

JRC TECHNICAL REPORTS



A proposal on cancer data quality checks:

one common procedure for European cancer registries



Carmen Martos, Emanuele Crocetti (Coordinator), Otto Visser, Brian Rous, Francesco Giusti and the *Cancer Data Quality Checks Working Group*

2018

Version 1.1 • July 2018

Joint Research Centre This publication is a Technical report by the Joint Research Centre (JRC), the European Commission's science and knowledge service. It aims to provide evidence-based scientific support to the European policymaking process. The scientific output expressed does not imply a policy position of the European Commission. Neither the European Commission nor any person acting on behalf of the Commission is responsible for the use that might be made of this publication.

Contact information

Carmen Martos Address: Joint Research Centre, Directorate F – Health, Consumers and Reference Materials, Health in Society Unit, Via Enrico Fermi 2749, TP 127, 21027 Ispra (VA), Italy E-mail: carmen.martos@ec.europa.eu Tel.:+390332789074

JRC Science Hub https://ec.europa.eu/jrc

JRC105078

EUR 29089 EN

PDF	ISBN 978-92-79-77889-6	ISSN 1831-9424	doi:10.2760/429053
Print	ISBN 978-92-79-77890-2	ISSN 1018-5593	doi:10.2760/50693

Luxembourg: Publications Office of the European Union, 2018

© European Union, 2018

Reuse is authorised provided the source is acknowledged. The reuse policy of European Commission documents is regulated by Decision 2011/833/EU (OJ L 330, 14.12.2011, p. 39).

For any use or reproduction of photos or other material that is not under the EU copyright, permission must be sought directly from the copyright holders.

How to cite this report: Carmen Martos, Emanuele Crocetti (Coordinator), Otto Visser, Brian Rous, Francesco Giusti and the Cancer Data Quality Checks Working Group, A proposal on cancer data quality checks: one common procedure for European cancer registries – version 1.1, EUR 29089 EN, Publications Office of the European Union, Luxembourg, 2018, ISBN 978-92-79-77889-6, doi:10.2760/429053, JRC105078.

All images © European Union 2018

Printed in Italy



A proposal on cancer data quality checks: **one common procedure** for **European cancer registries**

Carmen Martos, Emanuele Crocetti (Coordinator), Otto Visser, Brian Rous, Francesco Giusti and the *Cancer Data Quality Checks Working Group*

2018

Version 1.1 • July 2018





Table of Contents

Working group on cancer data quality checks							
Contributors		5					
1 . Introductio	on	7					
2. Case defini	ition and variable format quality checks	9					
2.1 . Case of	definition	9					
2.2. Variat	2.2. Variables and their format quality checks						
3 . List of qua	lity checks: internal consistency	15					
3.1 . Consi	3.1. Consistency within variables						
3.2 . Consi	stency between variables	26					
3.2.1.	Coherence of dates	26					
3.2.2.	Consistency between tumour data and demographic information	26					
	Consistency between age/topography/morphology	26					
	Consistency between sex/topography	28					
	Consistency between sex/morphology	29					
3.2.3.	Consistency between tumour variables	30					
	Consistency between basis of diagnosis/morphology/behaviour	30					
	Consistency between morphology/grade	31					
	Consistency between topography/laterality	33					
	Consistency between topography/morphology	33					
3.3. Specif	fic additional checks for survival analysis	40					
3.4. Other	additional checks on the extent of the disease	40					
4 . Quality Ch	ecklist for Multiple Primary Malignant Tumours	42					
5. References		47					
Appendix I: The Anatomical Therapeutic Chemical code, generic name of the drug and trade name 4							
Appendix II: TNM 6 edition stage grouping and corresponding T, N, M values							
Appendix III: TNM 7 edition stage grouping and corresponding T, N, M values							

Working group on cancer data quality checks

Claudia ALLEMANI CONCORD programme, London School of Hygiene Tropical Medicine, UK Manola BETTIO Joint Research Centre, European Commission Andrea BORDONI ENCR-IACR Steering Committee Riccardo CAPOCACCIA EUROCARE project Helena CARREIRA CONCORD programme, London School of Hygiene Tropical Medicine, UK Emanuele CROCETTI (coordinator) ENCR-GRELL Steering Committee Roberta DE ANGELIS EUROCARE project Tadeusz Dyba Joint Research Centre, European Commission Nadia DIMITROVA ENCR Steering Committee Gerda ENGHOLM Danish Cancer Society Gemma GATTA RARECARE project Anna GAVIN ENCR Steering Committee Francesco GIUSTI Joint Research Centre, European Commission Alexander KATALINIC ENCRSteeringCommittee Carmen MARTOS Joint Research Centre, European Commission Pamela MINICOZZI EUROCARE project Giorgia RANDI Joint Research Centre, European Commission Ivan RASHID Italian Association of Cancer Registries Stefano Rosso ENCR Steering Committee Brian Rous National Cancer Registration Service, England, UK Milena SANT EUROCARE project Eva STELIAROVA-FOUCHER International Agency for Research on Cancer Hans STORM ENCR-ANCR Steering Committee Otto VISSER Integraal Kankercentrum Nederland Lydia Vott Joint Research Centre, European Commission Helmut WALERIUS Directorate-General for Health and Consumer Protection, European Commission

Contributors

Daniela ALESSI Childhood Cancer Registry of Piedmont Eva ARDANAZ Navarra Cancer Registry Mariangela AUTELITANO Cancer Registry of Milan Shiva AYOUBI Swedish Cancer Registry Simone BOECKMANN Cancer Registry of Lower Saxony Bertrand CAMEY Registre Fribourgeois des Tumeurs Léonie Casagranda Association of the Childhood Cancer Registry of the Rhône-Alpes Region (ARCERRA) Helen CURRY West Midlands Regional Children's Tumour Registry Silvia Dehler Cancer Registry Zurich and Zug Nick DESSYPRIS Nationwide Registry for Childhood Haematological Malignancies-Brain Tumour (NaReCHeM-BT) Chakameh SAFAEI DIBA National Cancer Registry of Slovakia Silvia Ess St. Gallen-Appenzell & Grisons-Glarus Anna Clara FANETTI Registro Tumori della provincia di Sondrio-Sondrio Cancer Registry Gonçalo FORJAZ Azores Cancer Registry Zivana GAVRIC Cancer Registry Republika Srpska Adriano Giacomin Piedmont Cancer Registry, Province of Biella Stanisław Góźdź and Paweł MACEK Holycross Cancer Center Pascale GROSCLAUDE Registries of FRANCIM Network Gaël HAMMER Registre Morphologique des Tumeurs au Grand-Duché de Luxembourg Christine HEAD National Cancer Registration Service for England Miriam HOLZMANN Cancer Registry of Schleswig-Holstein, Germany Kris HENAU and Liesbet VAN EYCKEN Belgian Cancer Registry Zsuzsanna JAKAB Hungarian Childhood Cancer Registry Rosario JIMÉNEZ, Ana I. MARCOS and JOSÉ MARÍA DÍAZ Cancer Registry of Cuenca, Counselling of Health and Social Affairs, Castilla-La Mancha, Spain

Margit Mägi, TIIU AARELEID, Kaire INNOS and Mati RAHU Estonian Cancer Registry

Working group on cancer data quality checks | Contributors | 5

Rafael MARCOS-GRAGERA Girona Cancer Registry Luigino DAL MASO Friuli Venezia Giulia Cancer Registry Martin MEYER Population Based Cancer Registry Bavaria, Germany Ana Miranda Lisbon Cancer Institute-South Regional Cancer Registry Silvia PATRIARCA Piedmont Cancer Registry Josefina PERUCHA GONZÁLEZ La Rioja Cancer Registry Meike RESSING Cancer Registry Rhineland-Palatinate Anne SCHMIDT Cancer Registry Thurgau Giedre SMAILYTE Lithuanian Cancer Registry Giovanna TAGLIABUE Lombardy Cancer Registry, Varese Province Ana TORRELLA RAMOS Castellón Cancer Registry (Comunidad Valenciana), Spain SIGRÚN STEFÁNSDÓTTIR Icelandic Cancer Registry Zdravka VALERIANOVA and Minka YORDANOVA Bulgarian National Cancer Registry Maja PRIMIC ŽAKELJ Cancer Registry of Republic of Slovenia Miroslav Zvolský Czech National Cancer Registry

L. Introduction

The aim of population-based cancer registries (CRs) is: *a*) to obtain information from all new cases in a well-defined geographic area to assess the magnitude of the cancer burden and its evolution, and *b*) to provide a basis for research on cancer causes and outcome (incidence, prevalence and survival). Therefore, CRs contribute to monitoring the impact and effectiveness of policy implementation through monitoring outcomes such as incidence, prevalence or survival. The reliability and utility of the information provided by CRs depends on the quality of the data collected.

Three aspects are usually regarded when evaluating the quality of the data in CRs: comparability, completeness and validity. An additional quality indicator- the timeliness of registry procedures-is also considered.

A variety of methods and tools have been used to check the data validity of CRs. Therefore, the European Network of Cancer Registries (ENCR) in cooperation with the Joint Research Centre (JRC) has been working to establish a comprehensive and standardised list of data quality checks to be adopted by European CRs and European projects that would address the current fragmented and sometimes conflicting situation regarding validation of data collected for different purposes. The adoption of a common list of variables, formats and standard data quality checks will improve the harmonisation of European cancer data and the adherence to standardised data quality procedures will give CRs the opportunity to participate easily in different international projects.

Three workshops on data quality checks took place in JRC-Ispra, on 2 July and 15 October 2013 and 4 June 2014 (http://www.encr.eu/). The outcome of the first two meetings was the creation of a comprehensive list of the existing data quality checks currently in place for various European projects, collected and summarised by the JRC. At the conclusion of the second workshop, agreements were reached for drafting a preliminary list of mandatory and basic variables and their formats.

Furthermore, and in view of the third workshop, the Working Group drafted a document 'A proposal on cancer data quality checks: one common procedure for European cancer registries' taking into account the case definition, the list of variables and their format agreed upon during the second workshop, as well as the existing edits and the expertise of cancer registry experts. This report was disseminated among the European CRs for consultation. The report was the object of discussion and final revision at the third workshop, and final agreements were reached concerning case definition, variables and their format and data quality control list.

This document is the result of a collaborative project between the ENCR, the JRC, the Working Group on Cancer Data Quality Checks and European cancer registries. The final outcome of the project was an ENCRendorsed recommendations document, to be issued and presented at the 2014 ENCR Scientific Meeting and General Assembly, 12-14 November 2014, at JRC-Ispra.

This report focuses on *case definitions* and *variable format* quality checks and *internal consistency* within and between collected variables. The proposed quality checklist allows the identification of: impossible codes or code combinations, unlikely codes or code combinations and possible but very rare code or code combinations. Finally, the list of drugs used for chemotherapy, hormonal therapy, targeted therapy, immunotherapy and other therapies used in cancer treatment was revised and included in the *Appendices*: the list of drugs contains the Anatomical Therapeutic Chemical (ATC) code as well as the generic and trade names.

With respect to the first version (1.0), in this updated version (1.1) of the report, the quality checklist of warnings for multiple primary malignant tumours (MPMT) according to the current International Rules for Primary Cancers published in 2004 have been added (Chapter 4). Moreover, *Table* **6** (Valid combinations for behaviour and topography/morphology) in the previous version of the report (1.0) has been discarded.

Case definition and variable format . quality checks

The cancer data quality check list included in this report is based on the following case definition for CRs and European projects.

2.1. Case definition

- All primary malignant tumours (behaviour = 3), including basal cell and squamous cell carcinomas of skin.
- Benign tumours of the central nervous system (CNS).
- Uncertain behaviour tumours of CNS and urinary bladder.
- In situ tumours: breast, cervix, colon, rectum, urinary bladder and melanoma of the skin.

An extent of this case definition could be considered according to the European CRs needs in the future.

2.2. Variables and their format quality checks

During the second workshop on quality checks agreements were reached for the mandatory and basic variable list and their format. A revision of the list of variables and formats was made during the third workshop. *Table 1* shows the list of the variables: description, format, mandatory or nonmandatory status, missing/unknown values and the allowed values on which quality checks are based.

Variable description	Format	Mandatory	Missing/un- known values	Allowed values		
(Checkflag) The ENCR-JRC QC list	F1	Yes	Not allowed	Allowed values: 0, 1 $0 \rightarrow \text{Not checked}$ $1 \rightarrow \text{Checked}$		
Patient identification number	A20	Yes (accord- ing to registry coding)	Not allowed	Not allowed to have duplicate combination of the two variables: Patient identification number		
Tumour sequence number	F2	Yes (accord- ing to registry coding)	Not allowed	+Tumour sequence number in the same dataset		
Day of birth	A2 DD	Y	99	Range of allowed values: from 01 to 31 and 99		
Month of birth	A2 MM	Y	99	Range of allowed values: from 01 to 12 and 99 Warning for value = 99		
Year of birth	F4 YYYY	Y	9999	Range of allowed values: >1842 and ≤ the current year Warning for value = 9999		
F: Numeric variable A: Alphanumeric variable Y=ves N=no						

Table 1. Quality checks for the variables and their formats.

2. Case definition and variable format quality checks | 9

Variable description	Format	Mandatory	Missing/un- known values	Allowed values
Sex	F1	Y	9	Allowed values: 1, 2, 3, 9 $1 \rightarrow Male$ $2 \rightarrow Female$ $3 \rightarrow Other$ $9 \rightarrow Unknown$ Warning for value = 9
Day: date of incidence	A2 DD	Y	99	Range of allowed values: from 01 to 31 and 99
Month: date of incidence	A2 MM	Y	99	Range of allowed values: from 01 to 12 and 99 Warning for value = 99
Year: date of incidence ENCR recommendation for incidence date http://www.encr.eu/sites/default /files/pdf/incideng.pdf	F4 YYYY	Y	Not allowed	Range of allowed values: >1941 and ≤ the current year
Day of case registration	A2 DD	Ν	99	Range of allowed values: from 01 to 31 and 99
Month of case registration	A2 MM	Ν	99	Range of allowed values: from 01 to 12 and 99
Year of case registration	F4 YYYY	Ν	9999	Range of allowed values: >1941 and ≤ the current year
Age at diagnosis in years	F3	Y*	999	Range of allowed values: ≥0 and <121 Warning for value = 999 if com- pleted dates are not available
Basis of diagnosis (BoD) ENCR recommendations http://www.encr.eu/sites/default /files/pdf/basisd.pdf	F1	Y	9	Allowed values: 0, 1, 2, 4, 5, 6, 7, 9 $0 \rightarrow$ Death certificate only (DCO) $1 \rightarrow$ Clinical $2 \rightarrow$ Clinical investigation $4 \rightarrow$ Specific tumour markers $5 \rightarrow$ Cytology $6 \rightarrow$ Histology of a metastasis $7 \rightarrow$ Histology of a primary tumour $9 \rightarrow$ Unknown Warning for value = 9
ICD-O-3 topography (topography of the metastasis is not admitted)	A4	Y	Not allowed	Valid code in ICD-O-3. Warning for undefined topography when BoD is 5 or 7 C809; C76 (C760, C761, C762, C763, C764, C765, C767 and C768); C14 (C140, C148); C26 (C260, C268, C269); C39 (C390, C398, C399); C559; C579; C639; C689; C729; C759

F: Numeric variable A: Alphanumeric variable Y = yes N = no

* If complete date of birth and/or date of incidence are missing or unknown.

