applicable. **Example**: patient's sex (variable "Sex") is allowed only four numeric values: 1=male, 2=female, 3=other or 9=unknown. Every other value will produce an error (sex=4 will produce E-OUTR, value out of range; sex=X or 11 will produce E-FORM, format error, etc.);
- the field length, which must be within the allowed range. **Example**: the maximum length for Patient identification code (variable PAT) is 50 characters; the maximum length for Morphology (variable *Morpho*) is 4 characters (*Morpho*=82633 will produce E-FORM, format error, etc.);
- the validity of dates, including checking that dates are not set in the future. **Example**: YoI=2028 will produce E-OUTR, value out of range, being the current year the maximum allowed value;
- records failing the edits described in the 2023 JRC Technical Report.

All warning and error messages produced by the QCS in each validation run are saved in three specific output files (as an example, the names below are relative to the *incidence* file):

- 1) *QCS-Incidence-Output.pdf* contains a summary of the execution process (date and time, name of the processed file, number of rows, total number of errors and warnings) and the detailed list of all errors and warnings detected for each record of the input data file;
- 2) *QCS-Incidence-Output.txt* has the same content of the PDF file, but in text format;
- 3) QCS-Incidence-Output.csv contains all errors and warnings in a format easily readable by automated procedures. This file can be used to load the results of the validation process in a database, or to perform detailed statistics and analyses. It can be imported by most software packages to allow for advanced data manipulation, such as linkage with the original file using the unique combination of patient/ tumour ID. Warnings for multiple primary tumours are also included in this file. This format can be particularly useful if the input file generates a large number of errors.

5 Using the software

5.1 Running the software

- Please navigate to the folder in which you installed the software;
- Double click on the JRC-ENCR-QCS.bat file to launch the software (in case of any issue, it is possible to try running the QCS with 2GB of RAM memory by launching file JRC-ENCR-QCS-2GB.bat); please note that the software runs only double clicking on the file ending in .bat;

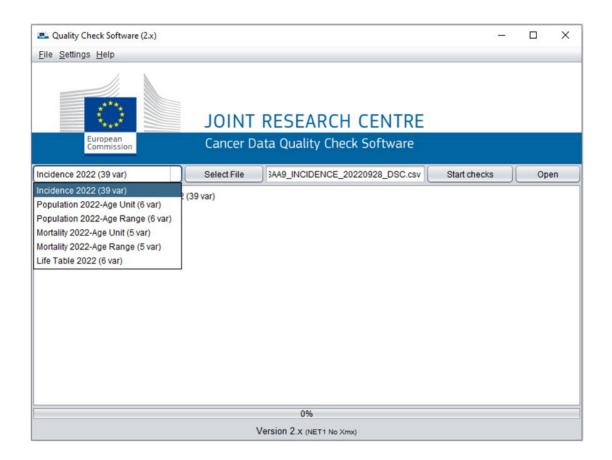
The first time the application is launched a warning window opens for acceptance of the software license. If the license is not accepted, the application will close. Launch the application again and accept the license to run the software.

🖭 Quality (License	×	
File Setting		_	
	STANDARD END-USER LICENCE AGREEMENT	1	
	JRC-ENCR-QCS (2.1.1 - EXECUTABLE) By Using the Software, the User accepts to be bound by the following terms and conditions:		
	The European Union (hereinafter "the Licensor") is the owner of the copyright and other intellectual and industrial property rights and know-how related to the "JRC-ENCR Cancer Registries Data Quality Check		
	Software" (hereinafter "the Software" or "JRC-ENCR-QCS") over which it has the power of disposal		
Incidence 20	regardless of geographical or other limitations. JRC-ENCR-QCS is developed by the Joint Research Centre of the European Commission for the purpose of facilitating data checking and quality controls by		Open
You selecte	European cancer registries, within the framework of the European Commission's and the European Network of Cancer Registries' joint efforts towards simplifying and improving the collection, reliability.		
	consistency, quality and comparability of cancer data in Europe.		
	1. DEFINITIONS		
	"Computer" means an electronic device that is capable of accepting, accessing, transmitting, processing or executing information, instructions or other input in digital or similar form. "Effective Date" means the date on which the Software was received by the User from the Licensor for		
	installation on a Computer. "Software" means the executable version of the software titled JRC-ENCR Cancer Registries Data		
	Quality Check Software, as well as any updated versions thereof, including the contents of all computer		
	files transmitted to the User (provided either by electronic download, on physical media or any other method of distribution) and any accompanying supporting or supplementary material, such as manuals,		
	release notes, specifications and other bundled files or documentation.		
	"Secondary Platform" means a software product owned or operated legally by the User, that is composed of a distinct and self-standing source code, to which one or more separable and self-standing modules		
	or components can be added or integrated, in order to deliver additional functionalities which supplement the core functionality of that software product.	v	
	I have read and understood the above license terms and I agree with them	e	

• To quit the QCS just close the window or select the "Exit" item in File menu.

5.2 Checking the files

Select the type of file you want to check from the drop down menu as shown below.

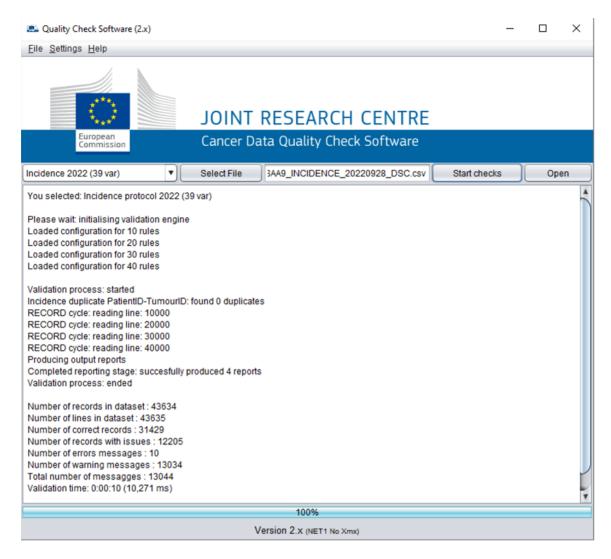


For checking the incidence file according to the ECIS data call protocol:

- Select the "Incidence 2022 (39 var)" option from the drop down menu;
- Press the "Select File" button;
- A file browsing window appears;
- Navigate to the folder where the incidence file to be checked is located, select it (only files with extension csv or txt are accepted) and press "*Open*";
- The full path of the file you have chosen will be displayed in the text box on the left of the "*Start Checks*" button;
- Press the "Start Checks" button to start the validation process.

Please note that in each validation run the output files **will be overwritten**. To keep different QCS output files, please save in different folders or with different names.

While the validation process is running, the number of checked records keeps updating in the display text box:



Once the validation ends, the output window displays a short summary report about the completed process, including the total number of checked records and the total number of errors and warning messages.

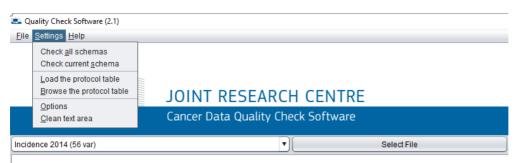
You can directly access the output files by selecting the "*Open*" tab in the user window, to open the "*incidence*" folder containing four reports (CSV, PDF, TXT). This folder is accessible in the "output" folder created at the end of the validation process (see screenshot below).

