Belgian Cancer Registry

Stereotactic Radiotherapy (SRT)
-
Registration form

Please register all SRT treatments started in a certain year by June 30th of the following year at the latest.

Please note that by the terms ‘current’ or ‘currently’, it is meant: ‘at the time of start of the SRT treatment that is being registered’.

Version 1.0 (04/02/2020)

REQ: Required variable
❍: Single-select variable: only one answer can be selected
❑: Multi-select variable: one or more answers can be selected

Administrative patient data

Hospital REQ: .....
Health insurance institution REQ: .....
National number for social security (INSZ/NISS)* REQ: .....

** if filled out in the WBCR application, the following variables will be completed automatically:*

Last name REQ: .....
First name REQ: .....
Postal code REQ: .....
City REQ: .....
Country REQ: .....
Health insurance number: .....
Date of birth REQ: ...../...../..... (dd/mm/yyyy)
Date of death: ...../...../..... (dd/mm/yyyy)
Sex REQ: ❍ Male
❍ Female

1. Diagnostics

Lesion to treat REQ: ❍ Primary tumour or relapse of primary tumour (Complete sections: 1A, 2A, 3A)
❍ Metastasis (Complete sections: 1B, 2B, 3B)
❍ Cranial arteriovenous malformation (AVM) (Complete sections: 1C, 2, 3)

A. Primary tumour or relapse of primary tumour

Indication REQ: ❍ Primary tumour
- Incidence date primary tumour REQ: ..... / ..... / ..... (dd/mm/yyyy)
- Clinical stage primary tumour (cTNM): cT: ..... cN: ..... cM: .....
- Pathological stage primary tumour (pTNM): pT: ..... pN: ..... pM: .....

❍ Relapse of primary tumour
- Incidence date initial primary tumour REQ: ..... / ..... / ..... (dd/mm/yyyy)
- Date of diagnosis of current relapse REQ: ..... / ..... / ..... (dd/mm/yyyy)
- Clinical stage of the relapse (rcTNM): rcT: ..... rcN: ..... rcM: .....

Basis for diagnosis primary tumour/relapse REQ: ❍ 1 - Autopsy
❍ 2 - Histology of primary tumour
❍ 3 - Histology metastasis
❍ 4 - Cytology/haematology
❍ 5 - Technical (e.g. CT scan, endoscopy, …)
❍ 6 - Clinical
❍ 7 - Tumour marker (e.g. PSA, HCG, AFP, Ig, …)
❍ Unknown

WHO score at start SRT treatment primary tumour/relapse [REQ]:

- ❍ 0 - Asymptomatic, normal activity
- ❍ 1 - Symptomatic, but ambulant
- ❍ 2 - Symptomatic, bedbound < 50% day
- ❍ 3 - Symptomatic, bedbound > 50% day
- ❍ 4 - Completely dependent, 100% bedbound
- ❍ Unknown

Primary tumour/relapse localisation [REQ]: .....

Laterality primary tumour/relapse [REQ]:
- ❍ Left
- ❍ Right
- ❍ Unpair organ
- ❍ Unknown

Histological diagnosis primary tumour/relapse [REQ]: .....

B. Metastasis

Incidence date primary tumour [REQ]: ..... / ..... / ..... (dd/mm/yyyy)

Localisation primary tumour [REQ]:
- ❍ Bladder
- ❍ Breast
- ❍ Cervix
- ❍ Colon
- ❍ Head and neck
- ❍ Kidney
- ❍ Lung
- ❍ Melanoma
- ❍ Oesophagus
- ❍ Pancreas
- ❍ Prostate
- ❍ Rectum
- ❍ Soft tissue
- ❍ Uterus
- ❍ Unknown
- ❍ Other; Specify [REQ]: .....

Current status primary tumour [REQ]:
- ❍ Controlled
- ❍ Not controlled

Prior metastatic event [REQ]?

❍ Unknown
❍ No
❍ Yes, oligometastatic disease (≤ 5 metastases)
❍ Yes, polymetastatic disease (> 5 metastases)

- Was prior metastases-directed treatment performed [REQ] ?

❍ Unknown
❍ No
❍ Yes *
❍ Yes, but information only partially available *

* - How many types of treatment did the patient receive for which information is available [REQ] ? °

❍ 1	❍ 6
❍ 2	❍ 7
❍ 3	❍ 8
❍ 4	❍ 9
❍ 5	❍ 10

° *Specify the following variables for each of the received treatments (pop-up):*

- Type of treatment [REQ] : ❍ Chemotherapy
❍ Hormonal therapy
❍ Immunotherapy
❍ Targeted therapy
❍ Surgery
❍ Radical radiotherapy
❍ Radiofrequency ablation (RFA)
❍ Other; Specify [REQ] : .....

- Start date [REQ] : ..... / ..... / ..... (dd/mm/yyyy)

- End date [REQ] : ..... / ..... / ..... (dd/mm/yyyy)

(When an exact date is not known, please enter 15/mm/yyyy if only month and year are known, or 1/07/yyyy if only the year is known.)