Variable description	Format	Mandatory	Missing/un- known values	Allowed valu	ies	
ICD-0-3 morphology	F4	Y	Not allowed	Valid code in ICD-0-3 and updated in 2011	Valid code in ICD-O-3 and updated in 2011 Warning for unde- fined morphology	
ICD-0-3 behaviour	F1	Y	Not allowed	Accepted value: 0-3	taking into account BoD (See <i>Figure 2</i> , p. 30)	
Incidental finding of cancer at the autopsy	F1	Y	9	Allowed value $0 \rightarrow No$ $1 \rightarrow Yes$ $9 \rightarrow Unknown Warning for$	es: 0, 1, 9 own value = 9	
ICD-O-3 grade	F1	Υ	9	Allowed values: 1-9 $1 \rightarrow$ Well differentiated, $2 \rightarrow$ Moderately differentiated $3 \rightarrow$ Poorly differentiated $4 \rightarrow$ Undifferentiated, anaplastic $5 \rightarrow$ T-cell; T-precursor $6 \rightarrow$ B-Cell; Pre-B; B-precursor $7 \rightarrow$ Null cell; Non T-non B $8 \rightarrow$ NK cell (natural killer cell) $9 \rightarrow$ Unknown		
Laterality of paired organs	F1	Ν	9	Allowed values: 0-4, 9 $0 \rightarrow \text{Not applicable}$ $1 \rightarrow \text{Right}$ $2 \rightarrow \text{Left}$ $3 \rightarrow \text{Unilateral NOS}$ $4 \rightarrow \text{Bilateral}$ $9 \rightarrow \text{Unknown}$		
Vital status at last contact	F1	Y	9	Allowed values: 1, 2, 9 $1 \rightarrow Alive$ $2 \rightarrow Dead$ $9 \rightarrow Unknown$ Warning for value = 9		
Day of the last known vital status	A2 DD	Y	99	Range of allo from 01 to	wed values: 31 and 99	
Month of the last known vital status	A2 MM	Y	99	Range of allo from 01 to Warning for	wed values: 12 and 99 value = 99	
Year of the last known vital status	F4 YYYY	Y	9999	Range of allo >1941 and Warning for	wed values: d ≤ the current year value = 9999	
Age at the last known vital status in years	F3	Y**	999	Range of allo ≥0 and < 1 Warning for	wed values: 21 value = 999	

F: Numeric variable A: Alphanumeric variable Y=yes N=no

** If complete date of birth, data of incidence and/or date of end of follow-up are missing or unknown.

2. Case definition and variable format quality checks | 11

Variable description	Format	Mandatory	Missing/un- known values	Allowed values
Duration of survival in days	F5	Y***	99999	≥0 Warning for value = 99999
Official underlying cause of death (ICD)	A5	Ν	99999	Valid code in ICD according to ICD edition
ICD edition used for coding cause of death	F2	Ν	99	Range of allowed values: ≥7 and ≤ 10 It has to be periodically updated
TNM stage, pathological primary site (pT)	A6	Ν	999999	Prefix modifiers will be considered: y: stage assessed after neo- adjuvant therapy; a: stage determined at autopsy (See <i>Table 2</i>)
TNM stage, pathological lymph nodes (pN)	A4	Ν	9999	(See Table 2)
TNM stage, pathological metastases (pM)	A4	Ν	9999	(See Table 2)
TNM stage, clinical primary site (cT)	A5	Ν	99999	(See Table 2)
TNM stage, clinical lymph nodes (cN)	A3	Ν	999	(See Table 2)
TNM stage, clinical metastases (cM)	A3	Ν	999	(See Table 2)
TNM stage grouping	A4	Ν	9999	Based on pathological TNM if it is available or clinical TNM when pathological TNM is not available (See <i>Table 2</i>)
TNM edition	F2	Ν	99	Allowed values: 6, 7, 99 It has to be periodically updated
Condensed TNM, T ENCR recommendations http://www.encr.eu/sites/default/ files/pdf/extentofdisease.pdf	A2	Ν	99	Allowed values: TL, TA, TX, 99 TL \rightarrow Localised TA \rightarrow Advanced TX \rightarrow Unknown
Condensed TNM, N ENCR recommendations http://www.encr.eu/sites/default/ files/pdf/extentofdisease.pdf	A2	Ν	99	Allowed values: N0, N1, NX, 99 N0 \rightarrow No regional nodes N1 \rightarrow Regional nodes NX \rightarrow Unknown
Condensed TNM, M ENCR recommendations http://www.encr.eu/sites/default/ files/pdf/extentofdisease.pdf	A2	Ν	99	Allowed values: M0, M1, MX, 99 M0 \rightarrow No distant metastasis M1 \rightarrow Distant metastasis MX \rightarrow Unknown

F: Numeric variable A: Alphanumeric variable Y=yes N=no

*** If complete date of incidence and/or date of end of follow-up are missing or unknown.

Variable description	Format	Mandatory	Missing/un- known values	Allowed values
Dukes' stage	A1	Ν	9	Allowed values: A, B, C, D, 9 $A \rightarrow$ Dukes' stage A $B \rightarrow$ Dukes' stage B $C \rightarrow$ Dukes' stage C $D \rightarrow$ Dukes' stage D $9 \rightarrow$ Dukes' stage unknown
FIGO stage	Α3	Ν	999	Allowed values: 0, I, II, III, IVA, IVB, 999 $0 \rightarrow$ FIGO stage 0 $I \rightarrow$ FIGO stage 1 $II \rightarrow$ FIGO stage II $II \rightarrow$ FIGO stage III $IVA \rightarrow$ FIGO stage IVA $IVB \rightarrow$ FIGO stage IVB $9 \rightarrow$ FIGO stage unknown
Summary extent of disease (EOD)	F1	Ν	9	Allowed values: 1, 2, 3, 4, 5, 9 $1 \rightarrow Confined$ $2 \rightarrow Adjacent tissues, and/or regional lymph-nodes 3 \rightarrow Distant organs4 \rightarrow Not confined but not specified whether code 2 or 3 applies 5 \rightarrow Not distant metastasis but not specified whether code 1 or 2 applies 9 \rightarrow Unknown$
Tumour size in mm with decimal	F5	Ν	999.9	>0 or 999.9
Number examined nodes	F2	Ν	99	From 0 to 99
Number metastatic nodes	F2	Ν	99	Number metastasis nodes ≤ Number examined nodes
Sentinel nodes	F1	Ν	9	Allowed values: 1, 2, 9 $1 \rightarrow \text{Done}$ $2 \rightarrow \text{Not done}$ $9 \rightarrow \text{Unknown}$
Metastasis in sentinel nodes	F1	Ν	9	Allowed values: 1, 2, 9 $1 \rightarrow Yes$ $2 \rightarrow No$ $9 \rightarrow Unknown$

F: Numeric variable

A: Alphanumeric variable Y=yes N=no

2. Case definition and variable format quality checks | 13

Variable description	Format	Mandatory	Missing/un- known values	Allowed values
C factor ENCR recommendations http://www.encr.eu/sites/default/ files/pdf/extentofdisease.pdf	F1	Ν	9	Allowed values: 1, 2, 3, 4, 5, 9 $1 \rightarrow C1$ Evidence from standard diagnostic methods only $2 \rightarrow C2$ Evidence obtained by special diagnostic means $3 \rightarrow C3$ Evidence from surgical exploration, including biopsy and cytology $4 \rightarrow C4$ Evidence following de- finitive surgery and pathological examination of the resected specimen $5 \rightarrow C5$ Evidence from autopsy $9 \rightarrow$ unknown
Surgery	F1	Ν	9	Allowed values: 1, 2, 9 $1 \rightarrow Yes$ $2 \rightarrow No$ $9 \rightarrow Unknown$
Chemotherapy	F1	Ν	9	Allowed values: 1, 2, 9 $1 \rightarrow Yes$ $2 \rightarrow No$ $9 \rightarrow Unknown$
Systemic therapy, other than chemotherapy	F1	Ν	9	Allowed values: 1, 2, 9 $1 \rightarrow Yes$ $2 \rightarrow No$ $9 \rightarrow Unknown$
Radiotherapy	F1	Ν	9	Allowed values: 1, 2, 9 $1 \rightarrow Yes$ $2 \rightarrow No$ $9 \rightarrow Unknown$
Hormone therapy	F1	Ν	9	Allowed values: 1, 2, 9 $1 \rightarrow Yes$ $2 \rightarrow No$ $9 \rightarrow Unknown$
Bone marrow transplantation	F1	Ν	9	Allowed values: 1, 2, 9 $1 \rightarrow Yes$ $2 \rightarrow No$ $9 \rightarrow Unknown$

F: Numeric variable A: Alphanumeric variable Y=yes N=no

S. List of quality checks: internal consistency

3.1. Consistency within variables

Most of the quality control checks for single variables concern its format and allowed values, detailed in *Table 1*. Nevertheless, other specific quality checks detailed below are required for dates and TNM/stage.

Regarding '*dates*', some simple rules are necessary when they are collected as three independent variables reporting 'day', 'month' and 'year' (*Figure 1*):

if the variable 'month' is equal to January (01), March (03), May (05), July (07), August (08), October (10) or December (12) the range of values for the variable day is from 1 to 31;

- if the variable 'month' is April (04), June (06), September (09) or November (11) the range of values for the variable 'day' is from 1 to 30;
- if the variable 'month' is February (02) the range of values for the variable 'day' is from 1 to 28, except for leap-years in which the range of values for the 'day' is from 1 to 29. The algorithm to define a leap-year is the following:
 - it is a year divisible by 4 (*i.e.* 2004, 2008, etc.). This rule does not apply to centennial years (those exactly divisible by 100 (*i.e.* 1900, 2100, etc.);
 - it is a centennial year (exactly divisible by 100) and it is also exactly divisible by 400 (like 2000, 2400).

Figure 1. Range of values for the variable 'day' according to variables 'month' and 'year'.



3. List of quality checks: internal consistency | 15

Age at diagnosis: measured as the age in years at the patient's last birthday. Age could be calculated if both incidence and birth dates are registered (or at least the incidence year and birth year). It is recommended using algorithms to impute the dates before calculating the age, when possible. The range of values must be between 0 and 120.

This variable is optional if at least both incidence and birth year are available, while it is mandatory when at least one of them is missing:

• If only year of diagnosis and birth are available, then age at diagnosis is computed as a difference:

Age at diagnosis = year of incidence - year of birth

• If the month and year of both dates are known, then age at diagnosis is computed as:

Age at diagnosis = [(year of incidence * 12 + month of incidence) - (year of birth * 12 + month of birth)] / 12 Integer

• If the month of diagnosis and birth are known and equal, and the day of diagnosis is earlier than the day of birth, then 1 is subtracted from the calculated age.

Once computed, the age at diagnosis should be compared with what was provided by the CRs, and be consistent according to the following rule:



TNM and stage grouping values depend on the cancer topography and the edition of the TNM classification. The TNM system includes both clinical (pre-treatment) and pathological (post-surgical histopathological) classifications. The clinical classification is designated as cTNM, and the pathological as pTNM.

Table 2 includes the valid values for T (extent of primary tumour), N (absence/presence and extent of regional lymph node metastasis) and M (absence/presence of distant metastasis) as well as the corresponding stage by topography and revision of TNM classification (6 and 7), according to the case definitions described in section 2.1 of this report.

T, N and M values are similar for clinical and pathological classifications with very few exceptions. Therefore, unless clearly specified, the T, N and M values included in *Table 2* are valid for both cTNM and pTNM classifications.

In addition, *Appendices II* and *III* contain detailed stage grouping as well as the corresponding T, N, M values for the TNM 6 and TNM 7 editions, respectively.

^{16 |} A proposal on cancer data quality checks: one common procedure for European cancer registries

 Table 2. Valid values for T, N, M and stage by cancer topography and TNM edition.§

Topography	TNM edition	т	Ν	М	Stage grouping
Lip and oral cavity C00, C02-C06	6	TX, T1, T2, T3, T4a, T4b	NX, N0, N1, N2, N2a, N2b, N2c, N3	MX, M0, M1	I, II, III, IVA, IVB, IVC
(except C051 and C052)	7	TX, T1, T2, T3, T4a, T4b	NX, NO, N1, N2, N2a, N2b, N2c, N3	M0, M1	I, II, III, IVA, IVB, IVC
Oropharynx C01, C051, C052, C090, C091, C099, C100, C102, C103		TX, T1, T2, T3, T4a, T4b	NX, NO, N1, N2, N2a, N2b, N2c, N3		I, II, III, IVA, IVB, IVC
Nasopharynx C11	6	TX, T1, T2, T2a, T2b, T3, T4	NX, NO, N1, N2, N3, N3a, N3b	MX, MO, M1	I, IIA, IIB, III, IVA, IVB, IVC
Hypopharynx C12, C13		TX, T1, T2, T3, T4a, T4b	NX, NO, N1, N2, N2a, N2b, N2c, N3		I, II, III, IVA, IVB, IVC
Oropharynx C01, C051, C052, C090, C091, C099, C100, C102, C103		TX, T1, T2, T3, T4a, T4b	NX, NO, N1, N2, N2a, N2b, N2c, N3		
Nasopharynx C11	7	TX, T1, T2, T3, T4	NX, NO, N1, N2, N3, N3a, N3b	M0, M1	I, II, III, IVA, IVB, IVC
Hypopharynx C12, C13		TX, T1, T2, T3, T4a, T4b	NX, NO, N1, N2, N2a, N2b, N2c, N3		
Major salivary glands	6	TX, T1, T2, T3, T4a, T4b	NX, NO, N1, N2, N2a, N2b, N2c, N3	MX, M0, M1	I, II, III, IVA, IVB, IVC
C07, C08	7	TX, T1, T2, T3, T4a, T4b	NX, NO, N1, N2, N2a, N2b, N2c, N3	M0, M1	I, II, III, IVA, IVB, IVC
Oesophagus C15	6	TX, T1, T2, T3, T4	NX, NO, N1	MX, M0, M1, M1a (for C153 and C155), M1b (for C153, C154 and C155)	I, IIA, IIB, III, IV, IVA, IVB
Oesophagus C15, C160	7	TX, T1, T2, T3, T4, T4a, T4b	NX, NO, N1, N2, N3	M0, M1	IA, IB, IIA, IIB, IIIA, IIIB, IIIC, IV
Stomach C16	6	TX, T1, T2, T2a, T2b, T3, T4	NX, NO, N1, N2, N3	MX, M0, M1	IA, IB, II, IIIA, IIIB, IV
Stomach C161-C164	7	TX, T1, T1a, T1b, T2, T3, T4, T4a, T4b	NX, NO, N1, N2, N3, N3a, N3b	M0, M1	IA, IB, IIA, IIB, IIIA, IIIB, IIIC, IV
Small intestine	6	TX, T1, T2, T3, T4	NX, NO, N1	MX, M0, M1	I, II, III, IV
	7	TX, T1, T1a, T1b, T2, T3, T4	NX, NO, N1, N2	M0, M1	I, IIA, IIB, IIIA, IIIB, IV
Appendix carcinoma C181	7	TX, Tis, T1, T2, T3, T4, T4a, T4b	NX, NO, N1, N2	M0, M1, M1a, M1b	0, I, IIA, IIB, IIC, IIIA, IIIB, IIIC, IVA, IVB, IVC

[§] In general, the classification applies to carcinomas, except for some topographies with specific morphologies that use separate classifications (*i.e.* appendix-carcinoid). There should be histological confirmation of the disease.

Topography	TNM edition	т	N	М	Stage grouping
Appendix carcinoid C181 (well differentiated neuroendocrine tumour)	7	TX, T1, T1a, T1b, T2, T3, T4	NX, NO, N1	M0, M1	I, II, III, IV
Colon and rectum C18, C19, C20	6	TX, Tis, T1, T2, T3, T4	NX, NO, N1, N2	MX, M0, M1	0, I, IIA, IIB, IIIA, IIIB, IIIC, IV
Colon and rectum C18 (excluded C181), C19, C20	7	TX, Tis, T1, T2, T3, T4, T4a, T4b	NX, NO, N1, N1a, N1b, N1c, N2, N2a, N2b	M0, M1, M1a, M1b	0, I, II, IIA, IIB, IIC, III, IIIA, IIIB, IIIC, IVA, IVB
Anal canal	6	TX, T1, T2, T3, T4	NX, NO, N1, N2, N3	MX, M0, M1	I, II, IIIA, IIIB, IV
(211	7	TX, T1, T2, T3, T4	NX, NO, N1, N2, N3	M0, M1	I, II, IIIA, IIIB, IV
Gastrointestinal stromal tumour (GIST) C15, C16, C170, C171, C172, C18, C20, C481 (Omen- tum, mesentery)	7	TX, T1, T2, T3, T4	NX, NO, N1	M0, M1	C16, C481 (omental GIST) IA, IB, II, IIIA, IIIB, IV C17, C15, C18, C20, C481 (mesentery) I, II, IIIA, IIIB, IV
Gastric, small and large Intestinal carcinoid tumours (appendix excluded)*	7	Stomach TX, T1, T2, T3, T4 Duodenum, ampul- la, jejunum, ileum TX, T1, T2, T3, T4 Large intestine TX, T1, T1a, T1b, T2, T3, T4	NX, NO, N1	M0, M1	I, IIA, IIB, IIIA, IIIB, IV
Liver and intra- hepatic bile ducts C220,C221	6	TX, T1, T2, T3, T4	NX, NO, N1	MX, M0, M1	I, II, IIIA, IIIB, IIIC, IV
Liver-hepatocel- lular carcinoma C220	_	TX, T1, T2, T3, T3a, T3b, T4		10.11	I, II, IIIA, IIIB, IIIC, IVA, IVB
Liver-intrahepatic bile ducts C221	/	TX, T1, T2a, T2b, T3, T4	NX, NU, NI	MU, MI	I, II, III, IVA, IVB
Gallbladder C23	6	TX, T1, T1a, T1b, T2, T3, T4	NX, NO, N1	MX, M0, M1	IA, IB, IIA, IIB, III, IV
	7	TX, T1, T1a, T1b, T2, T3, T4	NX, NO, N1	M0, M1	I, II, IIIA, IIIB, IVA, IVB

[§] In general, the classification applies to carcinomas, except for some topographies with specific morphologies that use separate classifications (*i.e.* appendix-carcinoid). There should be histological confirmation of the disease.

* Well-differentiated neuroendocrine tumours and Well-differentiated neuroendocrine carcinomas.