📕 İ 💆 📑 🖛 i U:\1b-Cancer	Information\QCS\jrc-qcs-2.x\output\incidence			
File Home Share	View			~
Pinto Quick Copy Paste	tut topy path aste shortcut	New Properties	Open Select a Select a Select n History en Select Select	one election
$\leftarrow \rightarrow \cdot \uparrow $ « QCS =	jrc-qcs-2.x > output > incidence	Search incidence		م
This PC	^ Name	Date modified	Туре	Size
3D Objects	QCS-Incidence-Output.csv	23/01/2024 16:35	Microsoft Excel C	2531 KB
📃 Desktop	QCS-Incidence-Output.pdf	23/01/2024 16:36	Adobe Acrobat D	2216 KB
Documents	QCS-Incidence-Output.txt	23/01/2024 16:35	Text Document	20027 KB
Downloads	QCS-Incidence-Output-Summary.txt	23/01/2024 16:35	Text Document	7 KB
h Music				
Pictures				
Videos				
🏭 (C:) Local Disk				
🛖 (H:) HomeDrive				
🛖 (O:) Common				
🛖 (P:) Public				
🛖 (S:) JRC Encrypted				
🛖 (U:) JRC.F.1				

Similarly, *mortality*, *population* and *life table* files can be checked by selecting the corresponding file types and related protocol from the drop down menu. The procedure for checking such files is the same as described above for incidence files.

5.2.1 Settings and options

The "Settings" menu enables to select additional QCS functionalities.

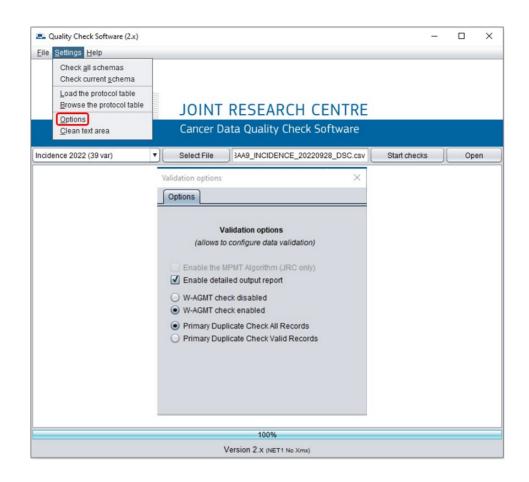


The following settings are available:

- *Check all schemas/Check current schema*. This functionality checks the existence of configuration files, the integrity of single files, the integrity of configuration files and returns the integrity status of either all schemas or the current schema;
- Load the protocol tables/Browse the protocol table. This functionality allows to load or browse the protocol table, listing all the protocol rules applied according to the selected protocol (see screenshot below);

<u>C</u> lean text area		ancer Data Quality Check Software											
dence 2020 (39 var)	Load	ded protocol Incidence 2020 (39 var): 41 m	ales (including function rules)								-		Open
selected: Incidence protocol 2	NICE	NAME	CLASS	RID	CODE	SID	TYPE (LEVEL)	SCOPE	CASE	TARGET	CYCLE	BLOCK	
	1	File size (not empty)	FileSizeRule	20	C-SIZE	11	FILE_FORMAT (0)	file_for	insensit	ALL_FIEL	PRE_REC	blocking 🔺	
	2	Incidence duplicate PatientID-Tum	BunchDuplicatesRule	31	E-DUPL	11	DUPLICATES (0)	all_rec	insensit	SOME_FI	PRE_REC		
	3	Incidence header (positional)	PositionalHeaderRule	40	E-HEAD	11	HEADER (1)	header	insensit	ALL_FIEL	PRE_REC		
	4	Record format (number of fields)	RecordFormatRule	50	E-RECO	11	RECORD FOR	record	insensit	ALL FIEL	RECORD	blocking	
	4	Fields format (datatype)	DataTypeRule	60	E-FORM	11	FIELD_FORMAT	record	insensit	ALL_FIEL	RECORD		
	4	Fields format (mandatory)	MandatoryRule	65		11	FIELD_FORMAT			ALL FIEL	RECORD		
	4	Fields format (max size)	MaxSizeRule	66	E-FORM	11	FIELD FORMAT	record	insensit	ALL FIEL	RECORD		
	4	Fields format (unknown value)	UnknownValueRule	67	W-UNKN	11	FIELD FORMAT	record	insensit	ALL_FIEL	RECORD		
	4	Fields format (important)	ImportantRule	68	W-MISS	11	FIELD FORMAT	record		ALL FIEL	RECORD		
	5	Range check	RangeRule	80	E-OUTR	11	RANGE (4)	record	field le	ALL FIEL	RECORD		
	6	TNM Edition not valid	TNMEditionRule	305	W-TNME	11	CROSS FIELD (record	insensit	SOME FI	RECORD		
	7	Age birth incidence 3	AgeBirthIncidenceRule3	300	E-AGED	11	CROSS FIELD (record	insensit	SOME FI	RECORD		
	7	Age birth incidence 4	AgeBirthIncidenceRule4	301	E-AGEC		CROSS_FIELD (insensit				
	7	Get age from month and year	AgeByMonthYear	104	W-NOMS	11	FUNCTION (5)	record	insensit				
	7	Age Tumour H2	AgeTumourRuleH 2	306	W-AGMT	11	CROSS FIELD (record	insensit	SOME FI	RECORD		
	7	Behaviour Morphology	BehaviourMorphologyRule	204	W-MOBE	11	CROSS FIELD (insensit				
	7	Date birth incidence 2	DateBirthIncidenceRule2	302	E-CoDA	11	CROSS_FIELD (insensit				
	7	Date of last vital status 2	DateLastVitalStatusRule2	304	E-CoDV	11	CROSS FIELD (insensit	SOME FI	RECORD		
	7	Diagnosis morphology 4	DiagnosisMorphologyRule4	308	W-BDMO	11	CROSS FIELD (record	insensit	SOME FI	RECORD		
	7	Diagnosis morphology 5	DiagnosisMorphologyRule5	309	W-BDMS	11	CROSS FIELD (record	insensit	SOME FI	RECORD		
	7	Disease extent 3	DiseaseExtentRule3	216	W-BDpT	11	CROSS FIELD (record	insensit	SOME FI	RECORD		
	7	Disease extent 4	DiseaseExtentRule4	217	W-BDpN	11	CROSS FIELD (record	insensit	SOME FI	RECORD		
	7	Disease extent 5	DiseaseExtentRule5	218	W-BDpM	11	CROSS FIELD (insensit				
	7	Disease extent 10	DiseaseExtentRule10	240	W-BTNM	11	CROSS FIELD (record	insensit	SOME FI	RECORD		
	7	Edition and COD (cause of death)	EditionCauseOfDeathRule	223	E-ECOD	11	CROSS FIELD (insensit				
	7	Grade and Behaviour	GradeBehaviourRule	224	W-BEGR	11	CROSS_FIELD (insensit				
	7	Grade and Morphology	GradeMorphologyRule	225	W-MOGR	11	CROSS_FIELD (record	insensit	SOME_FI	RECORD		
	7	Morphology and topography	MorphologyTopographyRule	226	W-MOTO	11	CROSS_FIELD (record	insensit	SOME_FI	RECORD		
	7	Sex Morphology	SexMorphologyRule	228	W-SEMO	11	CROSS_FIELD (record	insensit	SOME_FI	RECORD		
	7	Sex Topography	SexTopographyRule	229	E-SETO	11	CROSS_FIELD (record	insensit	SOME_FI	RECORD		
	7	TNM standard rule (Stage)	TNMStageRule	238	W-TNMS	11	CROSS FIELD (record	insensit	SOME FI	RECORD	*	

• *Options*. When selected, all validation options are shown. Tick/untick the box *Enable detailed output report* allows the creation of either a detailed or aggregated report. A detailed report is created as the default option.



• *Primary Duplicate Check All Records/Valid Records*. These options allow to have different conditions for checking multiple primary tumours:

With *Primary Duplicate Check All Records* this check is performed on all records except those with errors E-MISS (Value missing), E-FORM (Format error) and E-OUTR (Value out of range) on the morphology variable (Morpho).