Date of diagnosis of current metastasis, treated within the currently administered dosimetric plan [REQ]:
..... / ..... / ..... (dd/mm/yyyy)

Number of currently active oligometastatic lesion(s) [REQ] : .....

Localisation of currently active oligometastatic lesion(s) [REQ] :

- ❑ Adrenal metastases
- ❑ Bone (non-spinal) metastases
- ❑ Brain metastases
- ❑ Hepatic metastases
- ❑ Lung metastases
- ❑ Lymph node metastases
- ❑ (Para-) spinal metastases
- ❑ Other (oligo)metastatic lesion(s)

  Specify [REQ]: ……

WHO score at start SRT treatment of the current metastasis [REQ]:

- ❍ 0 - Asymptomatic, normal activity
- ❍ 1 - Symptomatic, but ambulant
- ❍ 2 - Symptomatic, bedbound < 50% day
- ❍ 3 - Symptomatic, bedbound > 50% day
- ❍ 4 - Completely dependent, 100% bedbound
- ❍ Unknown

C. Cranial AVM

Date of diagnosis of the AVM treated within the currently administered dosimetric plan [REQ]:

…… / …… / …… (dd/mm/yyyy)

WHO score at start SRT treatment of the AVM [REQ]:

- ❍ 0 - Asymptomatic, normal activity
- ❍ 1 - Symptomatic, but ambulant
- ❍ 2 - Symptomatic, bedbound < 50% day
- ❍ 3 - Symptomatic, bedbound > 50% day
- ❍ 4 - Completely dependent, 100% bedbound
- ❍ Unknown

2. Lesion specifications

Number of lesions in total to treat with SRT and/or SRS (cerebral lesions included) [REQ]: ……

Number of lesions treated within the currently administered dosimetric plan [REQ]: ……

Maximum diameter of the lesion(s) treated within the currently administered dosimetric plan [REQ]:

…… mm *(this is the sum of the (largest) diameters of all the lesions treated within the current plan)*

A. Primary tumour or relapse of primary tumour

Localisation of (the relapse of) the primary tumour lesion treated within the currently administered dosimetric plan [REQ]:

- ❍ Primary brain lesion
- ❍ Primary head & neck lesion
- ❍ Primary hepatic lesion
- ❍ Primary lung (peripheral) lesion
- ❍ Primary lung (central and/or >5 cm) lesion
- ❍ Primary pancreatic lesion
- ❍ Primary (para-) spinal lesion
- ❍ Primary prostate lesion
- ❍ Primary renal lesion
- ❍ Other primary lesion

  Specify [REQ]: ……

B. Metastasis

Localisation of the metastatic lesion(s) treated within the currently administered dosimetric plan [REQ]:

- ❍ Adrenal metastases
- ❍ Bone (non-spinal) metastases
- ❍ Brain metastases
- ❍ Hepatic metastases
- ❍ Lung metastases
- ❍ Lymph node metastases
- ❍ (Para-) spinal metastases
- ❍ Other (oligo)metastatic lesion

  Specify [REQ]: ……

3. Treatment specifications

Total dose delivered for the currently administered dosimetric plan [REQ] : …… Gy

Number of fractions delivered [REQ] : ……

Start date of RT for the currently administered dosimetric plan [REQ] : …… / …… / …… (dd/mm/yyyy)

End date of RT for the currently administered dosimetric plan [REQ] : …… / …… / …… (dd/mm/yyyy)

Centre where the RT was performed [REQ] : ……..

Centre that referred the patient to RT [REQ] : ……..

A. Primary tumour or relapse of primary tumour

Other treatment for the currently active locoregional primary lesion, administered within 90 days before or after SRT [REQ]:

❍ No

❍ Yes; Specify [REQ]:
- ❑ Chemotherapy
- ❑ Hormonal therapy
- ❑ Immunotherapy
- ❑ Targeted therapy
- ❑ Surgery
- ❑ Radical radiotherapy
- ❑ Radiofrequency ablation (RFA)
- ❑ Other; Specify [REQ]: ……

B. Metastasis

Current status primary tumour [REQ]:

❍ Controlled (complete 3B1)

❍ Not controlled (complete 3B2)

B1. Metastasis with controlled primary tumour

Other treatment for all currently active metastatic lesions, administered within 90 days before or after SRT [REQ]:

❍ No

❍ Yes; Specify [REQ]:
- ❑ Chemotherapy
- ❑ Hormonal therapy
- ❑ Immunotherapy
- ❑ Targeted therapy
- ❑ Surgery
- ❑ Radical radiotherapy
- ❑ Radiofrequency ablation (RFA)
- ❑ Other; Specify [REQ]: ……

B2. Metastasis with uncontrolled primary tumour

Other treatment for the currently active locoregional primary lesion and/or all currently active oligometastatic lesions, administered within 90 days before or after SRT [REQ]:

❍ No

❍ Yes; Specify [REQ]:
- ❑ Chemotherapy
- ❑ Hormonal therapy
- ❑ Immunotherapy
- ❑ Targeted therapy
- ❑ Surgery
- ❑ Radical radiotherapy
- ❑ Radiofrequency ablation (RFA)
- ❑ Other; Specify [REQ]: ……