Ta	ble	2. ((cont.)	§
		(

Topography	TNM edition	т	Ν	М	Stage grouping
Extrahepatic bile	6	TX, T1, T2, T3, T4	NX, NO, N1	MX, M0, M1	IA, IB, IIA, IIB, III, IV
C240	7	C240-Perihilar TX, T1, T2a, T2b, T3, T4 C240-Distal TX, T1, T2, T3, T4	NX, NO, N1	M0, M1	C240-Perihilar I, II, IIIA, IIIB, IVA, IVB C240-Distal IA, IB, IIA, IIB, III, IV
Ampulla of vater	6	TX, T1, T2, T3, T4	NX, NO, N1	MX, M0, M1	IA, IB, IIA, IIB, III, IV
	7	TX, T1, T2, T3, T4	NX, NO, N1	M0, M1	IA, IB, IIA, IIB, III, IV
Pancreas	6	TX, T1, T2, T3, T4	NX, NO, N1	MX, M0, M1	IA, IB, IIA, IIB, III, IV
25	7	TX, T1, T2, T3, T4	NX, NO, N1	M0, M1	IA, IB, IIA, IIB, III, IV
Supraglottis C321,C101		TX, T1, T2, T3, T4a, T4b			
Glottis C320	6	TX, T1, T1a, T1b, T2, T3, T4a, T4b	NX, NO, N1, N2, N2a, N2b, N2c, N3	MX, MO, M1	I, II, III, IVA, IVB, IVC
Subglottis C322		TX, T1, T2, T3, T4a, T4b			
Supraglottis C321,C101	7	TX, T1, T2, T3, T4a, T4b			
Glottis C320		7	TX, T1, T1a, T1b, T2, T3, T4a, T4b	NX, NO, N1, N2, N2a, N2b, N2c, N3	M0, M1
Subglottis C322		TX, T1, T2, T3, T4a, T4b			
Nasal cavity and paranasal sinuses	6	TX, T1, T2, T3, T4a, T4b	NX, NO, N1, N2, N2a, N2b, N2c, N3	MX, M0, M1	I, II, III, IVA, IVB, IVC
C300, C310, C311	7	TX, T1, T2, T3, T4a, T4b	NX, NO, N1, N2, N2a, N2b, N2c, N3	M0, M1	I, II, III, IVA, IVB, IVC
Malignant melanoma of aerodigestive tract (C00-C06, C10- C14, C30-C32)	7	TX, T3, T4a, T4b	NX, NO, N1	M0, M1	III, IVA, IVB, IVC
Lung C34	6	TX, T1, T2, T3, T4	NX, NO, N1, N2, N3	MX, M0, M1	IA, IB, IIA, IIB, IIIA, IIIB, IV
	7	TX, T1, T1a, T1b, T2, T2a, T2b, T3, T4	NX, NO, N1, N2, N3	M0, M1, M1a, M1b	IA, IB, IIA, IIB, IIIA, IIIB, IV
Pleural mesothelioma	6	TX, T1, T1a, T1b, T2, T3, T4	NX, NO, N1, N2, N3	MX, M0, M1	IA, IB, II, III, IV
C384	7	TX, T1, T1a, T1b, T2, T3, T4	NX, NO, N1, N2, N3	M0, M1	IA, IB, II, III, IV

[§] In general, the classification applies to carcinomas, except for some topographies with specific morphologies that use separate classifications (*i.e.* appendix-carcinoid). There should be histological confirmation of the disease.

Topography	TNM edition	T	N	М	Stage grouping
Bone C40,C41	6	TX, T1, T2, T3	NX, NO, N1	MX, M0, M1, M1a, M1b	IA, IB, IIA, IIB, III, IVA, IVB
	7	TX, T1, T2, T3	NX, NO, N1	M0, M1, M1a, M1b	IA, IB, II, III, IVA, IVB
Softtissues C381, C382, C383, C47, C480, C49, 9581/3, 8804/3, 9220/3, 9180/3,	6**	TX, T1, T1a, T1b, T2, T2a, T2b	NX, NO, N1	MX, M0, M1	IA, IB, IIA, IIB, III, IV
8810/3, 8890/3, 8850/3, 8830/3, 9150/3, 8990/3, 9540/3, 8900/3, 9040/3, 8800/3	7***	TX, T1, T1a, T1b, T2, T2a, T2b	NX, NO, N1	M0, M1	I, IIA, IIB, III, IV
Carcinoma of eyelid C441	6	TX, T1, T2, T3, T4	NX, NO, N1	NX, N0, N1 MX, M0, M1	
	7	TX, T1, T2a, T2b, T3a, T3b, T4	NX, NO, N1	M0, M1	IA, IB, IC, II, IIIA, IIIB, IIIC, IV
Carcinoma of conjunctiva	6	TX, T1, T2, T3, T4, T4a, T4b, T4c, T4d	NX, NO, N1	MX, M0, M1	No stage grouping recommended
C690	7	TX, T1, T2, T3, T4, T4a, T4b, T4c, T4d	NX, NO, N1	M0, M1	No stage grouping recommended
Malignant melanoma of	6	TX, T1, T2, T3, T4	NX, NO, N1	MX, M0, M1	No stage grouping recommended
conjunctiva C690	7	TX, T1, T1a, T1b, T1c, T1d, T2, T2a, T2b, T2c, T2d, T3, T3a, T3b, T3c, T3d, T4	NX, NO, N1	M0, M1	No stage grouping recommended
Malignant melanoma of uvea C693, C694	6	TX, T1, T1a, T1b, T1c, T2, T2a, T2b, T2c, T3, T3a, T4	NX, NO, N1	MX, M0, M1	I, II, III, IV
	7	TX, T1, T1a, T1b, T1c, T1d, T2, T2a, T2b, T2c, T2d, T3, T3a, T3b, T3c, T3d, T4, T4a, T4b, T4c, T4d, T4e	NX, NO, N1	M0, M1	I, IIA, IIB, IIIA, IIIB, IIIC, IV

[§] In general, the classification applies to carcinomas, except for some topographies with specific morphologies that use separate classifications (*i.e.* appendix-carcinoid). There should be histological confirmation of the disease.

** The following histological types are not included: Kaposi sarcoma, dermatofibrosarcoma (protuberans), fibromatosis (desmoid tumour), and sarcoma arising from the dura mater, brain, hollow viscera or parenchymatous organs (with the exception of breast sarcomas) and angiosarcoma.

*** The following histological types are not included: Kaposi sarcoma, dermatofibrosarcoma (protuberans), fibromatosis (desmoid tumour), sarcoma arising from the dura mater, brain, hollow viscera or parenchymatous organs (with the exception of breast sarcomas), angiosarcoma and gastrointestinal stromal tumours.

Topography	TNM edition	т	Ν	М	Stage grouping	
Sarcoma of orbit C696	6	TX, T1, T2, T3, T4	NX, NO, N1	MX, M0, M1	No stage grouping recommended	
	7	TX, T1, T2, T3, T4	NX, NO, N1	M0, M1	No stage grouping recommended	
Carcinoma of lachrymal gland	6	TX, T1, T2, T3, T3a, T3b, T4	NX, NO, N1	MX, M0, M1	No stage grouping recommended	
C695	7	TX, T1, T2, T3, T4, T4a, T4b, T4c	NX, NO, N1	M0, M1	No stage grouping recommended	
Retinoblastoma C692	6	Clinical T TX, T1, T1a, T1b, T2, T2a, T2b, T2c, T3, T4 Pathological T pTX, pT0, pT1, pT2, pT2a, pT2b, pT2c, pT3, pT3a, pT3b, pT3c, pT4	NX, NO, N1	Clinical M MX, M0, M1 Pathological M pMX, pM0, pM1, pM1a, pM1b	No stage grouping recommended	
	7	Clinical T TX, T1, T1a, T1b, T1c, T2, T2a, T2b, T3, T3a, T3b, T4, T4a, T4b, T4c, T4d Pathological T pTX, pT0, pT1, pT2, pT2a, pT2b, pT2c, pT3, pT3a, pT3b, pT3c, pT4	Clinical N NX, NO, N1 Pathological N pNX, pNO, pN1, pN2	Clinical M M0, M1 Pathological M pM0, pM1, pM1a, pM1b, pM1c, pM1d, pM1e	No stage grouping recommended	
Carcinoma of skin	6	TX, T1, T2, T3, T4	NX, NO, N1	MX, M0, M1	I, II, III, IV	
C632	7	TX, T1, T2, T3, T4	NX, N0, N1, N2, N3	M0, M1	I, II, III, IV	
Malignant melanoma of skin C44, C510, C609, C632	6	Extent of tumour is classified after excision: pTX, pTis, pT1, pT1a, pT1b, pT2, pT2a, pT2b, pT3, pT3a, pT3b, pT4, pT4a, pT4b	NX, NO, N1, N1a, N1b, N2, N2a, N2b, N2c, N3	MX, M0, M1, M1a, M1b, M1c	0, I, IA, IB, IIA, IIB, IIC, III, IIIA, IIIB, IIIC, IV	
	7	Extent of tumour is classified after excision: pTX, pTis, pT1, pT1a, pT1b, pT2, pT2a, pT2b, pT3, pT3a, pT3b, pT4, pT4a, pT4b	NX, NO, N1, N1a, N1b, N2, N2a, N2b, N2c, N3	M0, M1, M1a, M1b, M1c	0, I, IA, IB, IIA, IIB, IIC, III, IIIA, IIIB, IIIC, IV	

[§] In general, the classification applies to carcinomas, except for some topographies with specific morphologies that use separate classifications (*i.e.* appendix-carcinoid). There should be histological confirmation of the disease.

Topography	TNM edition	т	Ν	М	Stage grouping	
Merkel cell carcinoma of kin C44, C632	7	TX, T1, T2, T3, T4	NX, NO, N1, N1a, N1b, N2	M0, M1, M1a, M1b, M1c	I, IA, IB, IIA, IIB, IIC, IIIA, IIIB, IV	
Vulva C51	6	TX, T1, T1a, T1b, T2, T3, T4	NX, NO, N1, N2	MX, M0, M1	I, IA, IB, II, III, IVA, IVB	
	7	TX, T1, T1a, T1b, T2, T3	NX, NO, N1, N1a, N1b, N2, N2a, N2b, N2c, N3	M0, M1	I, IA, IB, II, IIIA, IIIB, IIIC, IVA, IVB	
Vagina	6	TX, T1, T2, T3, T4	NX, NO, N1	MX, M0, M1	I, II, III, IVA, IVB	
C52	7	TX, T1, T2, T3, T4	NX, NO, N1	M0, M1	I, II, III, IVA, IVB	
Cervix uteri C53	6	TX, Tis, T1, T1a, T1a1, T1a2, T1b, T1b1, T1b2, T2, T2a, T2b, T3, T3a, T3b, T4	NX, NO, N1	MX, M0, M1	0, IA, IA1, IA2, IB, IB1, IB2, IIA, IIB, IIIA, IIIB, IVA, IVB	
	7	TX, Tis, T1, T1a, T1a1, T1a2, T1b, T1b1, T1b2, T2, T2a, T2a1, T2a2, T2b, T3, T3a, T3b, T4	NX, NO, N1	M0, M1	0, I, IA, IA1, IA2, IB, IB1, IB2, II, IIA, IIA1, IIA2, IIB, III, IIIA, IIIB, IVA, IVB	
Corpus uteri C541, C55	6	TX, T1, T1a, T1b, T1c, T2, T2a, T2b, T3, T3a, T3b, T4	NX, NO, N1	MX, M0, M1	IA, IB, IC, IIA, IIB, IIIA, IIIB, IIIC, IVA, IVB	
	7	TX, T1, T1a, T1b, T2, T3, T3a, T3b, T4	NX, NO, N1	M0, M1	IA, IB, II, IIIA, IIIB, IIIC, IVA, IVB	
Uterine sarcoma C53, C540, C543 (8890/3, 8930/3, 8933/3)	7	8890/3, 8930/3 T1, T1a, T1b, T2, T2a, T2b, T3, T3a, T3b, T4 8933/3 T1, T1a, T1b, T1c, T2, T2a, T2b, T3, T3a, T3b, T4	NX, NO, N1	M0, M1	I, IA, IB, IC (only for 8993/3), II, IIA, IIB, IIIA, IIIB, IIIC, IVA, IVB	
Ovary C56	6	TX, T1, T1a, T1b, T1c, T2, T2a, T2b, T2c, T3, T3a, T3b	NX, NO, N1	MX, M0, M1	IA, IB, IC, IIA, IIB, IIC, IIIA, IIIB, IIIC, IV	
	7	TX, T1, T1a, T1b, T1c, T2, T2a, T2b, T2c, T3, T3a, T3b, T3c	NX, NO, N1	M0, M1	IA, IB, IC, IIA, IIB, IIC, IIIA, IIIB, IIIC, IV	

[§] In general, the classification applies to carcinomas, except for some topographies with specific morphologies that use separate classifications (*i.e.* appendix-carcinoid). There should be histological confirmation of the disease.

Topography	TNM edition	т	Ν	М	Stage grouping
Fallopian tube C570	6	TX, T1, T1a, T1b, T1c, T2, T2a, T2b, T2c, T3, T3a, T3b, T3c	NX, NO, N1	MX, MO, M1	IA, IB, IC, IIA, IIB, IIC, IIIA, IIIB, IIIC, IV
	7	TX, T1, T1a, T1b, T1c, T2, T2a, T2b, T2c, T3, T3a, T3b, T3c	NX, NO, N1	M0, M1	IA, IB, IC, IIA, IIB, IIC, IIIA, IIIB, IIIC, IV
Gestational trophoblastic tumours C58	6	TX, T1, T2	-	MX, M0, M1, M1a, M1b	I, IA, IB, II, IIA, IIB, III, IIIA, IIIB, IV, IVA, IVB
	7	TX, T1, T2	-	M0, M1, M1a, M1b	I, IA, IB, II, IIA, IIB, III, IIIA, IIIB, IV, IVA, IVB
Breast C50	6	TX, Tis, T1, T1mic, T1a, T1b, T1c, T2, T3, T4, T4a, T4b, T4c, T4d	NX, NO, N1, N2, N2a, N2b, N3, N3a, N3b, N3c Pathological N: pNX, pN0, pN1, pN1mi, pN1a, pN1b, pN1c, pN2, pN2a, pN2b, pN3, pN3a, pN3b, pN3c	MX, MO, M1	0, I, IIA, IIB, IIIA, IIIB, IIIC, IV
	7	TX, Tis, T0, T1, T1mi, T1a, T1b, T1c, T2, T3, T4, T4a, T4b, T4c, T4d	NX, NO, N1, N2, N2a, N2b, N3, N3a, N3b, N3c Pathological N: pNX, pNO, pN1, pN1mi, pN1a, pN1b, pN1c, pN2, pN2a, pN2b, pN3, pN3a, pN3b, pN3c	M0, M1	0, IA, IB, IIA, IIB, IIIA, IIIB, IIIC, IV
Penis	6	TX, T1, T2, T3, T4	NX, N0, N1, N2, N3	MX, M0, M1	I, II, III, IV
00	7	TX, T1, T1a, T1b, T2, T3, T4	NX, NO, N1, N2, N3	M0, M1	I, II, IIIA, IIIB, IV
Prostate C61	6	TX, T1, T1a, T1b, T1c, T2, T2a, T2b, T2c, T3, T3a, T3b, T4	NX, NO, N1	MX, M0, M1, M1a, M1b, M1c	I, II, III, IV
	7	TX, T1, T1a, T1b, T1c, T2, T2a, T2b, T2c, T3, T3a, T3b, T4	NX, NO, N1	M0, M1, M1a, M1b, M1c	I, II, III, IV

[§] In general, the classification applies to carcinomas, except for some topographies with specific morphologies that use separate classifications (*i.e.* appendix-carcinoid). There should be histological confirmation of the disease.

Topography	TNM edition	n T N M Sta		N M Stage groupi			M Stage grouping	
Testis C62	6	Extent of tumour → after radical orchiectomy pTX, pT1, pT2, pT3, pT4	NX, NO, N1, N2, N3	MX, M0, M1, M1a, M1b	I, IA, IB, IS, II, IIA, IIB, IIC, III, IIIA, IIIB, IIIC			
	7	Extent of tumour → after radical or- chiectomy, except for T4 pTX, pT1, pT2, pT3, pT4	NX, NO, N1, N2, N3	M0, M1, M1a, M1b	I, IA, IB, IS, II, IIA, IIB, IIC, III, IIIA, IIIB, IIIC			
Kidney C64	6	TX, T1, T1a, T1b, T2, T3, T3a, T3b, T3c, T4	NX, NO, N1, N2	MX, M0, M1	I, II, III, IV			
	7	TX, T1, T1a, T1b, T2, T2a, T2b, T3, T3a, T3b, T3c, T4	NX, NO, N1	M0, M1	I, II, III, IV			
Renal pelvis	6	TX, T1, T2, T3, T4	NX, N0, N1, N2, N3	MX, M0, M1	I, II, III, IV			
C65, C66	7	TX, T1, T2, T3, T4	NX, N0, N1, N2, N3	M0, M1	I, II, III, IV			
Urinary bladder C67	6	TX, Ta, Tis, T1, T2, T2a, T2b, T3, T3a, T3b, T4, T4a, T4b	NX, NO, N1, N2, N3	MX, M0, M1	0a, 0is, I, II, III, IV			
	7	TX, Ta, Tis, T1, T2, T2a, T2b, T3, T3a, T3b, T4, T4a, T4b	NX, NO, N1, N2, N3	M0, M1	0a, 0is, I, II, III, IV			
Urethra (C680) and transitional	6	TX, T1, T2, T3, T4	NX, NO, N1, N2	MX, M0, M1	I, II, III, IV			
cell carcinomas of prostate (C619)	7	TX, T1, T2, T3, T4	NX, NO, N1, N2	M0, M1	I, II, III, IV			

[§] In general, the classification applies to carcinomas, except for some topographies with specific morphologies that use separate classifications (*i.e.* appendix-carcinoid). There should be histological confirmation of the disease.

Topography	TNM edition	Т	Ν	М	Stage grouping
Thyroid gland C73	6	TX, T1, T2, T3, T4a, T4b	NX, NO, N1, N1a, N1b	MX, M0, M1	Papillary/Follicular <45 years I, II Papillary/Follicular 45 years and older Medullary I, II, III, IVA, IVB, IVC Anaplastic/ Undifferentiated IVA, IVB, IVC
	7	TX, T1, T1a, T1b, T2, T3, T4a, T4b	NX, NO, N1, N1a, N1b	M0, M1	Papillary/Follicular <45 years I, II Papillary/Follicular 45 years and Medullary I, II, III, IVA, IVB, IVC Anaplastic/ Undifferentiated IVA, IVB, IVC
Adrenal cortex tumours (C740)	7	TX, T1, T2, T3, T4	NX, NO, N1	M0, M1	I, II, III, IV

[§] In general, the classification applies to carcinomas, except for some topographies with specific morphologies that use separate classifications (*i.e.* appendix-carcinoid). There should be histological confirmation of the disease.

3. List of quality checks: internal consistency | 25

3.2. Consistency between variables

3.2.1. Coherence between dates

The following three dates are included in the list of agreed upon variables: *date of birth*, *date of incidence* and *date of the last known vital status*. The proposed rules below check for coherence between these dates:

- Date of birth ≤ Date of incidence. This rule is valid unless the case was diagnosed in utero if the diagnosis is in utero, the difference in months between dates should be no more than nine.
- Date of birth ≤ Date of the last known vital status.
- Date of incidence ≤ Date of the last known vital status.

