

# A Novel Standard for Systematic Reporting of Neuroblastoma Surgery: The International Neuroblastoma Surgical Report Form (INSRF)

*A Joint Initiative by the Pediatric Oncological Cooperative Groups SIOPEX\*, COG\*\*, and GPOH\*\*\**

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 on behalf of the Surgical and Medical Committees of SIOPEX\*, COG\*\* and GPOH\*\*\*

**Objective:** To create the first structured surgical report form for NBL with international consensus, to permit standardized documentation of all NBL-related surgical procedures and their outcomes.

**Summary of Background Data:** NBL, the most common extracranial solid malignant tumor in children, covers a wide spectrum of tumors with significant differences in anatomical localization, organ or vessel involvement, and tumor biology. Complete surgical resection of the primary tumor is an important part of NBL treatment, but maybe hazardous, prone to complications and its role in high-risk disease remains debated. Various surgical guidelines exist within the protocols of the different cooperative groups, although there is no standardized operative report form to document the surgical treatment of NBL.

**Methods:** After analyzing the treatment protocols of the SIOP Europe International Neuroblastoma Study Group, Children's Oncology Group,

and Gesellschaft fuer Paediatrische Onkologie und Haematologie – German Association of Pediatric Oncology and Haematology pediatric cooperative groups, important variables were defined to completely describe surgical biopsy and resection of NBL and their outcomes. All variables were discussed within the Surgical Committees of SIOP Europe International Neuroblastoma Study Group, Children's Oncology Group, and Gesellschaft fuer Paediatrische Onkologie und Haematologie – German Association of Pediatric Oncology and Haematology. Thereafter, joint meetings were organized to obtain intercontinental consensus.

**Results:** The “International Neuroblastoma Surgical Report Form” provides a structured reporting tool for all NBL surgery, in every anatomical region, documenting all Image Defined Risk Factors and structures involved, with obligatory reporting of intraoperative and 30 day-postoperative complications.

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\*SIOPEX: SIOP Europe International Neuroblastoma Study Group.

\*\*COG: Children's Oncology Group.

\*\*\*GPOH: Gesellschaft fuer Paediatrische Onkologie und Haematologie – German Association of Pediatric Oncology and Haematology.

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**Conclusion:** The International Neuroblastoma Surgical Report Form is the first universal form for the structured and uniform reporting of NBL-related surgical procedures and their outcomes, aiming to facilitate the postoperative communication, treatment planning and analysis of surgical treatment of NBL.