The option *Primary Duplicate Check Valid Records* is more selective because multiple primary tumours checks are performed only on records without the following errors/warnings: E-SETO (Topography + Sex not valid), E-AGED (DoI - DoB different from Age), E-AGEC (Age is invalid + impossible to calculate age from DoI – DoB), E-CoDA (DoB + DoI not coherent), W-AGMT (Unlikely Age + tumour type), W-MOTO (Morphology + Topography not valid) and errors as E-MISS, E-FORM and E-OUTR involving the variables topography (Topo) and morphology (Morpho) (see Annex 3 for the definition of error and warning codes)

• *Clear text area* in the *Settings* menu deletes all text from the dialog box.

5.2.2 Help menu

This functionality includes a link to the folder with information on the QCS, a contact e-mail and the link to the QCS page on the ENCR website.

🛃 Help 🛛 🕹
ENCR QCS Help
Please find below a button leading to the folder with the relevant documents for the QCS. You can look for updates at the address: https://encr.eu/tools-for-registries
In case of questions, you are welcome to contact us writing at JRC-ENCR@ec.europa.eu

In the "*Help*" menu you can also find the "*About*" item, with copyright statement, the license and the list of *jar* libraries.

💼 About	×											
JRC-ENCR Qua	ality Check Software											
Developed by the European Commission Joint Research Centre. Directorate F - Health, Consumers and Reference Materials. Disease Prevention Unit.												
In support of the	European Network of Cancer Registries (ENCR).											
Version: Version Build: 2024-01-26-												
Copyright 2016-	2024 European Union											
	License											
	License Libraries ×											

5.3 Output files

Each validation run generates four output files in PDF, TXT and CSV format. The output files are located in the related subfolders for *incidence*, *population*, *mortality* or *life table*, inside the "*output*" folder. For example, output files for incidence are located in the subfolder "**\JRC- ENCR-QCS-V2.x\output\incidence**".

5.3.1 Details of pdf/txt output file

The file "QCS-Incidence-Output-Summary.txt" contains only a summary of the execution process (date and time, name of the processed file, number of rows, total number of errors and warnings). The other pdf/txt files include also the detailed list of all errors and warnings detected for each record of the input data file.

The following screenshots refer to the different sections of the *QCS-Incidence-Output.txt* (or *pdf*) file.

Detail of upper section

QUALITY CHECK SOFTWARE REPORT - INCIDENCE PROCESSING PARAMETERS File process start : 2023-10-30 11:0:47.748 File process end : 2023-10-30 11:0:54.749 Validated by : OCS Version 2.1 (run 1698660055579) File Processed: U:\1b-Cancer Information\QCS\jrc-qcs-2.1\Test.csv (CC89315B5D) Validation options: : 11 : Incidence 2020 (39 var) Validation protocol ID Name of validation protocol Print detailed TXT report : true Use reference data : true Strip all quotes from dataset : false : true Print cross-record matching info ENCR Incidence options: Acceptance MP criterion Age-Tumour rule (W-AGMT) Apply MPMT merging algorithm Acceptance MP criterion : 2 : hybrid 2023 (obi 558) : false PROCESSING STATISTICS Number of records read : 25573 Total number of errors : 246 Total number of warnings : 7273 Total number of records with issues : 6552 Total number of rejected records : 0 Preliminary rules : nothing critical KEY TO ERROR AND WARNING CODES E-AGEC: Age is invalid + impossible to calculate age from DoI - DoB E-AGED: DoI - DoB different from Age E-AUVS: Autopsy+Vital status+Surv time+Date of Incidence/Date of Followup not valid (Tab. 12) E-BDVS: BoD+Vital status+Surv time+Date of Inidence/Date of Followup not valid (Tab. 11) E-CoDA: DoB + DoI not coherent (p.16) E-CoDV: Date of last known vital status not valid

Processing parameters:

- File process start, File process end;
- Validated by: The QCS version used to produce the report;
- *File processed*. The name and the path of the file checked by the software.

Processing statistics:

- Number of records read, Total numbers of errors, Total number of warnings;
- Total number of records with issues: each record can have more errors and/or warnings;
- *Total number of records rejected*: records are rejected whenever the header is correct but some of the variables are not present, not even left blank or with missing value.

Key to error and warning codes:

 Errors and warnings are referenced by codes, described by short labels and accompanied by the reference to the relevant table or page from the JRC Technical Report "A common data quality check procedure for European cancer registries". See also Annex 3 – List of error and warning codes.

Detail of second part (summary of errors and warnings)

SUMMARY OF WARNINGS BY CODE	7 239
E-AUVS E-BDVS SUMMARY OF WARNINGS BY CODE W-AGMT	239
E-BDVS SUMMARY OF WARNINGS BY CODE	*************
SUMMARY OF WARNINGS BY CODE	**********
w-AGMT	
W-AGMT	
	20
W-BDMO	236
W-BDMS	43
W-BDpM	1415
W-BDpN	123
W-BDpT	140
	282
	45
W-MOGR	718
W-MOTO W-MPCR	102
W-MPCK	20
W-TBGR	45
W-TNMM	511
W-TNMS	3555

<u>Summary of errors warnings by code</u>: see Annex3 – List of error and warning codes.

Detail of Multiple Primary Tumours checks: W-MPCR and W-MPMT sections

The QCS 2.x performs MPMTs quality checks for computing incidence according to the 2004 International Rules for Multiple Primary Cancers, also referenced in the JRC Technical Report (Chapter 4, p. 34-39).

In addition to checks on multiple primary tumours with behavior 3 (malignant), the QCS 2.x performs also checks on tumours with behavior <3 using new and more precise filtering criteria. For example, multiple primary tumours with topography codes C50 or C65-C67 and behaviour 2 (in situ), or tumours with codes C70-C72, C751-C753 and behaviour 0 (benign) or 1 (uncertain) and specific morphologies, are also checked.

These filtering criteria are detailed in the configuration file located in the QCS 2.x subfolder "...\jrc-qcs-2.x\config\rules\encr2014\PrimaryMultipleFilter.csv".

MULTIPLE PRIMARY MALIGNANT TUMOUR CHECK (W-MPCR) lum 572375 Tum 572375 PAT 234281 Duplicated record: -----BoD Topo Morpho Beh Sex DoI Dob DoB 2 C619 8000 3 1 12/2008 9/1924 -----PAT 234281 Tum 572375 Duplicated record: Tum 572373 _____ BoD Topo Morpho Beh Sex DoI DoB -----..... 2 (343 8000 3 1 1/2012 9/1924 ----------MULTIPLE PRIMARY MALIGNANT TUMOUR CHECK (W-MPMT) _____ Tum 165025 Tum 426753 PAT 135639 Duplicated record: ----------BoD Topo Morpho Beh Sex DoI DoB 3 2 2 4/2007 7 C509 8503 8/1936 ------_____ Tum -25 Tum 165025 PAT 135639 Duplicated record: BoD Topo Morpho Beh Sex DoI DoB 7 C509 8480 3 2 9/2008 8/1936 _____

For each MPMT warning the following variables are reported: PAT, Tum, BoD (basis of diagnosis), Topo (topography), Morpho (morphology), Beh (behaviour), Sex, DoI (date of incidence), DoB (date of birth).

The two MPMT output sections list all multiple primary tumours detected by the QCS 2.1 and reports two types of warnings, **W-MPCR** or **W-MPMT**, according to some special conditions defined for subgroups of topography/morphology codes.

- The warning **W-MPMT** applies, under certain conditions, only to subgroups of multiple primary solid tumours with specific topography codes:
 - tumours diagnosed in paired organs (e.g. C50, C62, C69 etc.);
 - tumours which occur in some 4th character subcategory of colon (C18.0-C18.7) and skin (C44.0-C44.7);
 - tumours with different first 3 digits in the site code, belonging to groups of topography codes considered as a single site according to Table 14 of the JRC Technical report 2023 (e.g. C01, C02; C00, C03-C06; C65-C68 etc.).

All equivalent tumours (same topography and morphology) with **W-MPMT** warnings in the

same patient should be counted only once for incidence analysis, according to the 2004 International Rules for Multiple Primary Cancers and the ENCR recommendations as referenced in the JRC Technical report. See some examples at point 6) of Chapter 6.