For each comparison between dates, if years are known but month(s) is (are) unknown / missing, then the years are only compared. Similarly, if the day(s) is (are) unknown/missing, only the years and months are compared.

3.2.2. Consistency between tumour data and demographic information

• Consistency between age/topography/ morphology.

Some cancers occur almost exclusively in certain age groups such as retinoblastoma (tumour of young children) or prostate cancer in older men; therefore, some combinations age/topography/morphology are unlikely and should result in a warning.

Nevertheless there are other age and tumour type combinations that are not unlikely *but* rare. A warning for these combinations improves the precision of these rare tumours. After the confirmation by CRs, a flag will show that the case has been checked. *Table* **3** shows unlikely and rare combinations by age group and tumour type.

Table 3. Unlikely and rare combinations of age and tumour type.

Age group [years]	Morphology	Topography		
0-2	Hodgkin lymphoma: 9650-9667	Hodgkin lymphoma: 9650-9667		
> 9	Neuroblastoma and ganglioneuroblast	-		
> 5	Retinoblastoma: 9510-9514	-		
> 8	Wilms' tumour, rhabdoid, and clear	8960, 8964	-	
	cell sarcoma	8963	C649	
0-8	Renal carcinoma: 8010-8041, 8050-80 8143, 8155, 8190-8201, 8210, 8211, 8 8260-8263, 8290, 8310, 8320, 8323, 8 8510, 8550, 8560-8573	975, 8082, 8120-8122, 8130-8141, 3221-8231, 8240, 8241, 8244-8246, 3401, 8430, 8440, 8480-8490, 8504,	C649	
	8312		-	

Age group [years]	Morphology	Topography	
> 5	Hepatoblastoma: 8970		-
0-8	Hepatic carcinoma	8010-8041, 8050-8075, 8082, 8120-8122, 8140, 8141, 8143, 8155, 8190-8201, 8210, 8211, 8230, 8231, 8240, 8241, 8244-8246, 8260- 8263, 8310, 8320, 8323, 8401, 8430, 8440, 8480-8490, 8504, 8510, 8550, 8560-8573	C220,C221
0.5		8160-8180	-
0-5	Osteosarcomas: 9180-9187, 9192-919	5	-
0-5	Chondrosarcoma	9220-9230	
		9240	C400-C419
0-3	Ewing sarcoma: 9260, 9364		-
> 7	Malignant extra-cranial and extra-gona 9072, 9080-9085, 9100-9105	C00-C55, C57-C61, C63-C69, C73-C750, C754-C768, C80	
0-14	Gonadal carcinoma	8010-8041, 8050-8075, 8082, 8120- 8122, 8130-8141, 8143, 8190-8201, 8210, 8211, 8221-8241, 8244-8246, 8260-8263, 8290, 8310, 8320, 8323, 8380-8384, 8430, 8440, 8480-8490, 8504, 8510, 8550, 8560-8573, 9014, 9015	C56, C62
		8313, 8441, 8450, 8460-8471, 9000	-
0-5	Thyroid carcinoma	8010-8041, 8050-8075, 8082, 8120- 8122, 8130-8141, 8155, 8190, 8200, 8201, 8211, 8230, 8231, 8244-8246, 8260-8263, 8290, 8310, 8320, 8323, 8430, 8440, 8480, 8481, 8510, 8560-8573	C73
		8330-8337, 8340-8347, 8350	-
0-5	Nasopharyngeal carcinoma: 8010-804 8122, 8130-8141, 8190, 8200, 8201, 8 8263, 8290, 8310, 8320, 8323, 8430, 8	C11	
0-4	Skincarcinoma:8010-8041, 8050-8075 8143, 8147, 8190, 8200, 8240, 8246, 8 8420, 8430, 8480, 8542, 8560, 8570-8	C44	
0-4	Carcinoma, NOS: 8010-8084, 8120-815 8315, 8320-8325, 8380-8384, 8430-8 8589, 8940-8941, 9000, 9010-9016, 9	C00-C10, C12-C21, C23-C39, C48, C50-C55, C57-C61, C63, C65-C72, C75-C76, C80	
0-14	Mesothelial neoplasms: 9050-9053		Any

3. List of quality checks: internal consistency | 27

Age group [years]	Morphology	Topography
0-14	Any	C17, C25
0-14	Choriocarcinoma: 9100	Any
< 20	Any	C15, C19, C20, 21, C23, C24, C384, C50-C55
	Less than 9590 (Haematological malignancies)	C17
	Any other than carcinoid tumours (8240-8245)	C18, C33, C34
< 25	Multiple myeloma: 9732 and Chronic lymphocytic leukaemia: 9823	Any
< 30	Chronic myeloid leukaemia: 9876, 9945	Any
	Any	C60
< 40	Adenocarcinoma:8140	C61
> 45	Choriocarcinoma: 9100	C58
>14	8910, 8960, 8970, 8981, 8991, 9072, 9470, 951_, 9687	Any
	Juvenile myelomonocytic leukaemia: 9946	Any

• Consistency between sex/topography.

Some sex/topography combinations are impossible. Invalid combinations are presented in *Table 4*.

 Table 4. Invalid sex and topography combinations.

Sex=	1 (male)	Sex=	2 (female)
C51	Vulva	C60	Penis
C52	Vagina	C61	Prostate gland
C53	Cervix uteri	C62	Testis
C54	Corpus uteri	C63	Other and unspecified male genital organs
C55	Uterus, NOS		
C56	Ovary		
C57	Other and unspecified female genital organs		
C58	Placenta		

• Consistency between sex/morphology.

Table 5 includes a list of unlikely sex /morphology combinations.

 Table 5. Unlikely sex and morphology combinations.

Sex = 1	(male)	Sex= 2 (female)
8313/3	Clear cell adenocarcinofibroma	9061/3	Seminoma, NOS
8380/3	Endometrioid adenocarcinoma, NOS	9062/3	Seminoma, anaplastic
8381/3	Endometrioid adenofibroma, malignant	9063/3	Spermatocytic seminoma
8382/3	Endometrioid adenocarcinoma, secretory variant		
8383/3	Endometrioid adenocarcinoma, ciliated cell variant		
8384/3	Adenocarcinoma, endocervical type		
8441/3	Serous cystadenocarcinoma, NOS		
8460/3	Papillary serous cystadenocarcinoma		
8471/3	Papillary mucinous cystadenocarcinoma		
8482/3	Mucinous adenocarcinoma, endocervical type		
8600/3	Thecoma, malignant		
8670/3	Steroid cell tumour, malignant		
8930/3	Endometrial stromal sarcoma, NOS		
8931/3	Endometrial stromal sarcoma, low grade		
8934/3	Carcinofibroma		
8950/3	Mullerian mixed tumour		
8951/3	Mesodermal mixed tumour		
9000/3	Brenner tumour, malignant		
9014/3	Serous adenocarcinofibroma		
9015/3	Mucinous adenocarcinofibroma		
9090/3	Struma ovary, malignant		

3.2.3. Consistency between tumour variables

• Consistency between basis of diagnosis/morphology/behaviour.

It is unlikely for specific morphologies not to have undergone a histological/cytological examination. Nevertheless, some combinations are considered as exceptions. ENCR recommendations have been followed for 'specific' morphology codes in absence of microscopic verification. Morphology codes for cases with 'death certificate only' (DCO) are allowed when they can be identified from the underlying cause of death code (International Classification of Diseases 10th Revision).

Figure **2** shows the accepted combinations between basis of diagnosis (BoD) and morphology. Combinations not included in *Figure* **2** need to be verified.

Figure 2. Valid combinations for basis of diagnosis and morphology.



* Since the determination that a neoplasm has not invaded surrounding tissue (in situ) is made via the microscope, cases coded in situ (behaviour=2) should have a basis of diagnosis=7 or 5.

• Consistency between morphology/grade.

Only malignant tumours (behaviour = 3) should be graded. The combination between a 'behaviour' code less than 3 and a 'grade' code less than 9 will be considered as an error. This edit is skipped if 'grade' is blank or missing.

Grade values and the allowed corresponding morphology codes are shown in *Table* **6**.

Table 6.	Valid	comhinations	for	mornholoav	, and	arade
rubic 0.	vullu	combinations.	,0,	morphology	unu	gruue

Grade →	5	6	8
Morphology	9700-9702, 9705, 9708, 9709, 9714, 9716, 9717, 9718, 9719, 9724, 9725, 9726, 9729, 9800, 9801, 9805-9807, 9809, 9820, 9827, 9831, 9834, 9837	9591, 9596, 9597, 9670, 9671, 9673, 9678-9680, 9684, 9687-9691, 9695, 9698, 9699, 9712, 9728, 9731, 9732, 9734, 9737, 9738, 9762, 9800, 9801, 9805-9808, 9811, 9812- 9818, 9820, 9823, 9826, 9833, 9836, 9940	9719, 9727, 9831, 9948

The combination between grades 5-8 and morphology out of the range 9590-9992 is impossible. Some terms in ICD-O-3 carry an implied statement of grade; therefore an appropriate grade code needs to be associated.

These combinations are specified in the following *Table 7*.

 Table 7. Morphology code and description, and correct associated grade for ICD-0-3 terms with implied statement of grade.

Morphology code	Morphology description	Grade
8020/3	Carcinoma, undifferentiated, NOS	4
8021/3	Carcinoma, anaplastic, NOS	4
8240/3	Neuroendocrine carcinoma, well-differentiated	1
8249/3	Neuroendocrine carcinoma, moderately differentiated	2
8331/3	Follicular adenocarcinoma, well-differentiated	1
8332/3	Follicular adenocarcinoma, moderately differentiated	2
8585/3	Well-differentiated thymic carcinoma	1
8631/3	Sertoli-Leydig cell tumour, poorly differentiated	3
8634/3	Sertoli-Leydig cell tumour, poorly differentiated, with heterologous elements	3
8805/3	Undifferentiated sarcoma	4
8851/3	Liposarcoma, well-differentiated	1
9062/3	Seminoma, anaplastic	4
9082/3	Malignant teratoma, undifferentiated	4
9362/3	Pineal parenchymal tumour of intermediate differentiation	2, 3
9382/3	Anaplastic oligoastrocytoma	3
9390/3	Choroid plexus papilloma, anaplastic	3
9401/3	Astrocytoma, anaplastic	3
9440/3	Glioblastoma	4
9451/3	Oligodendroglioma, anaplastic	3
9511/3	Retinoblastoma, differentiated	1
9512/3	Retinoblastoma, undifferentiated	4

• Consistency between topography/laterality.

Laterality' that means 'bilateral and separated topographies' should be coded for those paired organs for which such information may be relevant for clinical or epidemiological reasons. Therefore, laterality has a valid code from 1 to 4 for only the following topographies:

List of paired organs for which it is suggested to collect laterality:

- Co7 Parotid gland • Co9 Tonsil • C300 Nasal cavity • C340, C341, Lung C343, C348, C349 • C384 Pleura • C400 Long bones of upper limb and scapula • C401 Short bones of upper limb • C402 Long bones of lower limb • C403 Short bones of lower limb • C413 Rib and clavicle • C414 Pelvic bones (excluding sacrum, coccyx, and symphysis pubis) • C441 Skin of eyelid • C442 Skin of external ear • C446 Skin of arm and shoulder • C447 Skin of leg and hip • C50 Breast • C56 Ovary • C570 Fallopian tube • C62 Testis • C630 Epididymis
- C649 Kidney

• C570	Fallopian tube
• C62	Testis
• C630	Epididymis
• C649	Kidney
• C659	Renal pelvis
• C66	Ureter
• C69	Eye
• C74	Suprarenal gland

Laterality is usually 1 for the topography C342, except for rare cases with situs inversus.

• Consistency between topography/morphology.

The topography/ morphology combinations include those morphologies commonly identified in specific primary topography (allowed topography codes) as well as the ones occurring only rarely or never in some specific primary topographies (not allowed topography codes). *Table* **8** reports allowed / refused combinations.

Table 8.	Morphology	codes and	allowed/refused	topography	codes
----------	------------	-----------	-----------------	------------	-------

Morphology codes	Allowed topography codes	Not allowed topography codes
8000-8005		C420, C421, C77
8010-8589		C38, C40-C42, C47, C480, C49, C70- C72, C77
8015	C53	
8077	C00-C15, C21, C30-C32, C44, C51- C53, C60	
8080	C51, C60	
8081	C00, C300, C44, C51, C60, C632, C690, C691	
8082	C00-C14, C16, C30-C34, C44, C53, C65-C68, C80	
8090-8095, 8097, 8100-8103, 8110	C300, C44, C51, C60, C632	
8098	C53	
8120, 8122, 8130, 8131	C56, C65-C68, C80	
8121	C300, C31, C65-C68	
8124	C212	
8142	C16	
8144	C15-C26, C30, C31, C52, C53, C56, C67, C80	
8145	C15-C20, C80	
8147	C00-C14, C30-C32, C50, C61	
8148	C15-C25, C61	
8150-8152, 8154, 8155	C25	
8153	C16, C170, C25, C80	
8156	C170, C25, C80	
8160, 8161	C221, C239, C240	
8162	C240	
8163	C22-C25	
8170-8175	C220	
8180	C221, C220	
8201	C15-C26, C50, C61, C80	
8210	C15-C26	
Morphology codes	Allowed topography codes	Not allowed topography codes
-------------------------------------------------	--------------------------------------------------	------------------------------
8211	C15-C26, C50, C61, C80	
8213	C18	
8214	C16	
8215	C211	
8220, 8221	C18-C20	
8243	C18, C56, C80	
8247	C300, C44, C51, C60, C632, C80	
8250-8254	C34	
8261, 8262	C15-C26, C52-C57	
8263	C15-C26, C52-C57, C64	
8265	C18-C20	
8270-8272, 8280, 8281, 8300	C751	
8290	C07, C08, C64, C73, C740, C751, C80	
8312, 8316-8320	C64	
8313	C56	
8314, 8315	C50	
8322	C750	
8330-8332, 8335-8337, 8340-8347, 8350	C73	
8370	C740	
8380-8383	C481, C482, C52-C57, C80	
8384	C53	
8390, 8400, 8402-8410, 8413	C300, C44, C51, C60, C632	
8401	C300, C44, C50, C51, C60, C632	
8420	C442	
8440	C07, C08, C25, C481, C482, C54, C56, C57, C80	
8441, 8460	C481, C482, C54, C56, C57, C80	
8442, 8444, 8450, 8451, 8461-8463, 8471-8473	C481, C482, C56, C57	
8452, 8453	C24, C25	

Morphology codes	Allowed topography codes	Not allowed topography codes
8470	C181, C25, C56, C57, C80	
8500	C07, C08, C24, C25, C50, C61, C80	
8501-8508, 8512-8514, 8520-8524, 8530, 8540, 8541, 8543	C50	
8510	C16, C18, C50, C80	
8525	C003-C005, C01-C08, C300, C31	
8542	C300, C44, C51, C60, C632	
8550, 8551	C003-C005, C01-C08, C25, C30-C34, C61, C80	
8580-8586	C37	
8588, 8589	C73	
8590-8650	C56, C62	
8670	C56	
8690, 8691	C755	
8692	C754	
8700	C741	
8710, 8711		C420, C421, C77
8720		C38, C40-C42, C47-C49, C77
8721-8723, 8730	C21, C300, C44, C51, C60, C632, C69, C80	
8728	C70	
8740, 8761	C44	
8741, 8743, 8745	C300, C44, C51, C60, C632, C690	
8742	C44, C51, C60, C632	
8744	C446, C447	
8746	C00-C06,C09-C11,C15,C20,C21,C30, C31,C680	
8770-8772	C300, C44, C51, C60, C632, C690, C80	
8773, 8774	C693, C694	
8780	C44	

Morphology codes	Allowed topography codes	Not allowed topography codes
8800-8811, 8814-8831, 8840-8921, 8963, 8990, 8991, 9040-9043, 9120- 9150, 9170, 9540, 9550, 9561, 9580, 9581		C420, C421, C77
8812	C40, C41	
8832, 8833	C44, C51, C60, C632	
8930, 8931	C481, C482, C52-C57	
8933, 8934	C52-C57	
8936	C15-C20, C25, C26, C481, C482, C80	
8940	C003-C005, C04-C08, C300, C44	
8941	C003-C005, C04-C08, C300	
8950, 8951	C481, C482, C52-C57, C80	
8959, 8960, 8964	C64	
8970	C220	
8971	C25	
8972, 8973	C34	
8983	C50	
9000	C56	
9013-9015	C481, C482, C56-C57, C80	
9020	C50	
9044	C49, C80	
9050-9053	C380, C384, C481, C482, C637, C80	
9060	C381-C383, C480, C56, C71, C751, C753	
9061-9063	C381-C383, C480, C62	
9064, 9065	C381-C383, C480, C495, C56, C62, C71, C751, C753, C80	
9070-9073, 9080-9085, 9101, 9102	C381-C383, C480, C495, C52-C57, C62, C71, C72, C751, C753, C80	
9090, 9091	C56	
9100	C381-C383, C480, C56-C58, C62, C80	
9104, 9105	C58	
9124	C220	

3. List of quality checks: internal consistency | 37

Morphology codes	Allowed topography codes	Not allowed topography codes
9161	C71-C72	
9180	C40, C41, C480, C49, C50, C80	
9181-9187, 9250	C40, C41	
9192-9195, 9221	C40	
9220, 9230, 9231, 9240-9243	C300, C31, C323, C33, C40, C41, C480, C49, C80	
9251, 9252	C49	
9260, 9364		С70-С72
9261	C400, C402	
9270-9342	C03, C310, C410, C411	
9350	C751, C752	
9351, 9352	C752	
9360-9362	C753	
9370-9372	C11, C41, C49	
9380-9384, 9391-9393, 9400-9431, 9440-9460	C71, C72, C753	
9390	C715	
9394	C72	
9395	C753	
9432	C751	
9470-9472, 9474, 9480, 9493	C716	
9490, 9500, 9503	C381-C383, C47, C480, C71-C72, C741, C755, C80	
9492, 9505-9509	C71, C72, C753	
9501, 9502	C694, C71	
9510-9513	C692	
9521-9523	C300, C31, C722	
9530-9539	C70	
9560	C38, C47, C480, C71-C72, C80	
9582	C751	

Morphology codes	Allowed topography codes	Not allowed topography codes
9590-9596, 9670-9675, 9680-9688, 9690-9699, 9702, 9705, 9714, 9724, 9728, 9729, 9735, 9737, 9738, 9750-9760, 9762		C420, C80
9597, 9700, 9709, 9718, 9725, 9726	C300, C44, C51, C60, C632	
9650-9667	C024, C09-C11, C14, C220, C421, C422, C77	
9678	C380, C384, C481, C482	
9679	C379, C381, C383, C771	
9689	C422	
9701	C421, C44, C77	
9708	C44, C49	
9712	C49	
9716	C220, C42	
9717	C16-C18	
9719	C01-C06, C09-C14, C30-C32, C44, C696, C77	
9727 (BPDCN)*	C421, C44	
9731	C40, C41	
9732, 9733, 9742, 9800-9826, 9831- 9920, 9931-9967, 9975-9989, 9991, 9992	C421	
9734		C40, C41, C420, C421, C80
9741	C220, C42, C44, C77	
9761	C420	
9764	C17	
9827	C421, C77	
9930		C420, C421, C80

* In ICD-O-3, 9727 was used for precursor cell lymphoblastic lymphoma, NOS; in the 2011 updates to ICD-O-3, 9727 is used for blastic plasmacytoid dendritic cell neoplasm (BPDCN). The topography codes allowed refer to BPDCN only.