**Keywords:** biopsy, Clavien-Dindo classification, complication, high-risk, neuroblastoma, operation, outcome, postoperative, quality, reporting, resection, standardization, surgery, tumor  
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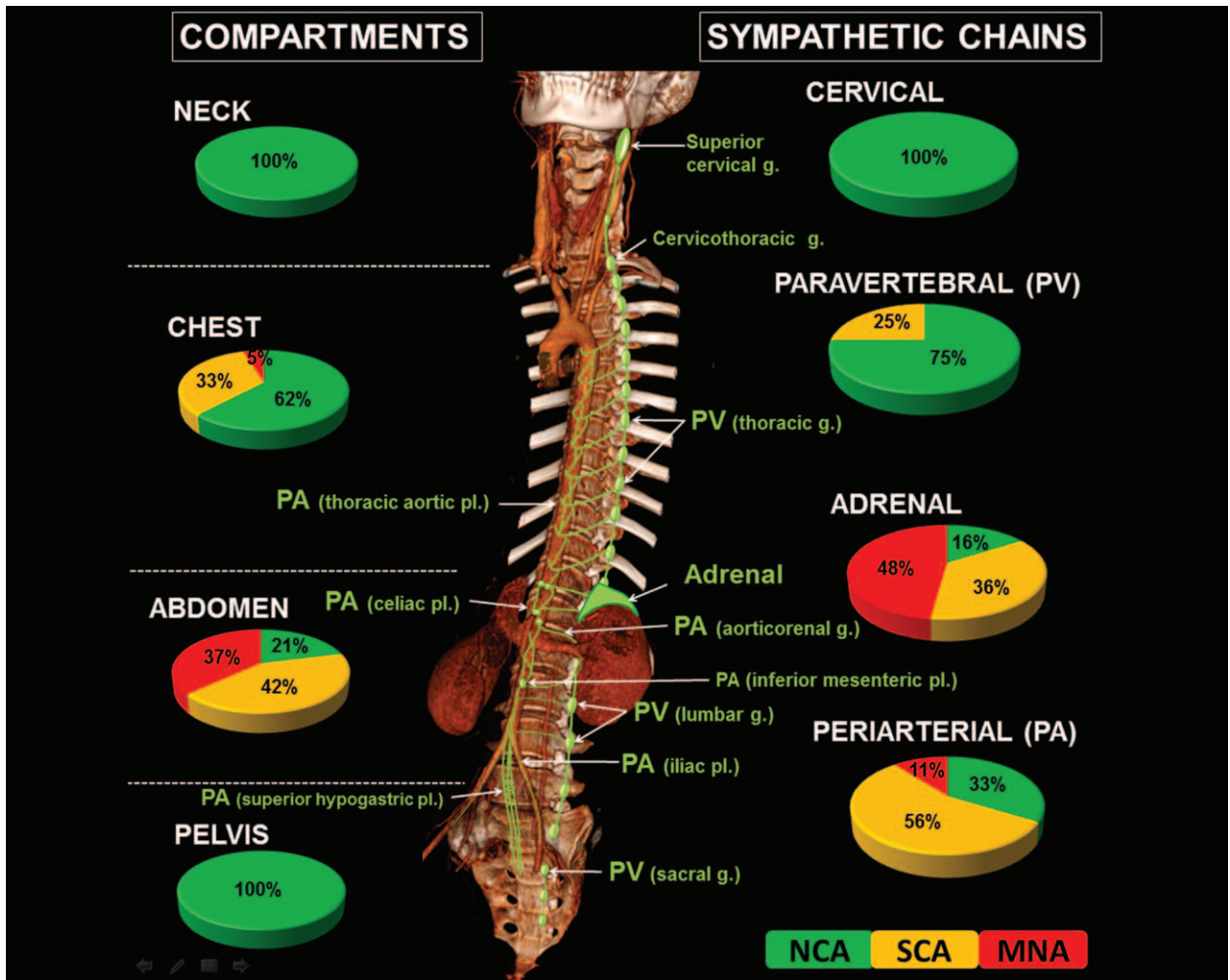
Neuroblastoma (NBL), an embryonal sympathetic nervous system tumor, is the most common cancer in infants and the most common extracranial solid malignant tumor in children,<sup>1–4</sup> with an overall age-standardized incidence rate of 10.2–10.9 cases per million children in the USA and Europe, respectively.<sup>2,3</sup> NBL accounts for 6% of all childhood cancers<sup>1</sup> and 12%–15% of cancer-related deaths under the age of 15 years.<sup>3,5</sup>

NBL has a broad spectrum of clinical behavior that correlates with a number of clinical and biological features.<sup>6</sup> After international

consensus, NBL is treated multimodally, according to the pretreatment International NBL Risk Group (INRG) classification system.<sup>6</sup>

This system identifies 4 risk categories (very low-, low-, intermediate-, and high-risk) taking into account the tumor stage, age at diagnosis, histology, grade, Neuroblastoma Myc (MYCN) oncogene status, chromosomal aberrations, and tumor cell ploidy.<sup>6</sup> To uniformly stage NBL before the initiation of treatment, the INRG-Staging-System was developed, based upon tumor imaging and on the absence or presence of preoperative “Image-Defined Risk Factors” (IDRFs).<sup>7</sup> These IDRFs describe the various locations where NBL may occur and the vital structures (organs, vascular, and neural structures) that may be involved by the tumor.<sup>8</sup>