• The warning **W-MPCR** applies to all tumours not belonging to the subgroups above, e.g. solid tumours with different topography codes such as C61 etc. and all hematological malignancies regardless of topography code.

Equivalent tumours with **W-MPCR** warnings in the same patient should be further checked by the cancer registry to decide if only one should be considered for incidence analysis. See some examples at point 6) of Chapter 6.

Detail on section with errors and warnings

PAT	208387				Tum 24				
BoD	Торо	Morpho					Var_Name		
7	C710	9441	3	1	1/2008	2/1937	Surgery pT	9 x	W-SUpT W-SUpT
РАТ	521016				Tum 65	5821			
BoD	Торо	Morpho	Beh	Sex	DoI	DoB	Var_Name	Var_Value	
7	C492	8920	3			4/2008	Surgery pT		W-SUpT W-SUpT
РАТ	1388134				Tum 15				
BoD	Торо	Morpho	Beh	Sex		DoB		Var_Value	Error_Cod
7	C221	8000			11/2018		Topo Morpho TNM_ed Stage pT pN pM cT cN cN cM 	C221 8000 8 3B × × 0 2 1 0 9 ×	W-TINMM W-TINMM W-TINMM W-TINMM W-TINMM W-TINMM W-TINMM W-TINMM W-TINMM W-TINMM W-SUPT W-SUPT

Errors and warnings: key variables are reported for each error and warning, namely: *PAT*, *Tum*, *Topo* (topography), *Morpho* (morphology), *Beh* (behaviour), Sex, *DoI* (date of incidence), *DoB* (date of birth), *Var_Name* and *Var_Value* (list of relevant variables which caused the warning/error with the corresponding name and value), *Error_Code* (code according to list in *Annex 3 – List of error and warning codes*).

5.3.2 Details of csv output files

The csv output file contains all errors and warnings in a format easily readable by automated procedures, which can be used to perform detailed statistics and analyses.

The following screenshots refer to the *QCS-Incidence-Output.csv* file:

Detail of the left part of the csv file

Line	Patient	Tumour	BoD	Торо	Morpho	Beh	Sex	Dol	DoB
4	1388134	1514686	7	C221	8000	3	2	Nov-18	Feb-49
6	2709	2726	5	C424	9836	3	2	Aug-04	Dec-93
9	2778	2793	5	C424	9836	3	1	Apr-05	Nov-99
14	2972	2975	5	C424	9872	3	1	Sep-05	Sep-02
23	3222	3208	7	C480	9500	3	1	Jun-06	Sep-03
24	3223	3209	5	C424	9874	3	1	Jan-06	Jul-92

Detail of the right part of the csv file

Error_Cod	Error_Description	Var1_Nam	Var1_Valu	Var2_Nam	Var2_Valu	Var3_Nam	Var3_Valu
W-TNMM	Morphology not addressed by the Topograp	Торо	C221	Morpho	8000	TNM_ed	8
W-MOTO	Morphology + Topography not valid (tab.8)	Morpho	9836	Торо	C424		
W-MOTO	Morphology + Topography not valid (tab.8)	Morpho	9836	Торо	C424		
W-MOTO	Morphology + Topography not valid (tab.8)	Morpho	9872	Торо	C424		
W-TNMM	Morphology not addressed by the Topograp	Торо	C480	Morpho	9500	TNM_ed	6
W-MOTO	Morphology + Topography not valid (tab.8)	Morpho	9874	Торо	C424		

6 How to interpret the incidence files' output generated by the QCS

Two types of messages are included in the output reports: W-YYYY (warning code) and E-YYYY (error code), where the code YYYY identifies the specific type of message.

This section describes how to interpret the outcomes of the QCS for some of the variables having an impact on the incidence estimation. Some examples of errors on key variables, of warnings on TNM classification variables and on multiple primary tumours are also reported.

1) Errors due to single variable format and values

• **E-OUTR**: out of range.

When the variables have values different from those allowed by the ECIS *Data Call Protocol*, the QCS returns the error E-OUTR.

PAT	000001				Tum 02				
BoD	Торо	Morpho	Beh	Sex	Dol	DoB	Var_Name	Var_Value	Error_Code
1	C427	9800	3	2	9/2010	2/1924	Торо	C427	E-OUTR

In this example the QCS gives the error E-OUTR on the variable *Topo* because topography C427 does not exist in the International Classification of Diseases for Oncology, third edition¹⁵ (ICD-O-3).

PAT 00	00002				Tum 01				
BoD	Торо	Morpho	Beh	Sex	Dol	DoB	Var_Name	Var_Value	Error_Code
7	C620	9999	9	1	10/2012	4/1935	Morpho	9999	E-OUTR
							Beh	9	E-OUTR

In this example the QCS returns the error E-OUTR on the variables *Morpho* and *Beh* because morphology 9999 does not exist in the ICD-O-3, and behaviour 9 is not allowed according to the ECIS data call protocol.

• **E-MISS**: missing value.

PAT 0	00003				Tum 01			
BoD	Торо	Morpho	Beh	Sex	DoI	DoB	Var_Name	Var_Value Error_Code
7	C187		3	1	8/2011	3/1945	Morpho	E-MISS

In this example the QCS returns the error E-MISS because the variable *morphology* (impacting on incidence calculations) is missing.

¹⁵ International Classification of Diseases for Oncology, Third Edition, First Revision. Geneva: World Health Organization, 2013

• **E-OUTR**: out of range value.

PAT 0	00002								
BoD	Торо	Morpho	Beh	Sex	DoI	DoB	Var_Name	Var_Value	Error_Code
7	C620	9999	9	1	10/2012	4/1935	Morpho	9999	E-OUTR
							Beh	9	E-OUTR

In this example the QCS returns the error E-OUTR on the variables *Morpho* and *Beh* because morphology 9999 does not exist in the ICD-O-3, and behaviour 9 is not allowed according to the ECIS data call protocol.

• **E-AGEC**: Age is invalid or missing, and it is impossible to calculate.

PAT 0	00004				Tum 01				
BoD	Торо	Morpho	Beh	Sex	DoI	DoB	Var_Name	Var_Value	Error_Code
7	C169	8140	3	2	11/2013	99/9999	Age YoB YoI	999 9999 2013	E-AGEC E-AGEC E-AGEC

In this example the QCS gives the error E-AGEC because the variable Age (impacting on incidence calculations) is unknown (999) and cannot be calculated from DoI (2013) – DoB (9999). All the related variables Age, YoB and YoI are reported with the same error.

• **E-FORM**: format error.

PAT	000005				 Tum 0	1			
BoD	Торо	Morpho	Beh	Sex	DoI	DoB	Var_Name	Var_Value	Error_Code
7	C443	80984	3	1	9/2011	2/1933	Morpho	80984	E-FORM

In this example the QCS gives the error E-FORM on variable *Morpho* (80984) because morphology should have four digits instead of five, according to the ICD-O-3.

2) Errors due to inconsistency of dates.

• **E-CoDA**: date of birth and date of incidence are not consistent.

PAT	000006				Tum 01				
BoD	Торо	Morpho	Beh	Sex	DoI	DoB	Var_Name	Var_Value	Error_Code
7	C159	8140	3	2	12/1992	8/2016	 УоВ	2016	E-CoDA

In this example the QCS gives the error E-CoDA on the variable *YoB* because the year of birth (2016) is set after the year of incidence (1992).

PAT 0	00007				 Tum 0	1			
BoD	Торо	Morpho	Beh	Sex	DoI	DoB	Var_Name	Var_Value	Error_Code
7	C741	9490	3	1	2/1992	3/1992	МоВ	3	E-CoDA

In this example the QCS gives the error E-CoDA because the birth (month 3) occurs after the incidence month (2) of the same year.