3.3. Specific checks for survival analysis

Follow-up time and extent of disease are two important components to evaluate and interpret cancer survival. • Consistency of vital status/autopsy, autopsy/basis of diagnosis and autopsy/ survival/dates of incidence and follow-up.

Vital status = 1 (alive)	Autopsy = 0 (not incidentally diagnosed at autopsy)	
	Basis of diagnosis ≠ 0 (DCO)	
Basis of diagnosis = 0 (DCO)	Status = 2 (dead)	
	Survival (in days) = 0	
	Date of incidence = Date of the end of follow-up	
	Tumour size/number of examined and metastatic nodes/cTNM/pTNM/ condensed TNM/EOD/stage = unknown	
Autopsy = 1 (yes)	Status = 2 (dead)	
	Survival (in days) = 0	
	Date of incidence = Date of the end of follow-up	

3.4. Other additional checks on the extent of the disease

Several variables have been included in order to retrieve information about the extent of disease: tumour size, number of examined and metastatic nodes, cTNM, pTNM, condensed_TNM, EOD (summary extent of disease) and stage grouping. *Appendices II* and *III* contain detailed TNM6 and TNM7 stage grouping, respectively, and corresponding T, N, M values.

Table **1** and *Table* **2** provide accepted values for T, N, M and stage grouping. Furthermore, the following additional checks are proposed to identify inconsistencies among variables related to survival:

- If C70, C71, C76, C42, C77 and C80, then the number of examined and metastatic nodes must be '99'.
- If TNM pT = Tis, then 'basis of diagnosis' = 7.
- TNM $cT \neq Tis$.
- If site = C80, tumour size = 999.9.
- If TNM pT ≠ TX or ≠ 999999, then 'basis of diagnosis' = 7.
- If TNM $pN \neq NX$ or \neq 999999, then 'basis of diagnosis' = 5 or 7.
- If TNM pM ≠ MX or ≠ 999 then 'basis of diagnosis' = 5 or 7 or 6.
- Inconsistencies between topographies and summary extent of disease (EOD)
 Topography = C809 and (TNM N = 0, TNM M = 0).
 Topography = C809 and (condensed N

= No, condensed M = Mo).

```
Topography: the fourth digit of the
      topography is = 8 and EOD = 1.
    Topography = Co69 and EOD = 1.
    Topography = C_{26} and EOD = 1.
    Topography = C_{39} and EOD = 1.
    Topography = C409 and EOD = 1.
    Topography = C419 and EOD = 1.
    Topography = C479 and EOD = 1.
    Topography = C499 and EOD = 1.
    Topography = C_{559} and EOD = 1.
    Topography = C_{579} and EOD = 1.
    Topography = C639 and EOD = 1.
    Topography = C809 and EOD = 1.
    Topography = C76 and EOD = 1.
    Topography = C_{77} and TNM N = NO.
    Topography = C_{77} and condensed N
      = No.
• Inconsistencies between behaviour and TNM/
  EOD
    Behaviour > 2 and pTNM T = Tis.
```

Behaviour = 6 and cTNM M = Mo.
Behaviour = 6 and pTNM M = Mo.
Behaviour = 6 and condensed M = Mo.

```
Behaviour = 6 and EOD = 1.
Behaviour = 6 and EOD = 2
```

```
(excluding nodes).
```

 Inconsistencies between EOD = 1 and TNM EOD = 1 and TNM N ≠ NO. EOD = 1 and TNM M ≠ MO.

- Inconsistencies between EOD = 1 and condensed TNM
 EOD = 1 and condensed T ≠ TL.
- EOD = 1 and condensed $N \neq N0$.
- EOD = 1 and condensed $M \neq MO$.
- Inconsistencies between EOD = 2 and TNM EOD = 2 and $TNM M \neq MO$. EOD = 2 and (pTNM T = Tis or TNM T = T1).
 - EOD = 2 and TNM N = NO.
- Inconsistencies between EOD = 2 and condensed TNM
- EOD = 2 and condensed M = M1.
- Inconsistencies between EOD = 3 and TNM EOD = 3 and TNM M = MO.
- Inconsistencies between EOD = 3 and condensed TNM
 EOD = 3 and condensed M = MO.
- Inconsistencies between TNM and condensed TNM
 TNM N ≠ NO and condensed N = NO.
 TNM N = NO and condensed N = N1.
 TNM M = MO and condensed M = M1.
 TNM M ≠ MO and condensed M = M0.
- Inconsistencies between N+ and stage Number of metastatic nodes > 0 and TNM N = NO.
 Number of metastatic nodes > 0 and condensed N = NO.

Number of metastatic nodes > 0 and EOD = 1.

Quality Checklist for Multiple Primary Malignant Tumours

The quality checklist of warnings for Multiple Primary Malignant (MPMTs) was developed by the JRC according to the current International Rules for Multiple Primary Cancers published in 2004 (<u>http://www.encr.eu/images/docs</u> /recommendations/MPrules_july2004.pdf).

This checklist has been addressed by the JRC at a later stage than for version 1.0 of this report, and is therefore included in the current version 1.1 to clarify the checks and the warning messages given by the JRC-ENCR Cancer Registries Data Quality Check Software. are the following:

Step 1. The two topographies are compared according to the current International Rules for Multiple Malignant Cancers published in 2004.

In addition to the groups of topography codes considered as a single site in table 1 of the 2004 international rules, **for checking** other groups are considered as a single topography (see *Table 9*).

C80 (unknown primary site) and C768 (overlapping lesion of ill-defined sites) are considered as a single site with any topography.

The steps for checking solid MPMTs

Definition Topography code C00 Lip C03 Gum C04 Floor of mouth C05 Palate C06 Other and unspecified parts of mouth C760 Head, face or neck, NOS C01 Base of tongue Other and unspecified parts of tongue C02 Head, face or neck, NOS C760 C07 Parotid gland C760 Head, face or neck, NOS C08 Other and unspecified major salivary glands Head, face or neck, NOS C760 C09 Tonsil C10 Oropharynx C12 Pyriform sinus C13 Hypopharynx C14 Other and ill-defined sites in lip, oral cavity and pharynx C760 Head, face or neck, NOS C11 Nasopharvnx C760 Head, face or neck, NOS

Table 9. Groups of topography codes considered as a single site for solid tumours

Note: topography codes C80 and C768 are considered as a single site in combination with any other topography.

Topography code	Definition
C15	Oesophagus
C268	Overlapping lesion of digestive system
C269	Gastrointestinal tract, NOS
C761	Thorax, NOS
C16	Stomach
C268	Overlapping lesion of digestive system
C269	Gastrointestinal tract, NOS
C762	Abdomen, NOS
C17	Small intestine
C26	Other and ill-defined digestive organs
C762	Abdomen, NOS
C18	Colon
C26	Other and ill-defined digestive organs
C762	Abdomen, NOS
C19	Rectosigmoid junction
C20	Rectum
C26	Other and ill-defined digestive organs
C762	Abdomen, NOS
C21	Anus and anal canal
C26	Other and ill-defined digestive organs
C76	Other and ill-defined sites
C22	Liver and intrahepatic bile ducts
C268	Overlapping lesion of digestive system
C762	Abdomen, NOS
C23	Gallbladder
C24	Other and unspecified parts of biliary tract
C268	Overlapping lesion of digestive system
C269	Gastrointestinal tract, NOS
C762	Abdomen, NOS
C25	Pancreas
C268	Overlapping lesion of digestive system
C762	Abdomen, NOS
C30	Nasal cavity and middle ear
C39	Other and ill-defined sites within respiratory system and intrathoracic organs
C760	Head, face or neck, NOS
C31	Accessory sinuses
C39	Other and ill-defined sites within respiratory system and intrathoracic organs
C760	Head, face or neck, NOS
C32	Larynx
C39	Other and ill-defined sites within respiratory system and intrathoracic organs
C760	Head, face or neck, NOS
C33	Trachea
C34	Bronchus and lung
C39	Other and ill-defined sites within respiratory system and intrathoracic organs
C760	Head, face or neck, NOS
C761	Thorax, NOS
C37	Thymus
C761	Thorax, NOS

Note: topography codes C80 and C768 are considered as a single site in combination with any other topography.

4. Quality Checklist for Multiple Primary Malignant Tumours | 43

Topography code	Definition
C38	Heart, mediastinum, and pleura
C398	Overlapping lesion of respiratory system and intrathoracic organs
C761	Thorax, NOS
C39	Other and ill-defined sites within respiratory system and intrathoracic organs
C760	Head, face or neck, NOS
C761	Thorax, NOS
C40	Bones, joints and articular cartilage of limbs
C41	Bones, joints and articular cartilage of other and unspecified sites
C76	Other and ill-defined sites
C44	Skin
C76	Other and ill-defined sites
C47	Peripheral nerves and autonomic nervous system
C76	Other and ill-defined sites
C48	Retroperitoneum and peritoneum
C76	Other and ill-defined sites
C49	Connective, subcutaneous and other soft tissues
C76	Other and ill-defined sites
C50	Breast
C761	Thorax, NOS
C51	Vulva
C578	Overlapping lesion of female genital organs
C579	Female genital tract, NOS
C763	Pelvis, NOS
C52	Vagina
C578	Overlapping lesion of female genital organs
C579	Female genital tract, NOS
C763	Pelvis, NOS
C53	Cervix uteri
C55	Uterus, NOS
C578	Overlapping lesion of female genital organs
C579	Female genital tract, NOS
C763	Pelvis, NOS
C54	Corpus uteri
C55	Uterus, NOS
C578	Overlapping lesion of female genital organs
C579	Female genital tract, NOS
C763	Pelvis, NOS
C56	Ovary
C578	Overlapping lesion of female genital organs
C579	Female genital tract, NOS
C763	Pelvis, NOS
C57	Other and unspecified female genital organs
C763	Pelvis, NOS
C58	Placenta
C763	Pelvis, NOS
C60	Penis
C638	Overlapping lesion of male genital organs
C639	Male genital organs, NOS
C763	Pelvis, NOS

Note: topography codes C80 and C768 are considered as a single site in combination with any other topography.

Topography code	Definition
C61	Prostate gland
C638	Overlapping lesion of male genital organs
C639	Male genital organs, NOS
C763	Pelvis, NOS
C62	Testis
C638	Overlapping lesion of male genital organs
C639	Male genital organs, NOS
C763	Pelvis, NOS
C63	Other and unspecified male genital organs
C763	Pelvis, NOS
C64	Kidney
C688	Overlapping lesion of urinary organs
C689	Urinary system, NOS
C76	Other and ill-defined sites
C65	Renal pelvis
C66	Ureter
C67	Bladder
C68	Other and unspecified urinary organs
C76	Other and ill-defined sites
C69	Eye and adnexa
C760	Head, face or neck, NOS
C70	Meninges
C760	Head, face or neck, NOS
C71	Brain
C760	Head, face or neck, NOS
C72	Spinal cord, cranial nerves, and other parts of central nervous system
C760	Head, face or neck, NOS
C73	Thyroid
C760	Head, face or neck, NOS
C74	Adrenal gland
C76	Other and ill-defined sites
C75	Other endocrine glands and related structures
C76	Other and ill-defined sites

Note: topography codes C80 and C768 are considered as a single site in combination with any other topography.

- a) If the two topographies are in the same group, the two morphologies are compared (**step 2**).
- b) If the two topographies are in different groups, no warning message is given.

Step 2. The two morphologies are compared according to the 2004 International Rules.

Some unspecified morphologies were included **for checking** in the groups defined in table 2 of the 2004 International Rules, and are considered as the same morphology (*Table 10*). Morphology codes 8000-8005 (Unspecified types of cancer) are considered as a single group in combination with any other morphology.

Note: For haematological malignancies, only morphologies are compared.

 Table 10. Groups of morphology codes considered as a single entity

Morphology code	Definition
8051-8084, 8120-8131 8010-8015, 8020-8022, 8050	Squamous and transitional cell carcinoma Unspecified carcinomas (NOS)
8090-8110 8010-8015, 8020-8022, 8050	Basal cell carcinomas Unspecified carcinomas (NOS)
8140-8149, 8160-8163, 8190-8221, 8260-8337, 8350-8552, 8570-8576, 8940-8941	Adenocarcinomas
8010-8015, 8020-8022, 8050	Unspecified carcinomas (NOS)
8030-8046, 8150-8157, 8170-8180, 8230-8255, 8340-8347, 8560-8562, 8580-8671 8010-8015, 8020-8022, 8050	Other specific carcinomas
8680-8713, 8800-8921, 8990-8991, 9040-9044, 9120-9125, 9130-9136, 9141-9252, 9370-9373, 9540-9582	Sarcomas and soft tissue tumours
9050-9055	Mesothelioma
9840, 9861-9931, 9945-9946, 9950, 9961-9967, 9980-9987, 9991-9992	Myeloid
9590-9591, 9596, 9727, 9760, 9800-9801, 9805- 9809, 9820, 9832, 9835, 9860, 9960, 9970, 9971, 9975, 9989	Unspecified types of haematopoietic and lymphoid tissues
9597, 9670-9699, 9712, 9728, 9731-9738, 9761- 9767, 9769, 9811-9818, 9823, 9826, 9833, 9836, 9940	B-cell neoplasms
9590-9591, 9596, 9727, 9760, 9800-9801, 9805- 9809, 9820, 9832, 9835, 9860, 9960, 9970, 9971, 9975, 9989	Unspecified types of haematopoietic and lymphoid tissues
9700-9709, 9714-9719, 9724-9726, 9729, 9768, 9827, 9831, 9834, 9837, 9948	T-cell and NK-cell neoplasms
9590-9591, 9596, 9727, 9760, 9800-9801, 9805- 9809, 9820, 9832, 9835, 9860, 9960, 9970, 9971, 9975, 9989	Unspecified types of haematopoietic and lymphoid tissues
9650-9667 9590-9591, 9596, 9727, 9760, 9800-9801, 9805- 9809, 9820, 9832, 9835, 9860, 9960, 9970, 9971, 9975, 9989	Hodgkin lymphoma Unspecified types of haematopoietic and lymphoid tissues
9740-9742 9590-9591, 9596, 9727, 9760, 9800-9801, 9805- 9809, 9820, 9832, 9835, 9860, 9960, 9970, 9971, 9975, 9989	Mast-cell tumours Unspecified types of haematopoietic and lymphoid tissues
9750-9759 9590-9591, 9596, 9727, 9760, 9800-9801, 9805- 9809, 9820, 9832, 9835, 9860, 9960, 9970, 9971, 9975, 9989	Histiocytes and Accessory Lymphoid cells Unspecified types of haematopoietic and lymphoid tissues
9140	Kaposi sarcoma
8720-8790, 8930-8936, 8950-8983, 9000-9030, 9060-9110, 9260-9365, 9380-9539	Other specified types of cancer

Note: morphology codes 8000-8005 (unspecified types of cancer) are considered as a single group in combination with any other morphology.



- Allemani C, Weir HK, Carreira H et al. and the CONCORD Working Group. Global surveillance of cancer survival 1995-2009: analysis of individual data for 25 676 887 patients from 279 populationbased registries in 67 countries (CON-CORD-2). Lancet, 2015, 385(9972):977-1010.
- Anatomical Therapeutic Chemical (ATC) classification system. WHO Collaborating Centre for Drug Statistics. Available in http://www.whocc.no/atc_ddd_index/.
- Bray F, Parkin DM. Evaluation of data quality in the cancer registry: Principles and methods. Part I: Comparability, validity and timeliness. *Eur J Cancer*, 2009, 45: 747-755.
- De Angelis R, Francisci S, Baili P *et al.* The EUROCARE-4 database on cancer survival in Europe: Data standardisation, quality control and methods of statistical analysis. *EurJ Cancer*, 2009, 45: 909-930.
- ENCR Recommendations. Available in http:// www.encr.eu/working-groups-and-recom mendations.
- EUROCARE. Checking procedures of the EUROCARE-4 Data Base. Available in http://www.eurocare.it/Eurocare4DataC hecking/tabid/81/Default.aspx.
- Ferlay J, Burkhard C, Whelan S, Parkin DM. Check and conversion programs for cancer registries (LARC/LACR tools for cancer registries). International Agency for Research on Cancer/International Association of Cancer Registries, IARC

Technical Report No. 42, Lyon, 2005.