As NBL originates from developmental anomalies of the neural crest, there is a wide spectrum of locations where the primary tumor may occur, related to the sympathetic nervous system anatomy.<sup>3</sup> The anatomical distribution of NBL can be described by compartmental classification (Fig. 1)<sup>4</sup> and most primary tumors arise in the abdominal compartment (adrenal gland 48%, extra-adrenal retroperitoneum 25%). The thoracic location of NBL is less frequent accounting for 16%–20%, and the cervical and pelvic compartments account for 3% to 5% of NBL, respectively.<sup>4</sup> Recent



**FIGURE 1.** Radiogenomics classification of neuroblastoma by anatomical localization of the primary tumor: compartmental versus sympathetic chain classification with respective genomic profile distribution.<sup>4</sup> (with permission of the authors).

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studies suggest that the sympathetic anatomical location of NBL may also be relevant as a prognostic factor.<sup>4</sup>

Local control of the primary tumor by complete surgical resection is an important part of NBL treatment. The role of surgical resection in high-risk NBL however remains debated as it is difficult to study, due to the relative rarity of NBL, its different localizations, heterogenous presentations<sup>4</sup> and internationally varying treatment protocols, incorporating postoperative radiation therapy to aid in local control. Especially for high-risk NBL, divergent results have been reported on the role, extent, and timing of surgical resection.<sup>9–11</sup> In addition, the reporting of surgical treatment of NBL is not standardized and may vary considerably.

As the surgical reporting of a postoperative residual primary tumor may influence postoperative irradiation planning in future high-risk NBL treatment protocols, the documentation of NBL surgery and its immediate outcomes must be optimized.

Improving the quality of surgical reporting will also facilitate the analysis of the role, extent, and timing of surgical treatment in NBL. Therefore we aimed to develop a uniform, structured report form to document every NBL-related surgical procedure.

## METHODS

A working party group within the Surgical Committee of SIOP Europe International Neuroblastoma Study Group (SIOPEN), under the joint leadership of the first and the last author, started in 2014 with the NBL-Surgical Report Form-project. After the approval of the members of the Surgical Committees of the Children's Oncology Group (COG) and of the German Association of Paediatric Oncology and Haematology (GPOH), the surgical guidelines in the different study protocols of SIOPEN, COG, and GPOH were analyzed to identify the variables needed to completely describe the mode and impact of surgical interventions (biopsy and resection) for NBL, and the outcomes of the surgical intervention. All crucial variables describing the timing, mode, management of IDRFs and the operative results were listed, defined and discussed within the working group of the SIOPEN Surgical Committee.

Thereafter, a first draft of a structured report form was developed, in different steps, discussing each step within the SIOPEN Surgical Committee, until consensus was reached on the Neuroblastoma Surgical Report Form (NSRF). In parallel, the same action was performed within the Surgical Committees of COG and GPOH. Over the years, several joint meetings were organized to discuss and standardize the NSRF draft together with the members of all 3 Surgical Committees to obtain intercontinental consensus. After its conception, the International Neuroblastoma Surgical Report Form (INSRF) was tested and used in clinical practice by the members of the core working party group, confirming its feasibility and completeness.

## RESULTS

The INSRF is the first universal report form to document every NBL-related surgical procedure (biopsy and resection) and contains 5 sections on 5 pages, see Fig. 2. The first 4 sections are ideally completed by the surgeon immediately after the surgical intervention.

The first section of the INSRF includes coded patient and treatment protocol details.

The second section involves the type, timing, and date of the intervention, the surgical approach (open or minimally invasive surgery) and the extent of the biopsy or resection, including 4 options, defined with international consensus: “complete resection;” “minimal residue” (defined as “less than 5 cubic centimeter of tumor remaining”); “incomplete resection” (defined as “5 or more

cubic centimeter residue”) or “other,” for which detailed information is then fully requested.

The localisation of the primary tumor is uniformly reported by the indication of the anatomical compartment(s) (cervical, cervico-thoracic, thoracic, thoraco-abdominal, abdominal/pelvic). Surgical metastatic sites are reported by organ. Preoperative plan discussion at a multidisciplinary tumor board is recorded.