• **E-CoDV**: date of the incidence and date of the last known vital status are not consistent.

PAT 0	8000				Tum 01				
BoD	Торо	Morpho	Beh	Sex	Dol	DoB	Var_Name	Var_Value	Error_Code
2	C549	8000	3	2	8/2013	4/1933	MoI YoI MoF YoF	8 2013 8 2012	E-CoDV E-CoDV E-CoDV E-CoDV E-CoDV

In this example the QCS gives the error E-CoDV because the year of incidence (*YoI* 2013) occurs later than the date of last known vital status (*YoF* 2012).

PAT	000009				Tum 01				
BoD	Торо	Morpho	Beh	Sex	DoI	DoB	Var_Name	Var_Value	Error_Code
2	C160	8000	3	1	6/2009	10/1924	MoI YoI MoF YoF	6 2009 4 2009	E-CoDV E-CoDV E-CoDV E-CoDV E-CoDV

In this example the QCS gives the error E-CoDV because the date of incidence (6/2009) occurs later than the date of last known vital status (4/2009).

- 3) Errors and warnings due to tumour and demographic variables combinations.
 - **E-SETO**: sex and topography combinations are not valid.

PAT 0	00010				Tum 01				
BoD	Торо	Morpho	Beh	Sex	Dol	DoB	Var_Name	Var_Value	Error_Code
2	C569	8000	3	1	10/2013	3/1935	Sex Topo	1 C569	E-SETO E-SETO

In this example the QCS returns the error E-SETO because the combination of topography=C569 (ovary) and sex=1 (men) is not valid.

• **W-AGMT**: age and morphology/topography combinations are unlikely.

PAT	000011				Tum 01				
BoD	Торо	Morpho	Beh	Sex	DoI	DoB	Var_Name	Var_Value	Error_Code
7	C424	9652	3	1	12/2003	10/2003	Age Morpho	0 9652	W-AGMT W-AGMT

In this example the QCS gives the warning W-AGMT because the morphology 9652 (Hodgkin lymphoma, mixed cellularity, NOS) is unlikely between ages 0-2 (see Table 2 of the JRC technical report).

PAT 00	00012				Tum 01				
BoD	Торо	Morpho	Beh	Sex	DoI	DoB	Var_Name	Var_Value	Error_Code
7	C619	8140	3	1	3/2007	5/1996	Age Topo Morpho	10 C619 8140	W-AGMT W-AGMT W-AGMT

In this example the QCS gives the warning W-AGMT because the topography C619 (prostate) in combination with morphology 8140/3 (adenocarcinoma, NOS) is unlikely under the age of 35 (see Table 2 of the JRC technical report).

4) Errors and warnings due to combinations of follow-up variables, basis of diagnosis and stage.

Three new rules have been implemented to check consistency between important variables related to survival.

• **E- AUVS (NEW):** autopsy, vital status and survival time combination not valid.

This new rule checks the consistency between the variables autopsy, vital status and survival time, according to Table 12 of the JRC technical report.

PAT 1	0855				Tum 13	439			
BoD	Торо	Morpho	Beh	Sex	Dol	DoB	Var_Name	Var_Value	Error_Code
7	C779	9663	3	2	1/2004	7/1929	Autopsy Vit_stat Surv_time YoI MoI YoF MoF	1 2 221 2004 1 2004 9	E-AUVS E-AUVS E-AUVS E-AUVS E-AUVS E-AUVS E-AUVS E-AUVS

In this example the QCS gives the error W-AUVS because the vital status is set to 2 (dead) but the survival time is not zero.

• **E-BDVS (NEW):** basis of diagnosis, vital status, survival time and stage combination not valid.

This new rule checks consistency between the variables basis of diagnosis, vital status, survival and stage according to Table 11 of the JRC technical report.

PAT	PAT 381477 Tum 426374										
BoD	Торо	Morpho	Beh	Sex	Dol	DoB	Var_Name	Var_Value	Error_Code		
0	C809	8000	3	2	8/2006	1/1917	BoD Vit_stat Surv_time YoI MoI YoF MoF Stage	0 2 0 2006 8 2006 8 4	E-BDVS E-BDVS E-BDVS E-BDVS E-BDVS E-BDVS E-BDVS E-BDVS E-BDVS		

In this example the QCS gives the error W-BDVS because basis of diagnosis set to 0 (DCO), vital status to 2 (dead) and survival time 0 are not coherent with stage 4. Stage should instead be unknown (9) or missing (blank).

• **E-VSBD (NEW):** vital status, basis of diagnosis and autopsy combination not valid.

This new rule checks consistency between the variables vital status, basis of diagnosis and autopsy, according to Table 10 of the JRC technical report.

PAT 3	3				Tum 374	41			
BoD	Торо	Morpho	Beh	Sex	Dol	DoB	Var_Name	Var_Value	Error_Code
0	C509	8500	3	2	12/2004	11/1948	Vit_stat BoD Autopsy	1 0 1	E-VSBD E-VSBD E-VSBD

In this example the QCS gives the error W-VSBD because vital status set to 1 (alive) is not coherent with basis of diagnosis 0 (DCO) and autopsy 1 (incidental finding of cancer at autopsy). Autopsy should not be 1 and Basis of Diagnosis should not be 0.

5) Errors and warnings due to combinations of other tumour variables.

• **W-MOBE**: morphology and behaviour combinations are not included in the ICD-O-3.

According to Rule F of the ICD-O-3 it is exceptionally possible to have a morphology and behaviour combination not listed in the ICD-O-3. Starting from version 2.0 the QCS reports as warnings such combinations, while previous versions reported as errors (E-MOBE). The list of accepted combinations of morphology and behaviour is reported in Table 9 of the JRC technical report 2023; these combinations are now allowed and do not produce W-MOBE warnings.

PAT 0	00013				Tum 01				
BoD	Торо	Morpho	Beh	Sex	Dol	DoB	Var_Name	Var_Value	Error_Code
7	C569	8621	3	2	4/2005	6/1982	Morpho Beh	8621 3	W-MOBE W-MOBE

In this example the QCS gives error the W-MOBE because morphology 8621 (granulosa cell-theca cell tumour) with behaviour 3 (malignant tumour) is not listed in the ICD-O-3. This combination is possible, but unlikely.

PAT 0	00014				Tum 01				
BoD	Торо	Morpho	Beh	Sex	DoI	DoB	Var_Name	Var_Value	Error_Code
7	C421	9950	1	1	12/1989	3/1921	Morpho Beh	9950 1	W-MOBE W-MOBE

In this example the QCS gives the W-MOBE warning because morphology 9950 (polycythaemia vera) has behaviour 3 (malignant tumour) in ICD-O-3. This morphology (polycythaemia vera) changed from borderline tumour (behaviour 1) in ICD-O-2¹⁶ to malignant tumour (behaviour 3) in ICD-O-3.

W-BDMO: morphology too specific according to the basis of diagnosis.
 This rule has been updated according to Table 4 of the JRC technical report.

PAT 3	11683				Tum 72	Tum 729324				
BoD	Торо	Morpho	Beh	Sex	Dol	DoB	Var_Name	Var_Value	Error_Code	
2	C710	9530	0	2	6/2013	7/1936	BoD Morpho Topo Beh	2 9530 C710 0	W-BDMO W-BDMO W-BDMO W-BDMO	

In the example above the QCS returns the warning W-BDMO because the combination of morpho/beh 9530/0 (Meningioma, NOS) with topography C71 (Brain) and basis of diagnosis 2 (clinical) is not allowed.

¹⁶ International Classification of Diseases for Oncology, Second Edition. Geneva: World Health Organization, 1990.

• **W-BDMS**: morphology not specific enough according to the basis of diagnosis. This rule has been updated according to Table 4 of the JRC technical report.