- Ferretti S, Giacomin A and Airtum Working Group. *Cancer registration handbook*. Available in http://www.registritumori.it/cms/pubblicazioni/cancerregistration-handbook-2010.
 - North American Association of Central Cancer Registries. NAACCR V14 Metafile Edit Detail Report– November 26, 2013. Available in http://www.naaccr.org/wp-content/ uploads/2016/11/NAACCR-v14-Beta-Edits-Detail-Report.pdf.
- Parkin DM, Bray F. Evaluation of data quality in the cancer registry: Principles and methods. Part II: Completeness. *EurJ Cancer*, 2009, 45:756-764.
- Percy C, Fritz A, Jack A, Shanmugarathan S, Sobin L, Parkin DM, Whelan S. *International Classification of Diseases for Oncology (ICD-O).* World Health Organization, 3rd ed., December 2000.
- Sobin LH, Gospodarowicz MK, Wittekind Ch, eds. TNM Classification of Malignant Tumors. 7th ed., Wiley-Blackwell, Oxford, 2009.
- Sobin LH, Wittekind Ch. TNM Classification of Malignant Tumours. 6th ed., John Wiley & Sons, Hoboken, New Jersey, 2002.
- *TNM Classification Help.* Manual for Cancer Staging. Available in cancerstaging. blogspot.com

Appendix I: *The Anatomical Therapeutic Chemical* (ATC) code, generic and trade names

Chemotherapy

ATC code	Generic name	Trade name
L01XX32	Bortezomib	VELCADE
L01XX17	Topotecan	HYCAMTIN
		TOPOTECAN
L01XX19	Irinotecan	IRINOTECAN HYDROCH
		САМРТО
		IRINOTECAN ACTAVIS
L01XX05	Hydroxycarbamide	HYDREA
L01XX14	Tretinoin	VESANOID
L01XX02	Asparaginase	KIDROLASE
LO1XX11	Estramustine	ESTRACYT
L01XA03	Oxaliplatin	ELOXATIN
		OXALIPLATINE ACTAV
		OXALIPLATINE
L01XA02	Carboplatin	CARBOPLATIN
		PARAPLATIN
L01XA01	Cisplatin	SINPLATIN
		CISPLATIN
		PLATIDIAM
		PLATINEX
L01BC06	Capecitabine	XELODA
L01BC05	Gemcitabine	GEMZAR
L01BC02	Fluorouracil	FLUOROURACIL ACCOR
		FLUOROURACIL 5
		5-FU
		FLUOROURACIL

Chemotherapy (cont.)

ATC code	Generic name	Trade name
L01BC01	Cytarabine	ALEXAN
		CYTARABINE
		CYTOSAR
L01BC53	Tegafur combinations	UFT
L01BA04	Pemetrexed	ALIMTA
L01BA01	Methotrexate	METHOTREXATE
L01BA03	Raltitrexed	TOMUDEX
L01BB05	Fludarabine	FLUDARA
L01BB02	Mercaptopurine	PURI NETHOL
L01BB07	Nelarabine	ATRIANCE
L01BB04	Cladribine	LITAK10
L01BB03	Tioguanine	LANVIS
L01CD02	Docetaxel	TAXOTERE
		DOCETAX
L01CD01	Paclitaxel	SINDAXEL
		PACLITAXEL
		TAXOL
		GENEXOL
		PACLITEVA
		PACLITAXIN
L01CA04	Vinorelbine	VINORELBIN ACTAVIS
		NAVELBIN
		VINORELBIN EBEWE
L01CA02	Vincristine	CYTOCRISTIN
		VINCRISTIN
L01CA01	Vinblastine	CYTOBLASTIN
		VINBLASTIN

Appendix I: The Anatomical Therapeutic Chemical code, generic name of the drug and trade names | 49

Chemotherapy (cont.)

ATC code	Generic name	Trade name
L01CB01	Etoposide	ETOSID
		ETOPOSIDE
		LASTET
		VEPESID
L01CB02	Teniposide	VUMON
L01CX01	Trabectedin	YONDELIS
L01AX03	Temozolomide	TEMODAL
L01AX04	Dacarbazine	DACARBAZIN
L01AA06	Ifosfamide	HOLOXAN
L01AA01	Cyclophosphamide	ENDOXAN
L01AA03	Melphalan	ALKERAN
L01AA02	Chlorambucil	LEUKERAN
L01AD01	Carmustine	BCNU
L01AD02	Lomustine	CCUN
L01AB01	Busulfan	MYLERAN
L01DB03	Epirubicin	FARMORUBICIN
		EPIRUBICIN
		EPILEM
		EPISINDAN
L01DB06	Idarubicin	ZAVEDOS
L01DB07	Mitoxantrone	MITOXANTRON
		ONCOTRONE
		NOVANTRONE
L01DB01	Doxorubicin	DOXORUBICIN
		CAELYX
L01DC03	Mitomycin	MITOMYCIN C
L01DC01	Bleomycin	BLEOCIN
L01DA01	Dactinomycin	COSMEGEN LYOVAC

Hormonal therapy

ATC code	Generic name	Trade name
L02BG04	Letrozole	FEMARA
L02BG03	Anastrozole	ARIMIDEX
		ANAROMAT
L02BG06	Exemestane	AROMASIN
L02BG01	Aminogluthetimide	AMINOGLUTETHIMID
		ORIMETEN
L02BA03	Fulvestrant	FASLODEX
LO2BA01	Tamoxifen	NOLVADEX
		TAMOXIFEN
		TAMIFEN
L02BB03	Bicalutamide	BICUSAN
		CASODEX
L02BB01	Flutamide	FLUTASIN
		FLUCINOM
		FLUTAMIDE
LO2AEO3	Goserelin	ZOLADEX
L02AE02	Leuprorelin	LUCRIN DEPOT
		ELIGARD
L02AE01	Buserelin	SUPREFACT
L02AE04	Triptorelin	DECAPEPTYL
		DIPHERELINE
		DIPHERELINE SR
LO2AB01	Megestrol	MEGACE
L02AB02	Medroxyprogesterone	MEDROXYPROGESTERON
		MPA
G03HA01	Cyproterone	ANDROCUR
G03DA02	Medroxyprogesterone	FARLUTAL

Appendix I: The Anatomical Therapeutic Chemical code, generic name of the drug and trade names | 51

Targeted therapy			
ATC code	Generic name	Trade name	
L01XC03	Trastuzumab	HERCEPTIN	
L01XC02	Rituximab	MABTHERA	
L01XC07	Bevacizumab	AVASTIN	
L01XC06	Cetuximab	ERBITUX	
L01XC08	Panitumumab	VECTIBIX	
L01XC04	Alemtuzumab	MABCAMPATH	
L01XE01	Imatinib	GLIVEC	
L01XE04	Sunitinib	SUTENT	
L01XE03	Erlotinib	TARCEVA	
L01XE05	Sorafenib	NEXAVAR	
L01XE07	Lapatinib	TYVERB	
L01XE06	Dasatinib	SPRYCEL	
L01XE08	Nilotinib	TASIGNA	

Systemic therapies other than chemotherapy and hormonal therapy

Immunotherapy			
ATC code	Generic name	Trade name	
L03AA13	Pegfilgrastim	NEULASTA	
L03AA02	2 Filgrastim		
		TEVAGRASTIM	
L03AA10	Lenograstim	GRANOCYTE	
L03AB01	Interferon alfa natural	ROFERON A	
L03AB05	Interferon alfa-2b	REALDIRON	
		INTRON A	

Other therapies			
ATC code	Generic name	Trade name	
A04AA01 Ondansetron	ZOFRAN		

Other therapies <i>(cont.)</i>			
ATC code	Generic name	Trade name	
A04AA02	Granisetron	KYTRIL	
		RASETRON	
A04AA04	Dolasetron	ANZEMET	
A04AA05	Palonosetron	ALOXI	
A04AA03	Tropisetron	NAVOBAN	
B03XA01	Erythropoietin	NEO RECORMON	
		EPREX	
B03XA03	Pegzerepoetinalfa	MIRCERA	
B03XA02	Darbepoetinalfa	ARANESP	
M05BA08	Zoledronic acid	ZOMETA	
M05BA06	Ibandronic acid	BONDRONAT	
M05BA03	Pamidronic acid	PAMITOR	
		AREDIA	
M05BA02	Clodronic acid	SINDRONAT	
		BONEFOS	
		OSTAC	
H02AB07	Prednisone	DEHYDROCORTISON	
H02AB06	Prednisolone	PREDNISOLON CORTIC	
		PREDNISOLON	
H02AB02	Dexamethasone	DEXAMETHASONE	
		DEXAVEN	
		PREDNISOLON F	
H02AB04	Methylprednisolone	METHYLPREDN.SOPHAR	
		SOLU MEDROL	
		DEPO MEDROL	
		MEDROL	
		METHYLPREDN.CORTIC	
		METHYLPREDNISOLON	

Systemic therapies other than chemotherapy and hormonal therapy (cont.)

Appendix I: The Anatomical Therapeutic Chemical code, generic name of the drug and trade names | 53

Appendix II: TNM 6 edition stage grouping and corresponding T, N, M values

Li	p and Oral Cavity COO, CO	02-C06 (except C051 and C05	2)
Stage	т	Ν	М
0	Tis	NO	МО
1	Τ1	NO	MO
Ш	Т2	NO	MO
III	T1,T2	N1	MO
	Т3	N0, N1	МО
IVA	T1, T2, T3	N2	MO
	T4a	N0, N1, N2	MO
IVB	Any T	N3	MO
	T4b	Any N	MO
IVC	Any T	Any N	M1

Oropharynx and Hyp	oopharynx C01, C051, C05	52, C090, C091, C099, C100, C	102, C103, C12, C13
Stage	т	Ν	М
0	Tis	NO	MO
1	Τ1	NO	МО
Ш	Τ2	NO	МО
ш	T1,T2	N1	MO
	Т3	N0, N1	МО
IVA	T1, T2, T3	N2	МО
	T4a	N0, N1, N2	МО
IVB	T4b	Any N	MO
	Any T	N3	МО
IVC	Any T	Any N	M1

Nasopharynx C11				
Stage	т	Ν	М	
0	Tis	NO	MO	
1	Τ1	NO	MO	
IIA	T2a	NO	MO	
IIB	Τ1	N1	MO	
	T2a	N1	MO	
	T2b	N0, N1	MO	
III	Τ1	N2	MO	
	T2a, T2b	N2	MO	
	Т3	N0, N1, N2	MO	
IVA	T4	N0, N1, N2	MO	
IVB	Any T	N3	МО	
IVC	Any T	Any N	M1	

Larynx C320, C321, C322, C101			
Stage	Т	Ν	М
0	Tis	NO	МО
1	Т1	NO	МО
Ш	Τ2	NO	МО
Ш	T1,T2	N1	МО
	Т3	N0, N1	МО
IVA	T1, T2, T3	N2	МО
	T4a	N0, N1, N2	МО
IVB	T4b	Any N	МО
	Any T	N3	МО
IVC	Any T	Any N	M1

Nasal Cavity and Paranasal Sinuses C300, C310, C311			
Stage	Т	Ν	М
0	Tis	NO	МО
1	Т1	NO	МО
Ш	Т2	NO	МО
Ш	T1,T2	N1	МО
	Т3	N0, N1	MO
IVA	T1, T2, T3	N2	МО
	T4a	N0, N1, N2	MO
IVB	T4b	Any N	МО
	Any T	N3	МО
IVC	Any T	Any N	M1

Salivary Glands C07, C08			
Stage	т	Ν	М
1	Τ1	NO	МО
Ш	Т2	NO	MO
Ш	Т3	NO	MO
	T1, T2, T3	N1	MO
IVA	T1, T2, T3	N2	MO
	T4a	N0, N1, N2	MO
IVB	T4b	Any N	MO
	Any T	N3	MO
IVC	Any T	Any N	M1

Thyroid Gland C73			
Stage	т	Ν	М
	Papillary or Follicu	lar, under 45 years	
1	Any T	Any N	МО
Ш	Any T	Any N	M1
	Papillary or Follicular, 45 yea	ars and older, and Medullary	
1	Τ1	NO	МО
Ш	Т2	NO	MO
ш	Т3	NO	МО
	T1, T2, T3	Nla	МО
IVA	T1, T2, T3	N1b	MO
	T4a	N0, N1	МО
IVB	T4b	Any N	МО
IVC	Any T	Any N	M1
Anaplastic/Undifferentiated (all cases are stage IV)			
IVA	T4a	Any N	МО
IVB	T4b	Any N	МО
IVC	Any T	Any N	M1

Oesophagus C15				
Stage	т	Ν	М	
0	Tis	NO	МО	
1	Τ1	NO	MO	
IIA	Т2, Т3	NO	МО	
IIB	T1,T2	N1	MO	
Ш	Т3	N1	MO	
	T4	Any N	MO	
IV	Any T	Any N	M1	
IVA	Any T	Any N	Mla	
IVB	Any T	Any N	M1b	

Stomach C16			
Stage	т	Ν	М
0	Tis	NO	МО
IA	Τ1	NO	MO
IB	Τ1	N1	МО
	T2a/b	NO	MO
Ш	Τ1	N2	MO
	T2a/b	N1	МО
	Т3	NO	МО
IIIA	T2a/b	N2	МО
	Т3	N1	МО
	T4	NO	МО
IIIB	Т3	N2	МО
IV	T4	N1, N2, N3	МО
	T1, T2, T3	N3	МО
	Any T	Any N	M1

Small Intestine C17				
Stage	Т	Ν	М	
0	Tis	NO	МО	
1	T1, T2	NO	МО	
Ш	T3, T4	NO	МО	
Ш	Any T	N1	MO	
IV	Any T	Any N	M1	

Colon and Rectum C18-C20				
Stage	Т	Ν	М	
0	Tis	NO	МО	
1	T1,T2	NO	MO	
IIA	Т3	NO	МО	
IIB	Τ4	NO	MO	
IIIA	T1,T2	N1	MO	
IIIB	T3, T4	N1	MO	
IIIC	Any T	N2	MO	
IV	Any T	Any N	M1	

Anal Canal C211				
Stage	Т	Ν	М	
0	Tis	NO	МО	
1	Т1	NO	MO	
Ш	Т2	NO	MO	
	Т3	NO	MO	
IIIA	T1, T2, T3	N1	MO	
	Τ4	NO	MO	
IIIB	Τ4	N1	МО	
	Any T	N2, N3	МО	
IV	Any T	Any N	M1	

Liver C220, C221				
Stage	т	Ν	М	
1	Τ1	NO	MO	
Ш	Т2	NO	МО	
IIIA	Т3	NO	МО	
IIIB	T4	NO	МО	
шс	Any T	N1	мо	
IV	Any T	Any N	M1	

Appendix II: TNM 6 edition stage grouping and corresponding T, N, M values | 59

Gallbladder C23				
Stage	Т	Ν	М	
0	Tis	NO	MO	
IA	Т1	NO	MO	
IB	Т2	NO	МО	
IIA	Т3	NO	MO	
IIB	T1, T2, T3	N1	MO	
Ш	T4	Any N	МО	
IV	Any T	Any N	M1	

Extrahepatic Bile Ducts C240				
Stage	т	Ν	М	
0	Tis	NO	МО	
IA	Т1	NO	MO	
IB	Т2	NO	МО	
IIA	Т3	NO	MO	
IIB	T1, T2, T3	N1	МО	
ш	T4	Any N	МО	
IV	Any T	Any N	M1	

Ampulla of Vater C241				
Stage	т	Ν	М	
0	Tis	NO	МО	
IA	Τ1	NO	МО	
IB	Τ2	NO	МО	
IIA	Т3	NO	МО	
IIB	T1, T2, T3	N1	МО	
Ш	T4	Any N	MO	
IV	Any T	Any N	M1	

Pancreas C25				
Stage	Т	Ν	М	
0	Tis	NO	MO	
IA	Т1	NO	MO	
IB	Τ2	NO	МО	
IIA	Т3	NO	MO	
IIB	T1, T2, T3	N1	MO	
ш	Τ4	Any N	MO	
IV	Any T	Any N	Ml	

Lung C34				
Stage	т	Ν	М	
Occult carcinoma	ТХ	NO	МО	
0	Tis	NO	MO	
IA	Τ1	NO	МО	
IB	Т2	NO	MO	
IIA	Τ1	N1	МО	
IIB	Τ2	N1	МО	
	Т3	NO	MO	
IIIA	T1, T2	N2	МО	
	Т3	N1, N2	МО	
IIIB	Any T	N3	МО	
	T4	Any N	MO	
IV	Any T	Any N	M1	

Pleural Mesothelioma C384				
Stage	Т	Ν	М	
IA	Tla	NO	MO	
IB	T1b	NO	MO	
Ш	Т2	NO	MO	
Ш	T1,T2	N1	MO	
	T1, T2	N2	MO	
	Т3	N0, N1, N2	MO	
IV	T4	Any N	MO	
	Any T	N3	MO	
	Any T	Any N	M1	

Bone C40, C41				
Stage	т	Ν	М	Grade (G)
IA	Τ1	NO, NX	МО	1,2
IB	Т2	NO, NX	МО	1,2
IIA	Τ1	NO, NX	МО	3, 4
IIB	Т2	NO, NX	MO	3, 4
ш	Т3	NO, NX	МО	Any G
IVA	Any T	NO, NX	Mla	Any G
IVB	Any T	N1	Any M	Any G
	Any T	Any N	M1b	Any G