In addition, section 2 incorporates the description of all preoperative post-chemotherapy IDRFs, according to the definitions of terms to describe relationships between primary tumor and vital structures stated in the INRG Guidelines for Imaging and Staging of Neuroblastic Tumors.<sup>8</sup>

The surgical findings are documented in the third section of the INSRF. Here, the surgeon indicates and documents all organs and structures involved, and their individual intraoperative management, by the use of agreed-upon and well-defined surgical terminology. To document vessel involvement, the surgeon can differentiate between 2 types: “adherence” is defined as vessel involvement of <50% of the circumference, and “encasement” as 50%–100% of the vessel circumference.<sup>12</sup> If a blood vessel was injured, the surgeon specifies how this injury was managed during the surgical intervention and if macroscopic residual tumor remained at this location or not. The side of the vessel is indicated and in case of bilateral involvement, detailed specifications per side can be added as free text. In the same fashion, organ involvement is to be reported. When appropriate, the side of involvement is documented and in case of bilateral involvement detailed specifications per side can be added as free text. The type of organ involvement is stated as “adherence” (defined as close contact between tumor and organ but with a distinct plane of dissection) or “infiltration” (no distinct dissection plane, necessitating partial (even minimal) organ resection). The surgeon then documents how the organ involvement was treated by the use of 1 out of 3 options (“no organ resection” – “partial organ resection,” or “complete organ resection”) and whether macroscopic residual tumor was left or not. Liver involvement is described using the segmental classification, as described by Couinaud.<sup>13</sup>

The fourth section of the INSRF documents all intraoperative complications and the management/intervention(s) performed to solve the complication, with enough free text space to add detailed specifications or remarks.

Possible intraoperative deviations from the preoperative plan discussed at a multidisciplinary tumor board can be indicated and described in detail.

The last section of the INSRF is to be completed by the surgeon 1 month after the operation. This final part describes the postoperative complications, related to the surgery, during the first 30 days after the operation, see Fig. 2. To standardize the reporting of postoperative outcomes, the Clavien-Dindo classification was adopted, with space added in the form for free text specifications.<sup>14,15</sup> In addition, the surgeon is expected to report any unscheduled delay of the postoperative chemotherapy regimen(s) due to the surgery or its complications within the first 30 days after the surgical intervention and to specify how long this delay lasted (in days).<sup>11</sup>

## DISCUSSION

We created the first INSRF with intercontinental consensus for the structured and uniform reporting of all NBL-related surgical procedures and their outcomes. Due to the various locations of NBL and frequent encasement or infiltration of vital structures and organs, the surgical treatment of NBL may be highly challenging with important operative morbidity and even mortality.<sup>9,16</sup> In localized NBL, the impact of surgical resection is well established and

International Neuroblastoma Surgical Report Form (INSRF)

Form to be filled out by the surgeon for every surgical procedure

Patient Initials: .....  
 ID system: COG / GPOH / SIOPEN  
 ID number: .....

DATE OF INTERVENTION  
 .. / .. / . . . . (DD/MM/YYYY)

PROTOCOL

- SIOPEN HRNBL2
- LINES
- NBL 2015
- COG PROTOCOL:
- .....

BIOPSY

- Diagnosis
- Relapse / Progression
- Other: ...
  
- Needle core biopsy 14-16 Ga, US guided
- Open
- MIS : Laparoscopic / Thoracoscopic
- Endoscopic: ...
- Other: .....

RESECTION

- Timing**
- Diagnosis
  - During Induction (primary site resection)
  - After Induction chemotherapy
  - After High dose chemotherapy
  - Relapse/progression
  - Other
- Open**
- Laparotomy
  - Thoracophrenolaparotomy
  - Thoracotomy
  - Clamshell
  - Trap-door
  - Transmanubrial
  - Cervical incision
- MIS**
- Laparoscopy
  - Thoracoscopy
  - Both
- Extent of resection**
- Complete excision
  - Minimal Residual Tumor < 5 cm<sup>3</sup> \*
  - Incomplete excision > 5 cm<sup>3</sup> \*
  - Other: ...