PAT 1	0542				Tum 13	3064			
BoD	Торо	Morpho	Beh	Sex	Dol	DoB	Var_Name	Var_Value	Error_Code
5	C421	9866	3	1	9/2004	3/1936	BoD Morpho Beh	5 9866 3	W-BDMS W-BDMS W-BDMS

In this example the QCS gives the warning W-BDMS because the combination of *morpho/beh* 9866/3 (Acute promyelocytic leukemia,) with basis of diagnosis 5 (cytology) is not allowed.

PAT 3	0407				Tum 33	085			
BoD	Торо	Morpho	Beh	Sex	Dol	DoB	Var_Name	Var_Value	Error_Code
7	C421	9896	3	2	4/2004	9/1953	BoD Morpho Beh	7 9896 3	W-BDMS W-BDMS W-BDMS

In this example the QCS gives the warning W-BDMS because the combination of *morpho/beh* 9896/3 (Acute myeloid leukemia) with basis of diagnosis 7 (histology) is not allowed.

• **W-BTNM**: behaviour and TNM combination not valid.

This rule has been updated according to the additional checks suggested at p. 33 of the JRC technical report.

PAT 0	00021				Tum 01				
BoD	Торо	Morpho	Beh	Sex	Dol	DoB	Var_Name	Var_Value	Error_Code
7	C629	9061	3	1	2/2011	9/1991	Beh pT cT	3 is 9	W-BTNM W-BTNM W-BTNM

In this example the QCS gives the warning W-BTNM because behaviour 3 (malignant tumour) is not coherent with pathological T (pT) set to *is* (carcinoma in situ).

PAT 4	164				Tum 42	09				
BoD	Торо	Morpho	Beh	Sex	Dol	DoB	Var_Name	Var_Value	Error_Code	
7	C445	8720	2	1	10/2004	12/1962	Beh pT cT Stage	2 is X 9	W-BTNM W-BTNM W-BTNM W-BTNM	

In the example above the QCS gives the warning W-BTNM because behavior 2 (in situ) should have stage set to 0.

 W-TBGR (NEW): topography, morphology, behaviour and grade combinations are unlikely.

This new rule follows Table 5 of the JRC technical report, checking the grade of some tumours with behavior <3 for specific morphologies. In addition to the combinations reported in Table 5, the grade of CNS tumours with behaviours 0 and 1 is also checked according to the new ENCR Recommendations for coding tumours of the CNS (to be published).

PAT 5	104				17				
BoD	Торо	Morpho	Beh	Sex	Dol	DoB	Var_Name	Var_Value	Error_Code
7	C679	8130	2	1	2/2004	7/1953	Topo Morpho Beh Grade	C679 8130 2 2	W-TBGR W-TBGR W-TBGR W-TBGR

In this example the QCS gives the warning W-TBGR because topography C679 (Bladder, NOS) with behavior 2 and morphology 8130 should have grade set to 1 or 3.

• **W-MOGR**: morphology, behaviour and grade combinations are unlikely.

This rule has been updated according to Tables 6 and 7 of the JRC technical report.

PAT	000022				Tum 01				
BoD	Торо	Morpho	Beh	Sex	Dol	DoB	Var_Name	Var_Value	Error_Code
7	C569	8620	3	2	5/2012	7/1954	Grade Morpho Beh	5 8620 3	W-MOGR W-MOGR W-MOGR

In the example above the QCS gives the warning W-MOGR because grade 5 (T-cell) is used to denote cell lineage for haematological malignancies (leukaemia and lymphoma), while morphology 8620 (granulosa cell tumour, malignant) is not an hematological malignancy. In general, the combination between grades 5-8 and morphology out of the range 9590-9993 is impossible, and produces the warning W-MOGR, except for 9801/34.

PAT	000023				Tum 01				
BoD	Торо	Morpho	Beh	Sex	Dol	DoB	Var_Name	Var_Value	Error_Code
7	C445	9709	3	1	11/2013	4/1935	Grade Morpho Beh	6 9709 3	W-MOGR W-MOGR W-MOGR

In this example, the QCS gives the warning W-MOGR because morphology 9709 (Cutaneous T-cell lymphoma, NOS) should have grade 5 (T-cell) instead of 6.

• **W-MOTO**: morphology and topography combinations are unlikely.

This rule has been updated according to Tables 8 of the JRC technical report.

PAT	000024				Tum 01				
BoD	Торо	Morpho	Beh	Sex	DoI	DoB	Var_Name	Var_Value	Error_Code
7	C779	8070	3	1	12/2008	10/1946	Morpho Topo	8070 C779	W-MOTO W-MOTO

The QCS gives the warning W-MOTO because the combination of topography C779 (Lymph node, NOS) and morphology 8070 (squamous cell carcinoma, NOS) is not allowed; this combination is probably a metastasis and the topography should be coded as C809.

PAT	000025				Tum 01				
BoD	Торо	Morpho	Beh	Sex	Dol	DoB	Var_Name	Var_Value Error_Cod	le
7	C539	8120	3	2	11/2007	9/1959	Morpho Topo	8120 W-MOTO C539 W-MOTO	

In the example above the QCS gives the warning W-MOTO because the combination of topography C539 (cervix uteri) and morphology 8120 (transitional cell carcinoma, NOS) is very rare.

• **W-TNMM**: TNM and stage are present, but morphology is not included in the TNM.

The groups of morphology codes used in the TNM classification rules have been updated to TNM edition 6, 7 and 8, according to Table 15 of the JRC technical report.

PAT 000026 Tum 01									
BoD	Торо	Morpho	Beh	Sex	DoI	DoB	Var_Name	Var_Value	Error_Code
7	C505	9120	3	2	11/2007	3/1971	Торо	C505	W-TNMM
							Morpho	9120	W-TNMM
							TNM_ed	6	W-TNMM
							Stage	IIB	W-TNMM
							рт	3	W-TNMM
							pN	0	W-TNMM
							рМ	0	W-TNMM
							cT	9	W-TNMM
							cN	9	W-TNMM
							сМ	9	W-TNMM

In the example above the QCS returns the warning W-TNMM because the tumour is a breast angiosarcoma (morphology 9120) with stage IIB. When topography is set to C50 (breast) only carcinomas should be staged.

• **W-TNMS**: TNM and stage are not consistent.

PAT 0	00027				Tum 01				
BoD	Торо	Morpho	Beh	Sex	DoI	DoB	Var_Name	Var_Value	Error_Code
7	C502	8140	3	2	8/2013	6/1965	Topo Morpho TNM_ed Stage pT pN pM cT cN cC cM Grade Age Beh	C502 8140 7 IIIA 3 1 9 9 9 9 3 48 3	W-TNMS W-TNMS W-TNMS W-TNMS W-TNMS W-TNMS W-TNMS W-TNMS W-TNMS W-TNMS W-TNMS W-TNMS W-TNMS W-TNMS W-TNMS

In the example above the QCS returns the warning W-TNMS because the tumour is a breast carcinoma with pT=3, pN=1, pM=1 and Stage=IIIA. This combination is not consistent, according to TNM Ed.7 *Stage* should be IV.

6) W-MPMT and W-MPCR: Warnings for multiple primary tumours.

The MPMTs checks performed by the QCS produce two types of warnings, W-MPMT and W-MPCR, according to some special conditions for topography/morphology codes as explained at § 5.3.1 above.

If a patient has two or more multiple primary tumours that produce a warning **W-MPMT**, only one tumour should be considered for incidence analysis according to 2004 International Rules for Multiple Primary cancer.

If a patient has two or more multiple primary tumours that produce a warning **W-MPCR**, probably the multiple records refer to the same tumour, therefore these records should be reviewed by CRs.

Below some examples:

PAT	00002	8				Tum 01
BoD	Торо	Morpho	Beh	Sex	Dol	DoB
7	C717	8000	3	2	12/2016	12/1954
PAT	00002	8				Tum 02
BoD	Торо	Morpho	Beh	Sex	Dol	DoB
7	C717	9590	3	2	11/2016	12/1954

In example above, the QCS gives the warning W-MPCR because probably the two records refer to the same tumour (unspecific morphology 8000, topography code C71).