Soft Tissues C381, C382, C383, C47-C49					
Stage	т	Ν	М	Grade (G)	
IA	Tla	NO, NX	МО	1,2	
	T1b	NO, NX	МО	1,2	
IB	T2a	NO, NX	МО	1,2	
	T2b	NO, NX	МО	1,2	
IIA	Tla	NO, NX	МО	3, 4	
	T1b	NO, NX	МО	3, 4	
IIB	T2a	NO, NX	МО	3, 4	
ш	T2b	NO, NX	МО	3, 4	
IV	Any T	N1	МО	Any G	
	Any T	Any N	M1	Any G	

Carcinoma of Skin (excluding eyelid, vulva, and penis) C440, C442-C449, C632			
Stage	т	Ν	М
0	Tis	NO	MO
I	T1	NO	MO
Ш	T2, T3	NO	MO
111	T4	NO	MO
	Any T	N1	MO
IV	Any T	Any N	M1

	Malignant Melanoma of Skir	n C44, C510, C609, C632	
Stage	т	Ν	М
0	pTis	NO	MO
1	pT1	NO	MO
IA	pTla	NO	MO
IB	pT1b	NO	MO
	pT2a	NO	MO
IIA	pT2b	NO	MO
	pT3a	NO	MO
IIB	pT3b	NO	MO
	pT4a	NO	MO
ΙΙС	pT4b	NO	MO
ш	Any pT	N1, N2, N3	MO
IIIA	pT1a-pT4a	N1a, N2a	MO
IIIB	pT1a-pT4a	N1b, N2b, N2c	MO
	pT1b-pT4b	N1a, N2a, N2c	MO
IIIC	pT1b-pT4b	N1b, N2b	MO
	Any pT	N3	MO
IV	Any pT	Any N	M1

Breast Tumours C50				
Stage	т	Ν	М	
0	Tis	NO	MO	
1	Τ1	NO	MO	
IIA	ТО	N1	MO	
	Τ1	N1	MO	
	Т2	NO	МО	
IIB	Т2	N1	MO	
	Т3	NO	МО	
IIIA	ТО	N2	MO	
	Τ1	N2	МО	
	Т2	N2	MO	
	Т3	N1, N2	МО	
IIIB	T4	N0, N1, N2	МО	
IIIC	Any T	N3	МО	
IV	Any T	Any N	M1	

Vulva C51				
Stage	т	Ν	М	
0	Tis	NO	МО	
1	Т1	NO	МО	
IA	Tla	NO	МО	
IB	T1b	NO	МО	
Ш	Т2	NO	МО	
Ш	T1,T2	N1	МО	
	Т3	N0, N1	MO	
IVA	T1, T2, T3	N2	МО	
	T4	Any N	МО	
IVB	Any T	Any N	M1	

Vagina C52				
Stage	т	Ν	М	
0	Tis	NO	MO	
1	Τ1	NO	MO	
Ш	Т2	NO	MO	
ш	Т3	NO	MO	
	T1, T2, T3	N1	MO	
IVA	T4	Any N	MO	
IVB	Any T	Any N	M1	

Cervix Uteri C53				
Stage	Т	Ν	М	
0	Tis	NO	МО	
IA	Tla	NO	MO	
IA1	Tlal	NO	МО	
IA2	T1a2	NO	MO	
IB	T1b	NO	МО	
IB1	T1b1	NO	МО	
IB2	T1b2	NO	МО	
IIA	T2a	NO	MO	
IIB	T2b	NO	МО	
IIIA	T3a	NO	MO	
IIIB	T1, T2, T3a	N1	MO	
	T3b	Any N	МО	
IVA	T4	Any N	МО	
IVB	Any T	Any N	M1	

Corpus Uteri C54				
Stage	т	Ν	М	
0	Tis	NO	МО	
IA	Tla	NO	MO	
IB	T1b	NO	МО	
IC	Tlc	NO	MO	
IIA	T2a	NO	МО	
IIB	T2b	NO	MO	
IIIA	T3a	NO	MO	
IIIB	T3b	NO	MO	
IIIC	T1, T2, T3	N1	MO	
IVA	T4	Any N	MO	
IVB	Any T	Any N	M1	

Ovary C56				
Stage	т	Ν	М	
IA	Tla	NO	МО	
IB	T1b	NO	МО	
IC	Tlc	NO	МО	
IIA	T2a	NO	МО	
IIB	T2b	NO	МО	
IIC	T2c	NO	МО	
IIIA	T3a	NO	МО	
IIIB	T3b	NO	MO	
IIIC	T3c	NO	МО	
	Any T	N1	МО	
IV	Any T	Any N	M1	

Fallopian Tube C570				
Stage	т	Ν	М	
0	Tis	NO	MO	
IA	Tla	NO	MO	
IB	T1b	NO	MO	
IC	Tlc	NO	MO	
IIA	T2a	NO	MO	
IIB	T2b	NO	MO	
IIC	T2c	NO	МО	
IIIA	T3a	NO	MO	
IIIB	T3b	NO	MO	
IIIC	T3c	NO	MO	
	Any T	N1	MO	
IV	Any T	Any N	M1	

Gestational Trophoblastic Tumours C58				
Stage	Т	N – not applicable	м	Risk category
1	Τ1		МО	unknown
IA	Τ1		МО	low
IB	Τ1		МО	high
Ш	Т2		МО	unknown
IIA	Т2		MO	low
IIB	Т2		MO	high
ш	Any T		Mla	unknown
IIIA	Any T		Mla	low
IIIB	Any T		Mla	high
IV	Any T		M1b	unknown
IVA	Any T		M1b	low
IVB	Any T		M1b	high

Penis C60				
Stage	Т	Ν	М	
0	Tis	NO	MO	
	Та	NO	МО	
1	Τ1	NO	MO	
Ш	Τ1	N1	МО	
	Τ2	NO	МО	
	Τ2	N1	МО	
ш	T1, T2	N2	МО	
	Т3	N0, N1, N2	MO	
IV	T4	Any N	МО	
	Any T	N3	МО	
	Any T	Any N	M1	

Prostate C61				
Stage	т	Ν	М	Grade (G)
1	Tla	NO	МО	1
II	Tla	NO	МО	2, 3, 4
	T1b, T1c	NO	МО	Any G
	T1, T2	NO	МО	Any G
ш	Т3	NO	МО	Any G
IV	T4	NO	МО	Any G
	Any T	N1	МО	Any G
	Any T	Any N	M1	Any G

Testis C62				
Stage	т	Ν	М	Serum tumour markers
0	pTis	NO	МО	S0, SX
1	pT1-T4	NO	МО	SX
IA	pT1	NO	МО	50
IB	pT2	NO	МО	50
	pT3	NO	МО	50
	pT4	NO	МО	50
IS	Any pT, Tx	NO	МО	S1-S3
Ш	Any pT, Tx	N1-N3	МО	SX
IIA	Any pT, Tx	N1	МО	50
	Any pT, Tx	N1	МО	51
IIB	Any pT, Tx	N2	МО	50
	Any pT, Tx	N2	МО	51
IIC	Any pT, Tx	N3	МО	50
	Any pT, Tx	N3	МО	51
ш	Any pT, Tx	Any N	M1, M1a	SX
IIIA	Any pT, Tx	Any N	M1, M1a	50
	Any pT, Tx	Any N	M1, M1a	51
IIIB	Any pT, Tx	N1-N3	МО	52
	Any pT, Tx	Any N	M1, M1a	52
	Any pT, Tx	N1-N3	МО	53
	Any pT, Tx	Any N	M1, M1a	S3
	Any pT, Tx	Any N	M1b	Any S
Kidney C64				
------------	------------	--------	----	--
Stage	т	Ν	М	
1	Т1	NO	МО	
Ш	Т2	NO	MO	
Ш	Т3	NO	MO	
	T1, T2, T3	N1	МО	
IV	T4	N0, N1	MO	
	Any T	N2	MO	
	Any T	Any N	Ml	

Renal Pelvis and Ureter C65, C66				
Stage	Т	Ν	М	
0a	Та	NO	МО	
Ois	Tis	NO	MO	
1	Т1	NO	МО	
Ш	Τ2	NO	МО	
Ш	Т3	NO	МО	
IV	Τ4	NO	MO	
	Any T	N1, N2, N3	MO	
	Any T	Any N	M1	

Urinary Bladder C67				
Stage	т	Ν	М	
0a	Та	NO	МО	
Ois	Tis	NO	MO	
1	Τ1	NO	MO	
Ш	T2a, b	NO	MO	
Ш	T3a, b	NO	МО	
	T4a	NO	MO	
IV	T4b	NO	МО	
	Any T	N1, N2, N3	MO	
	Any T	Any N	M1	

Appendix II: TNM 6 edition stage grouping and corresponding T, N, M values | 71

Urethra C680				
Stage	т	Ν	М	
0a	Та	NO	MO	
Ois	Tis	NO	MO	
	Tispu	NO	MO	
	Tispd	NO	MO	
1	Τ1	NO	MO	
Ш	Т2	NO	MO	
Ш	T1,T2	N1	MO	
	Т3	N0, N1	MO	
IV	T4	N0, N1	MO	
	Any T	N2	MO	
	Any T	Any N	M1	

Malignant Melanoma of Uvea C693, C694				
Stage	т	Ν	М	
1	Т1	NO	МО	
Ш	Т2	NO	MO	
ш	T3, T4	NO	MO	
IV	Any T	N1	MO	
	Any T	Any N	M1	

Appendix III: *TNM 7 edition stage grouping and corresponding T, N, M values*

Li	Lip and Oral Cavity C00, C02-C06 (except C051 and C052)			
Stage	Т	Ν	М	
0	Tis	NO	МО	
1	Т1	NO	МО	
Ш	Т2	NO	МО	
III	T1, T2, T3	N1	МО	
	Т3	NO	МО	
IVA	T1, T2, T3, T4a	N2	МО	
	T4a	N0, N1	МО	
IVB	Any T	N3	МО	
	T4b	Any N	МО	
IVC	Any T	Any N	M1	

Oropharynx	and Hypopharynx C01, C0	pharynx C01, C051, C052, C09, C100, C102, C103, C12, C1	
Stage	Т	Ν	М
0	Tis	NO	MO
1	Т1	NO	MO
Ш	Т2	NO	МО
Ш	T1, T2, T3	N1	MO
	Т3	NO	МО
IVA	T1, T2, T3	N2	МО
	T4a	N0, N1, N2	МО
IVB	T4b	Any N	МО
	Any T	N3	МО
IVC	Any T	Any N	M1

Nasopharynx C11			
Stage	Т	Ν	М
0	Tis	NO	MO
1	Т1	NO	MO
II	Т2	N0, N1	MO
	Т1	N1	MO
Ш	T1, T2	N2	MO
	Т3	N0, N1, N2	MO
IVA	T4	N0, N1, N2	MO
IVB	Any T	N3	MO
IVC	Any T	Any N	M1

Larynx C320, C321, C322, C101				
Stage	т	Ν	М	
0	Tis	NO	МО	
1	Т1	NO	MO	
Ш	Т2	NO	МО	
Ш	T1,T2	N1	MO	
	Т3	N0, N1	MO	
IVA	T1, T2, T3	N2	МО	
	T4a, T4b	N0, N1	MO	
IVB	T4b	Any N	MO	
	Any T	N3	МО	
IVC	Any T	Any N	M1	

Nasal Cavity and Paranasal Sinuses C300, C310, C311				
Stage	т	Ν	М	
0	Tis	NO	MO	
1	Т1	NO	MO	
Ш	Т2	NO	MO	
Ш	T1, T2, T3	N1	MO	
	Т3	NO	MO	
IVA	T1, T2, T3	N2	MO	
	T4a	N0, N1, N2	МО	
IVB	T4b	Any N	MO	
	Any T	N3	MO	
IVC	Any T	Any N	M1	

Malignant Melanoma of Upper Aerodigestive Tract C00-C06, C10-C14, C30-C32				
Stage	т	Ν	М	
ш	Т3	NO	MO	
IVA	T3, T4a	N1	MO	
	T4a	NO	MO	
IVB	T4b	Any N	МО	
IVC	Any T	Any N	M1	

Major Salivary Glands C07, C08				
Stage	Т	Ν	М	
1	Τ1	NO	MO	
Ш	Τ2	NO	MO	
ш	Т3	NO	МО	
	T1, T2, T3	N1	МО	
IVA	T4a	NO	МО	
	T4a	N1	MO	
	T1, T2, T3, T4a	N2	МО	
IVB	T4b	Any N	MO	
	Any T	N3	МО	
IVC	Any T	Any N	M1	

	Thyroid Gla	ind C73	
Stage	т	Ν	М
	Papillary or Follicu	lar, under 45 years	
1	Any T	Any N	MO
Ш	Any T	Any N	M1
	Papillary or Follicula	r, 45 years and older	
1	T1a, T1b	NO	МО
Ш	Т2	NO	MO
ш	Т3	NO	MO
	T1, T2, T3	Nla	MO
IVA	T1, T2, T3	N1b	MO
	T4a	N0, N1	MO
IVB	T4b	Any N	МО
IVC	Any T	Any N	M1
	Medı	ıllary	
1	T1a, T1b	NO	МО
Ш	Т2, Т3	NO	МО
Ш	T1, T2, T3	Nla	МО
IVA	T1, T2, T3	N1b	MO
	T4a	Any N	MO
IVB	T4b	Any N	MO
IVC	Any T	Any N	M1
	Anaplastic/Undifferentiat	ed (all cases are stage IV)	
IVA	T4a	Any N	МО
IVB	T4b	Any N	MO
IVC	Any T	Any N	M1

Oesophagus C15, C160				
Stage	т	Ν	М	
0	Tis	NO	MO	
IA	Τ1	NO	MO	
IB	Т2	NO	MO	
IIA	Т3	NO	MO	
IIB	T1,T2	N1	MO	
IIIA	T4a	NO	MO	
	Т3	N1	MO	
	T1,T2	N2	MO	
IIIB	Т3	N2	МО	
IIIC	T4a	N1, N2	MO	
	T4b	Any N	MO	
	Any T	N3	MO	
IV	Any T	Any N	M1	

	Stomach	C161-C164	
Stage	Т	N	м
0	Tis	NO	МО
IA	Т1	NO	МО
IB	Т2	NO	МО
	Т1	N1	МО
IIA	Т3	NO	МО
	Т2	N1	MO
	Т1	N2	МО
IIB	T4a	NO	МО
	Т3	N1	МО
	T2	N2	МО
	Т1	N3	МО
IIIA	T4a	N1	MO
	Т3	N2	MO
	Т2	N3	MO
IIIB	T4b	N0, N1	МО
	T4a	N2	МО
	Т3	N3	МО
	T4a	N3	МО
	T4b	N2, N3	МО
IV	Any T	Any N	M1

Small Intestine C17				
Stage	Т	Ν	М	
0	Tis	NO	MO	
1	T1,T2	NO	MO	
IIA	Т3	NO	MO	
IIB	T4	NO	MO	
IIIA	Any T	N1	MO	
IIIB	Any T	N2	MO	
IV	Any T	Any N	M1	

	Арр	endix-Carcinoma C	181	
Stage	т	Ν	м	Grade (G)
0	Tis	NO	MO	
1	T1, T2	NO	MO	
IIA	Т3	NO	MO	
IIB	T4a	NO	MO	
IIC	T4b	NO	MO	
IIIA	T1,T2	N1	MO	
IIIB	T3, T4	N1	MO	
IIIC	Any T	N2	MO	
IVA	Any T	NO	Mla	G1
IVB	Any T	NO	Mla	G2, G3
	Any T	N1, N2	Mla	Any G
IVC	Any T	Any N	M1b	Any G

Note: G1 well-differentiated/mucinous low grade; G2 moderately differentiated/mucinous high grade; G3 poorly differentiated/mucinous high grade; G4 undifferentiated.

Арреі	ndix-Carcinoid (Well-differentiated neuroendocrine tumour)			
Stage	т	Ν	М	
I	T1	NO	MO	
Ш	T2, T3	NO	MO	
Ш	Τ4	NO	MO	
	Any T	N1	MO	
IV	Any T	Any N	M1	

	Colon and Rectum	C18-C20, excluded C181	
Stage	т	Ν	М
0	Tis	NO	МО
1	T1,T2	NO	МО
Ш	T3, T4	NO	МО
IIA	Т3	NO	MO
IIB	T4a	NO	МО
IIC	T4b	NO	MO
Ш	Any T	N1, N2	МО
IIIA	T1, T2	N1	MO
	Т1	N2a	MO
IIIB	T3, T4a	N1	MO
	Т2, Т3	N2a	MO
	T1,T2	N2b	MO
IIIC	T4a	N2a	MO
	T3, T4a	N2b	MO
	T4b	N1, N2	MO
IVA	Any T	Any N	Mla
IVB	Any T	Any N	M1b

Anal Canal C211				
Stage	т	Ν	М	
0	Tis	NO	МО	
1	Т1	NO	MO	
Ш	Т2, Т3	NO	MO	
IIIA	T1, T2, T3	N1	MO	
	T4	NO	MO	
IIIB	T4	N1	MO	
	Any T	N2, N3	MO	
IV	Any T	Any N	M1	

Gastrointestinal Stromal Tumour (GIST)				
Stage	Т	Ν	М	Mitotic rate
		Gastric GIST*		
IA	T1,T2	NO	МО	Low
IB	Т3	NO	МО	Low
Ш	T1,T2	NO	МО	High
	T4	NO	МО	Low
IIIA	Т3	NO	МО	High
IIIB	T4	NO	МО	High
IV	Any T	N1	МО	Any rate
	Any T	Any N	M1	Any rate
		Small Intestinal GIST*		
1	T1,T2	NO	МО	Low
Ш	Т3	NO	МО	Low
IIIA	Т1	NO	МО	High
	Τ4	NO	МО	Low
IIIB	T2, T3, T4	NO	МО	High
IV	Any T	N1	МО	Any rate
	Any T	Any N	M1	Any rate

* Staging criteria for gastric tumours can be applied in primary, solitary omental GISTs. Staging criteria for intestinal tumours can be applied to GISTs in less common sites, such as oesophagus, colon, rectum and mesentery.