PAT	00002	9				Tum	01
BoD	Торо	Morpho	Beh	Sex	DoI	DoB	
7	C679	8130	3	2	5/2003	1/1	930
PAT	00002	9				Tum	02
BoD	Торо	Morpho	Beh	Sex	DOI	DoB	
7	C809	8000	3	2	11/2010	1/1	930

In this example, the QCS gives the warning W-MPCR because the two records could refer to the same tumour (unspecific morphology 8000, unknown primary site C809).

PAT	00003	0				Tum 01
BoD	Торо	Morpho	Beh	Sex	Dol	DoB
7	C501	8500	3	2	1/2001	7/1960
PAT	00003	0				Tum 02
BoD	Торо	Morpho	Beh	Sex	DOI	DoB
7	C508	8520	3	2	10/2002	7/1960

In example above, the QCS gives the warning W-MPMT because according to the 2004 International Rules, the two records could refer to the same tumour of different laterality with same morphology, diagnosed in paired organs (Table 15 of the JRC Technical report). In this case only one tumour should be considered for incidence analysis (the first to be diagnosed in 2001).

PAT 1141980 Duplicated record:		Tum 1230495 Tum 1567154
BoD Topo Morpho	Beh Sex Dol	DoB
7 C182 8140	3 1 12/201	7 4/1960
PAT 1141980 Duplicated record:		Tum 1567154 Tum 1230495
BoD Topo Morpho	Beh Sex Dol	DoB
7 C184 8140	3 1 12/201	8 4/1960

In this example, the QCS gives the warning W-MPMT because according to the 2004 International Rules, the two topographies are the same (single site C18) and the two morphologies are in the same group (single entity).

In this case, only one tumour should be considered for incidence analysis (the first to be diagnosed in 2017).

Annex 1 – Major changes and improvements

Major updates and new features available from version 2.1 onwards are listed below (please refer to the change log file "change_log.txt" in the doc folder for more details):

GUI:

- A reminder to accept the user license has been added the first time the software is launched;
- Additional messages on the status of the validation process have been added;
- A short summary on the results of the validation process has been added at the end of each run;
- A new option "Remove all quotes from the dataset" has been added in the Settings/Options menu to help users in cleaning the dataset.

Output Reports:

- Details on the set of "Validation options" used in the last run have been added.
- A new TXT output file with a summary of the validation output (from version 2.1.1)

Code and configuration files:

- Upgrade of Log4j to version 2.17.1 to mitigate vulnerability risk;
- Better usage of the computer's memory during validation;

Univariate/multivariate checks have been updated according to the latest version of the JRC Technical report:

- Update of univariate checks, compliant to Table 1;
- Update of consistency checks between morphology and topography (rule W-MOTO) according to the updated Table 8;
- Update of consistency checks between morphology and grade of CNS tumours with Beh=3 (rule W-MOGR), according to the new ENCR Recommendations for coding tumours of the CNS (to be published);
- Update of consistency checks between morphology and grade of CNS tumours with Beh<3 (rule W-TBGR), according to the new ENCR Recommendations for coding tumours of the CNS (to be published);
- Update of consistency checks between Surgery, Basis of Diagnosis (BoD), Autopsy and the variable pT, according to the updated rules on p.33 of the JRC report;
- Update of consistency checks between age and tumour type (rule W-AGMT) according to the updated Table 2;
- Update of consistency checks between basis of diagnosis, morphology and behaviour by topography (rules W-BDMO, W-BDMS), according to Table 4 and to the 2022 ENCR Recommendations on Basis of Diagnosis;
- **NEW**: update of consistency checks between morphology and grade compliant to Table 5 and to the WHO grading system, table 27 of the ICD-O-3.1 (new rule **W-TBGR**) for CNS tumours;

- Update of consistency checks between morphology and grade compliant to Table 6 and 7 of the JRC Technical report for rule W-MOGR;
- Update of consistency checks between morphology and topography codes (rule W-MOTO) according to Table 8;
- Selected combinations of morphology/behaviour not included in any version of the ICD-O-3 are now accepted, according to Table 9;
- NEW: more precise consistency checks for follow-up variables, basis of diagnosis and TNM stage, according to Tables 10, 11 and 12 (new rules **E-VSBD**, **E-BDVS** and **E-AUVS** replacing the rule E-VBDA in previous versions of the QCS);
- **NEW**: multivariate checks for treatment variables, basis of diagnosis and TNM stage implemented according to rules reported at p.33 of the latest version of the JRC report (new rules **W-SUBD** and **W-SUpT**);
- **NEW:** update of consistency checks between behaviour and TNM/stage (rule **W-BTNM** now checking also the stage, see p.33 of the latest version of the JRC report);
- Update of consistency checks between TNM/stage and basis of diagnosis (rules W-BDpT, W-BDpN and W-BDpM, see p.33 of the latest version of the JRC report);
- Update of groups of morphology codes used to check consistency between TNM variables and stage according to TNM editions 6,7 and 8 (rules **W-TNMM** and **W-TNMS**);
- Update of NUTS codes for the geographical variables Geo_code according to the EUROSTAT Nuts 2021 classification (<u>https://ec.europa.eu/eurostat/web/nuts/background</u>);
- Update of MPMT checks according to the 2004 International Rules and Chapter 4 of the latest version of the JRC Technical report:
 - Groups of morphology codes considered as a single entity updated according to Table 15, following the adoption of the new ICD-O-3.2 codes;
 - Groups of topography codes considered as a single site updated according to Table 14;
 - **NEW**: more precise filtering criteria for MPMTs for records with behavior less than 3, implemented in the new configuration file located in the QCS 2.1.x *config* folder: "...\jrc-qcs-2.1.x\config\rules\encr2014\PrimaryMultipleFilter.csv";
- **NEW**: more precise identification and classification of MPMTs (old rule W-MPMT splitted into the new rules **W-MPMT** and **W-MPCR**).
- Update of morphology groups for checking multiple primaries according to the latest version of the JRC report (from QCS 2.1.1)
- Update of consistency checks between TNM values and basis of diagnosis (new rule W-BDpM from QCS 2.1.2)
- Update of topography codes according to ICD-O-3 (wrong code C74.4 removed from QCS 2.1.2)

Annex 2 – Known issues and bugs, and future improvements

Issues arising from cancer registries data validation through the QCS are continuously collected, to be fixed in next versions. The following lists includes already known issues:

- The QCS could not run properly on computers with very low RAM memory (less than 4GB). If the application does not start, or if it blocks during execution, make sure that your local machine supports at least 4GB of memory;
- To avoid memory issues, the QCS should be run clicking on a BAT file containing a "2GB" label in the file's name. This solution could be blocked by strict system policies, such as those applied by universities, corporate environments, governments etc. If you experience memory issues and can't run the BAT file with the "2GB" label, please ask support to your local system administrator and ask to be granted the necessary permissions for running the "2GB" file;
- The MPMT warning W-MPCR could be wrongly assigned to some rare combinations of solid/haematological MPMTs: in case, please notify;
- If the QCS is run while the PDF output report is still opened by the user, the PDF will **not** be updated. This happens on purpose, as a "security block" that avoids to delete the results of the previous run while the user is still using the software;
- If the input dataset has a critically wrong header (the first line), with errors or missing variables, the QCS might not correctly produce output reports;
- The QCS still checks for the previous version's warning W-SEMO, which is not relevant anymore for the ECIS data call protocol. Should your dataset trigger any W-SEMO message, please ignore related warnings.

Annex 3 – List of error and warning codes

This annex reports the list of error and warning codes included in the output files "*QCS- Incidence-Output.pdf*" and "*QCS-Incidence-Output.txt*". The table or page numbers referenced in the list are from the latest version of the JRC Technical report.

Error codes

E-AGEC: Age is invalid or missing, and cannot be calculated age by subtracting date of incidence from date of birth, since one or both dates are invalid or missing.

E-AGED: The age calculated as difference in years from (Date of incidence – Date of birth) differs from the value of variable *Age* by more than one year.

E-AUVS (NEW): the combination of Autopsy 1 with vital status and/or survival (Date of incidence – Date of follow-up in days) is not valid. Vital status should be 2 and Survival should be 0 (Table 12).

E-BDVS (NEW): the combination of Basis of diagnosis 0 (DCO) with vital status and/or survival and/or stage is not valid. Vital status should be 2, survival should be 0 and stage should be 9 or blank (Table 11).

E-CoDA: The date of birth is not consistent with the date of incidence, i.e. the latter occurs before the date of birth.

E-CoDV: The date of last known vital status is not valid (e.g. when the date of the incidence and the date of the last known vital status are not consistent).

E-DUPL: The same patient ID/tumour ID combination is repeated in two or more records.

E-ECOD: ICD¹⁷ edition and cause of death combination are not valid, e.g. cause of death 157 (pancreatic cancer) with ICD edition 10 (the correct value for pancreatic cancer is C25 for ICD-10, and 157 in ICD-7, ICD-8 and ICD-9). The check is performed for ICD editions from 7 to 10.

E-FORM: Format error, e.g. when a character value is used when a numeric one is required.

E-HEAD: Errors in the file header, e.g invalid number/order of columns, invalid header's separator etc.

E-MISS: Value missing, e.g. when variable *morphology* is unknown. This errors applies to variables whose invalid/missing/unknown values have an impact on incidence statistics.

E-OUTR: Value out of range; value is not in agreement with the ones allowed by the ECIS data call protocol (for instance, behavior equal to 6 or sex set to 4).

E-RECO: The record has the wrong number of fields.

E-SETO: Sex and topography combinations are not valid (Table 3).

E-VSBD (NEW): the combination of vital status 1 with autopsy and/or basis of diagnosis is not valid. Autopsy should be not 1 and Basis of diagnosis should be not 0 (Table 10).

¹⁷ International Classification of Diseases (https://www.who.int/standards/classifications/other-

classifications/international-classification-of-diseases-for-oncology)

Warning codes

W-AGMT: Unlikely age and morphology/topography combinations (Table 2).

W-BDMO: Morphology too specific according to the basis of diagnosis. See pages 17-18 for valid combinations of basis of diagnosis and morphology.

W-BDMS: Morphology not specific enough according to the basis of diagnosis. See page 19 for valid combinations of basis of diagnosis and morphology.

W-BDMU: Basis of diagnosis and morphology/behavior combination is unlikely. See page 30 for valid combinations of basis of diagnosis and morphology.

W-BDpM: Basis of diagnosis and pM combination is not valid. If pM is 1 then basis of diagnosis should be 7 or 6 (see page 33).

W-BDpN: Basis of diagnosis and pN combination is not valid. If pN is not NX and is not missing then basis of diagnosis should be or 7 (see page 33).

W-BDpT: Basis of diagnosis and pT combination is not valid. If pT is not TX and is not missing then basis of diagnosis should be 7 (see page 33).

W-BEGR: Behavior and grade combination is not valid (Table 7).

W-BTNM: Invalid behavior and TNM combination, e.g. *Behavior*=3 and *pT*=*Tis* (see page 33).

W-EDIM: TNM edition and pM are not consistent. The warning is returned when TNM edition is 7 or 8, and pM or cM are "X", since this value should be "0".

W-MISS: Value missing, e.g. when variable *Autopsy* is blank. This check applies to variables whose invalid/missing/unknown values is not impacting on incidence statistics.

W-MOBE: Morphology and behavior combinations not included in the ICD-O-3 (some combinations are allowed, according to Table 9).

W-MOGR: Morphology and grade combination is unlikely (Tables 6 and 7).

W-MOTO: Morphology and topography combination is unlikely (see table 8)

W-MPCR: Multiple primary tumour (see § 5.3 of this document)

W-MPMT: Multiple primary tumour (see § 5.3 of this document)

W-SEMO: Sex and morphology combination is unlikely, e.g. female with seminoma. See table 5 for the list of unlikely combinations.

W-TBGR (**NEW**): combination of morphology/behavior and grade not valid for tumours with behavior <3 (Table 5).

W-TNME: TNM and stage are present, but TNM edition is not valid or missing. The warning is returned since it is not possible to make a consistency check between TNM and stage.

W-TNMM: TNM and stage are present, but the morphology is not included in the TNM, e.g. when only carcinomas can be staged in a given topography, but stage is filled in for sarcomas.

W-TNMS: TNM and stage are not consistent, e.g. pT is 1, pN is 0, pM is 0 and stage is IV. In case both pathological (pT, pN and pM) and clinical (cT, cN and cM) TNM are provided for a tumour, the QCS will check the consistency between the pathological TNM and stage.

W-UNKN: missing/unknown value for variables with no impact on incidence calculations, which however could be important for quality evaluations or survival analysisa (e.g. basis of diagnosis=9, or year of follow up=9999).

Annex 4 – Running the JRC-ENCR QCS in background

Overview

The QCS can be run in different modes as listed below:

- **GUI** (standard execution): opens the main window and wait for user's actions;
- Silent (background process): run in background and validate the file passed as argument;
- **Introspection** (consolle): prints information about protocols and rules.

To view all supported arguments, please run the application with the **-h** option. The actual syntax may depend by your Java installation, and should be something like:

java -jar jrc-qcs-2.1.jar -h

The main supported runtime arguments are the following (see *Command line examples* below):

- -gui: opens the application in GUI mode (standard execution);
- -v: validates target dataset using the target validation protocol;
- **-p**: prints the list of rules defining target protocol;
- -s: prints the list of variables addressed by target protocol;
- -fi: prints all the information about the target variable (aka *field*);
- **-rf**: prints the list of rules addressing the target variable (aka *field*).

Command line examples

Validation of a dataset using protocol the 2022 ECIS data call protocol (i.e. protocol with ID = 11):

java -jar jrc-qcs-2.1.jar -v 11 C:/Home/incidence_1.csv

List of rules for the 2022 ECIS data call protocol (i.e. protocol with ID = 11):

java -jar jrc-qcs-2.1.jar **-p** 11

Description of the topography field in the 2022 ECIS data call protocol (field with ID = 80):

java -jar jrc-qcs-2.1.jar -fi 80

List of rules addressing the topography field in the 2022 ECIS data call protocol (field ID = 80):

java -jar jrc-qcs-2.1.jar **-rf** 80

Sample scripts

The *Examples* directory of the application contains two sample files showing examples of usage as a **background** process:

- **Run-qcs.bat**: example of executing the application in Windows OS;
- **run-qcs.sh**: example of executing the application in Linux OS.

Remark: the sample files listed above DO NOT provide complete management of possible execution errors, and DO NOT access (nor read, nor parse) the output reports produced at the end of the validation process. The actual management of the execution outcome MUST BE handled by the caller, with respect of his/her specific client's *execution context* (e.g. type of operative system, execution from webapp, execution as system service, etc.) and of the specific client's *needs and business* (e.g. validation of a single line, validation of big files, synchronous validation, asynchronous validation, etc.).

These sample files are provided only to show an example of executing the application as a background process and how to intercept the possible process outcomes.

Output reports

At the end of the validation process, the application should produce all output reports in path:

<application base path>/output

Guidelines

Some of the reports produced in the *output* directory are intended to be accessed directly by the final user, therefore are formatted in a human-friendly style (PDF or TXT). If the client application needs to read, parse, analyse or process the results of the validation process, usage of the following report is recommended:

• **QCS-Incidence-Output.csv**: read this file in order to acknowledge the detailed result of the validation process, line by line. This should be the core report when the application is run as a background process.