Gastric, Small, and Large Intestinal Carcinoid Tumours

(Well-differentiated Neuroendocrine Tumours and Well-differentiated Neuroendocrine Carcinomas)

Stage	т	Ν	м
1	Т1	NO	МО
IIA	Т2	NO	MO
IIB	Т3	NO	MO
IIIA	Τ4	NO	MO
IIIB	Any T	N1	МО
IV	Any T	Any N	M1

Liver-Hepatocellular Carcinoma C220				
Stage	т	Ν	М	
1	Τ1	NO	МО	
Ш	Т2	NO	МО	
IIIA	T3a	NO	МО	
IIIB	T3b	NO	MO	
IIIC	T4	N1	МО	
IVA	Any T	N1	MO	
IVB	Any T	Any N	M1	

Liver-Intrahepatic Bile Duct C221				
Stage	т	Ν	М	
1	Τ1	NO	MO	
Ш	Τ2	NO	МО	
Ш	Т3	NO	МО	
IVA	T4	NO	МО	
	Any T	N1	МО	
IVB	Any T	Any N	M1	

Gallbladder C23				
Stage	Т	Ν	М	
0	Tis	NO	MO	
1	Т1	NO	MO	
Ш	Т2	NO	MO	
IIIA	Т3	NO	MO	
IIIB	T1, T2, T3	N1	MO	
IVA	T4	Any N	MO	
IVB	Any T	Any N	M1	

Extrahepatic Bile Ducts-Perihiliar C240				
Stage	т	Ν	М	
0	Tis	NO	MO	
1	Τ1	NO	MO	
Ш	T2a, T2b	NO	МО	
IIIA	Т3	NO	MO	
IIIB	T1, T2, T3	N1	МО	
IVA	T4	N0, N1	MO	
IVB	Any T	Any N	M1	

Extrahepatic Bile Ducts-Distal C240				
Stage	т	Ν	М	
0	Tis	NO	MO	
IA	Τ1	NO	МО	
IB	Т2	NO	МО	
IIA	Т3	NO	MO	
IIB	T1, T2, T3	N1	МО	
ш	T4	Any N	MO	
IV	Any T	Any N	M1	

Ampulla of Vater C241				
Stage	т	Ν	М	
0	Tis	NO	MO	
IA	Τ1	NO	MO	
IB	Τ2	NO	МО	
IIA	Т3	NO	MO	
IIB	T1, T2, T3	N1	МО	
ш	T4	Any N	MO	
IV	Any T	Any N	M1	

Pancreas C25				
Stage	т	Ν	М	
0	Tis	NO	МО	
IA	Τ1	NO	MO	
IB	Т2	NO	MO	
IIA	Т3	NO	MO	
IIB	T1, T2, T3	N1	MO	
Ш	T4	Any N	MO	
IV	Any T	Any N	M1	

Lung C34				
Stage	Т	Ν	М	
Occult carcinoma	ТХ	NO	МО	
0	Tis	NO	MO	
IA	T1a, T1b	NO	МО	
IB	T2a	NO	МО	
IIA	T2b	NO	МО	
	T1a, T1b	N1	МО	
	T2a	N1	МО	
IIB	T2b	N1	МО	
	Т3	NO	MO	
IIIA	T1a, T1b, T2a, T2b	N2	МО	
	Т3	N1, N2	МО	
	T4	N0, N1	МО	
IIIB	Any T	N3	МО	
	Τ4	N2	МО	
IV	Any T	Any N	M1	

Pleural Mesothelioma C384					
Stage	Т	Ν	М		
IA	Tla	NO	MO		
IB	T1b	NO	MO		
Ш	Т2	NO	MO		
III	T1, T2	N1	MO		
	T1,T2	N2	MO		
	Т3	N0, N1, N2	MO		
IV	T4	Any N	MO		
	Any T	N3	MO		
	Any T	Any N	M1		

Bone C40, C41				
Stage	т	Ν	М	Grade (G)
IA	Τ1	NO	МО	1,2
IB	Т2	NO	МО	1,2
IIA	Τ1	NO	МО	3, 4
IIB	Т2	NO	МО	3, 4
ш	Т3	NO	МО	Any G
IVA	Any T	NO	Mla	Any G
IVB	Any T	N1	Any M	Any G
	Any T	Any N	M1b	Any G

Note: Use N0 for NX. For T1 and T2, use low grade if no grade is stated.

Soft Tissues C381-C383, C47, C480, C49					
Stage	т	Ν	М	Grade (G)	
IA	Tla	NO	МО	1,2	
	T1b	NO	МО	1,2	
IB	T2a	NO	МО	1,2	
	T2b	NO	МО	1,2	
IIA	Tla	NO	МО	3, 4	
	T1b	NO	МО	3, 4	
IIB	T2a	NO	МО	3, 4	
Ш	T2b	NO	МО	3, 4	
	Any T	N1	МО	Any G	
IV	Any T	Any N	M1	Any G	

Note: Use low grade for GX. Use N0 for NX.

Carcinoma of Skin of Eyelid C441				
Stage	Т	Ν	М	
0	Tis	NO	MO	
IA	Т1	NO	MO	
IB	T2a	NO	MO	
IC	T2b	NO	MO	
Ш	T3a	NO	MO	
IIIA	T3b	NO	MO	
IIIB	Any T	N1	MO	
IIIC	T4	Any N	MO	
IV	Any T	Any N	M1	

Carcinoma c	of Skin (excluding eyelid, vulva	a, and penis) C440, C442	-C449, C632
Stage	т	Ν	М
0	Tis	NO	МО
1	Τ1	NO	MO
Ш	Т2	NO	МО
Ш	Т3	NO	MO
	T1, T2, T3	N1	MO
IV	T1, T2, T3	N2, N3	МО
	T4	Any N	МО
	Any T	Any N	M1

Malignant Melanoma of Skin C44, C510, C609, C632				
Stage	Т	Ν	М	
0	pTis	NO	МО	
1	pT1	NO	MO	
IA	pT1a	NO	MO	
IB	pT1b	NO	MO	
	pT2a	NO	MO	
IIA	pT2b	NO	MO	
	pT3a	NO	МО	
IIB	pT3b	NO	MO	
	pT4a	NO	MO	
IIC	pT4b	NO	МО	
ш	Any pT	N1, N2, N3	MO	
IIIA	pT1a-pT4a	N1a, N2a	MO	
IIIB	pT1a-pT4a	N1b, N2b, N2c	MO	
	pT1b-pT4b	N1a, N2a, N2c	MO	
IIIC	pT1b-pT4b	N1b,N2b	MO	
	Any pT	N3	МО	
IV	Any pT	Any N	M1	

Malignant Melanoma of Uvea C693, C694					
Stage	Т	Ν	М		
1	Tla	NO	MO		
IIA	T1b-T1d, T2a	NO	MO		
IIB	T2b, T3a	NO	MO		
IIIA	T2c-T2d	NO	MO		
	T3b-T3c	NO	МО		
	T4a	NO	MO		
IIIB	T3d	NO	МО		
	T4b-T4c	NO	MO		
IIIC	T4d-T4e	NO	MO		
IV	Any T	N1	МО		
	Any T	Any N	M1		

Merkel Cell Carcinoma of Skin C440-C449, C632					
Stage	т	Ν	М		
0	Tis	NO	MO		
1	Т1	NO	MO		
IA	Т1	pNO	MO		
IB	Т1	cN0	MO		
IIA	Т2, Т3	pNO	MO		
IIB	Т2, Т3	cN0	MO		
IIC	T4	NO	MO		
IIIA	Any T	Nla	MO		
IIIB	Any T	N1b, N2	MO		
IV	Any T	Any N	M1		

Breast Tumours C50				
Stage	Т	Ν	М	
0	Tis	NO	МО	
IA	T1*	NO	MO	
IB	T0,T1*	N1mi	МО	
IIA	T0,T1*	N1	MO	
	Т2	NO	MO	
IIB	Т2	N1	MO	
	Т3	NO	MO	
IIIA	T0, T1*, T2	N2	MO	
	Т3	N1, N2	MO	
IIIB	T4	N0, N1, N2	MO	
IIIC	Any T	N3	MO	
IV	Any T	Any N	M1	

* T1 includes T1mi.

Vulva C51				
Stage	Т	Ν	М	
0	Tis	NO	MO	
I	Т1	NO	MO	
IA	Tla	NO	MO	
IB	T1b	NO	MO	
Ш	Т2	NO	MO	
IIIA	T1,T2	N1a, N1b	MO	
IIIB	T1,T2	N2a, N2b	МО	
IIIC	T1,T2	N2c	MO	
IVA	T1,T2	N3	МО	
	Т3	Any N	МО	
IVB	Any T	Any N	M1	

Vagina C52				
Stage	т	Ν	М	
0	Tis	NO	MO	
1	Τ1	NO	MO	
Ш	Τ2	NO	МО	
ш	Т3	NO	МО	
	T1, T2, T3	N1	MO	
IVA	Τ4	Any N	МО	
IVB	Any T	Any N	M1	

Cervix Uteri C53				
Stage	Т	Ν	М	
0	Tis	NO	МО	
1	Т1	NO	МО	
IA	Tla	NO	МО	
IA1	Tlal	NO	МО	
IA2	T1a2	NO	МО	
IB	T1b	NO	МО	
IB1	T1b1	NO	МО	
IB2	T1b2	NO	МО	
Ш	Т2	NO	МО	
IIA	T2a	NO	МО	
IIA1	T2a1	NO	МО	
IIA2	T2a2	NO	МО	
IIB	Т2Ь	NO	МО	
Ш	Т3	NO	МО	
IIIA	T3a	NO	МО	
IIIB 	T1, T2, T3	N1	МО	
	ТЗЬ	Any N	МО	
IVA	T4	Any N	МО	
IVB	Any T	Any N	M1	

Corpus Uteri C541, C55				
Stage	т	Ν	М	
IA	Tla	NO	MO	
IB	T1b	NO	MO	
Ш	Т2	NO	MO	
IIIA	T3a	NO	MO	
IIIB	T3b	NO	МО	
IIIC	T1, T2, T3	Nl	МО	
IVA	T4	Any N	МО	
IVB	Any T	Any N	M1	

Uterine Sarcoma: Leiomyosarcoma (8890/3), endometrial stromal sarcoma (8930/3) and adenosarcoma (8933/3) C53, C540, C543				
Stage	т	Ν	М	
1	Т1	NO	MO	
IA	Tla	NO	MO	
IB	T1b	NO	МО	
IC*	Tlc	NO	MO	
Ш	Т2	NO	MO	
IIA	T2a	NO	MO	
IIB	T2b	NO	MO	
IIIA	T3a	NO	МО	
IIIB	T3b	NO	MO	
IIIC	T1, T2, T3	N1	МО	
IVA	T4	Any N	MO	
IVB	Any T	Any N	M1	

* Stage IC does not apply for leiomyosarcoma and endometrial stromal sarcoma.

Ovary C56				
Stage	Т	Ν	М	
IA	Tla	NO	MO	
IB	T1b	NO	MO	
IC	Tlc	NO	MO	
IIA	T2a	NO	MO	
IIB	T2b	NO	MO	
ΙΙС	T2c	NO	MO	
IIIA	T3a	NO	MO	
IIIB	T3b	NO	MO	
IIIC	T3c	NO	MO	
	Any T	N1	MO	
IV	Any T	Any N	M1	

Fallopian Tube C570				
Stage	т	Ν	М	
0	Tis	NO	МО	
IA	Tla	NO	MO	
IB	T1b	NO	MO	
IC	T1c	NO	MO	
IIA	T2a	NO	MO	
IIB	T2b	NO	MO	
IIC	T2c	NO	MO	
IIIA	T3a	NO	MO	
IIIB	T3b	NO	MO	
IIIC	T3c	NO	MO	
	Any T	N1	MO	
IV	Any T	Any N	M1	

Gestational Trophoblastic Tumours C58				
Stage	т	N – not applicable	М	Risk category
1	Τ1		МО	unknown
IA	Τ1		МО	low
IB	Τ1		МО	high
Ш	Τ2		МО	unknown
IIA	Т2		МО	low
IIB	Т2		МО	high
ш	Any T		Mla	unknown
IIIA	Any T		Mla	low
IIIB	Any T		Mla	high
IV	Any T		M1b	unknown
IVA	Any T		M1b	low
IVB	Any T		M1b	high

Penis C60				
Stage	т	Ν	М	
0	Tis	NO	МО	
	Та	NO	МО	
1	Tla	NO	МО	
II	T1b	NO	МО	
	Τ2	N0, N1	МО	
	Т3	NO	МО	
IIIA	T1, T2, T3	N1	МО	
IIIB	T1, T2, T3	N2	МО	
IV	T4	Any N	МО	
	Any T	N3	МО	
	Any T	Any N	M1	

Prostate C61				
Stage	т	Ν	М	
1	T1,T2a	NO	MO	
Ш	T2b, T2c	NO	МО	
ш	Т3	NO	МО	
IV	T4	NO	МО	
	Any T	N1	МО	
	Any T	Any N	M1	

Testis C62				
Stage	т	Ν	М	Serum tumour markers
0	pTis	NO	МО	50, SX
1	pT1-T4	NO	МО	SX
ΙΑ	pT1	NO	МО	50
IB	рТ2-Т4	NO	МО	50
IS	Any pT/Tx	NO	МО	S1-S3
Ш	Any pT/Tx	N1-N3	МО	SX
IIA	Any pT/Tx	N1	МО	50
	Any pT/Tx	N1	МО	51
IIB	Any pT/Tx	N2	МО	50
	Any pT/Tx	N2	МО	S1
IIC	Any pT/Tx	N3	МО	50
	Any pT/Tx	N3	МО	S1
Ш	Any pT/Tx	Any N	Mla	SX
IIIA	Any pT/Tx	Any N	Mla	50
	Any pT/Tx	Any N	Mla	S1
IIIB	Any pT/Tx	N1-N3	МО	52
	Any pT/Tx	Any N	Mla	52
	Any pT/Tx	N1-N3	МО	53
	Any pT/Tx	Any N	Mla	53
	Any pT/Tx	Any N	M1b	Any S

Kidney C64				
Stage	т	Ν	М	
1	Τ1	NO	MO	
Ш	Т2	NO	MO	
III	Т3	Any N	МО	
	T1, T2, T3	N1	МО	
IV	T4	Any N	MO	
	Any T	N2	MO	
	Any T	Any N	M1	

Renal Pelvis and Ureter C65, C66				
Stage	т	Ν	М	
0a	Та	NO	МО	
Ois	Tis	NO	MO	
1	Τ1	NO	МО	
Ш	Τ2	NO	МО	
ш	Т3	NO	МО	
IV	T4	NO	MO	
	Any T	N1, N2, N3	MO	
	Any T	Any N	M1	

Urinary Bladder C67				
Stage	Т	Ν	М	
0a	Та	NO	MO	
Ois	Tis	NO	MO	
1	Т1	NO	MO	
Ш	T2a, T2b	NO	MO	
Ш	T3a, T3b	NO	MO	
	T4a	NO	MO	
IV	T4b	NO	MO	
	Any T	N1, N2, N3	MO	
	Any T	Any N	M1	

	Urethra C680 and C619 (transitional cell carcinomas)		
Stage	т	Ν	М
0a	Та	NO	MO
Ois	Tis	NO	MO
	Tispu	NO	MO
	Tispd	NO	MO
1	Т1	NO	MO
Ш	Т2	NO	MO
Ш	T1,T2	N1	MO
	Т3	N0, N1	MO
IV	T4	N0, N1	MO
	Any T	N2	MO
	Any T	Any N	M1

Adrenal Cortex Tumours C740				
Stage	т	Ν	М	
1	Τ1	NO	MO	
Ш	Τ2	NO	MO	
III	T1, T2	N1	MO	
	Т3	NO	МО	
IV	Т3	N1	MO	
	T4	Any N	MO	
	Any T	Any N	M1	

GETTING IN TOUCH WITH THE EU

In person

All over the European Union there are hundreds of Europe Direct information centres. You can find **the address of the centre nearest you at:** <u>http://europea.eu/contact</u>

On the phone or by email

Europe Direct is a service that answers your questions about the European Union. You can contact this service: - by freephone: 00 800 6 7 8 9 10 11 (certain operators may charge for these calls),

- at the following standard number: +32 22999696, or

- by electronic mail via: <u>http://europa.eu/contact</u>

FINDING INFORMATION ABOUT THE EU

Online

Information about the European Union in all the official languages of the EU is available on the Europa website at: <u>http://europa.eu</u>

EU publications

You can download or order free and priced EU publications from EU Bookshop at: <u>http://bookshop.europa.eu</u>. Multiple copies of free publications may be obtained by contacting Europe Direct or your local information centre (see <u>http://europa.eu/contact</u>).

JRC Mission

As the Commission's in-house science service, the Joint Research Centre's mission is to provide EU policies with independent, evidence-based scientific and technical support throughout the whole policy cycle.

Working in close cooperation with policy Directorates-General, the JRC addresses key societal challenges while stimulating innovation through developing new methods, tools and standards, and sharing its know-how with the Member States, the scientific community and international partners.

Serving society Stimulating innovation Supporting legislation



doi:10.2760/429053 ISBN 978-92-79-77889-6