TUMOR LOCALISATION (A-F) & PREOPERATIVE IDRF (POST-CHEMOTHERAPY):

EXTENT OF PRIMARY TUMOR – IDRF	Yes	No	N.A.
<b>A Ipsilateral tumor extension within two body compartments</b>			
A.1 Neck-chest			
A.2 Chest-abdomen			
A.3 Abdomen-pelvis			
<b>B Neck</b>			
B.1 Tumor encasing carotid and/or vertebral artery and/or internal jugular vein			
B.2 Tumor extending to base of skull			
B.3 Tumor compressing the trachea			
<b>C Cervico-thoracic junction</b>			
C.1 Tumor encasing brachial plexus roots			
C.2 Tumor encasing subclavian vessels and/or vertebral and/or carotid artery			
C.3 Tumor compressing the trachea			
<b>D Thorax</b>			
D.1 Tumor encasing the aorta and/or major branches			
D.2 Tumor compressing the trachea and/or principal bronchi			
D.3 Lower mediastinal tumor, infiltrating the costo-vertebral junction between T9 and T12			
<b>E Thoraco-abdominal</b>			
E.1 Tumor encasing the aorta and/or vena cava			
<b>F Abdomen/pelvis</b>			
F.1 Tumor infiltrating the porta hepatis (liver hilum) and/or hepatoduodenal ligament			
F.2 Tumor encasing branches of the superior mesenteric artery at the mesenteric root			
F.3 Tumor encasing the origin of the celiac axis and/or of the superior mesenteric artery			
F.4 Tumor invading one or both renal pedicles			
F.5 Tumor encasing the aorta and/or vena cava			
F.6 Tumor encasing the iliac vessels			
F.7 Tumor crossing the sciatic notch			
<b>G Intraspinal tumor extension whatever location provided that:</b>			
G.1 More than one third of the spinal canal in the axial plane is invaded and/or the perimedullary leptomeningeal spaces are not visible and/or the spinal cord signal is abnormal			
<b>H Infiltration of adjacent organs/structures</b>			
H.1 Pericardium			
H.2 Diaphragm			
H.3 Kidney			
H.4 Liver			
H.5 Duodeno-pancreatic block			
H.6 Mesentery			
H.7 Other organ (H.8) considered to be of similar significance			
H.8 Organ (H.7) infiltrated:			

SURGICAL METASTATIC SITE

- Brain
- Liver
- Lung
- Lymph nodes
- Other: ...

Pre-operative plan discussed at a Multidisciplinary Tumor Board (MTB):  NO /  YES

\* 5 cm<sup>3</sup> equals the content of a 5 ml syringe, a spherical tumor measuring 2.1 cm in one dimension, or a cubic tumor measuring 1.7 cm on one side.

FIGURE 2. The International Neuroblastoma Surgical Report Form (INSRF): paper form on 5 pages.

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International Neuroblastoma Surgical Report Form (INSRF)

VASCULAR INVOLVEMENT	Side*	Type** (SURGICAL FINDING):	SURGICAL INJURY	PEROPERATIVE SOLUTION	Macroscopic residual tumor
<input type="checkbox"/> Carotid artery	<input type="checkbox"/> Left <input type="checkbox"/> Right <input type="checkbox"/> Bilateral	<input type="checkbox"/> Adherence <input type="checkbox"/> Encasement	<input type="checkbox"/> No <input type="checkbox"/> Yes	<input type="checkbox"/> Repair <input type="checkbox"/> Reconstruction <input type="checkbox"/> Ligation	<input type="checkbox"/> No <input type="checkbox"/> Yes
<input type="checkbox"/> Internal Jugular vein	<input type="checkbox"/> Left <input type="checkbox"/> Right <input type="checkbox"/> Bilateral	<input type="checkbox"/> Adherence <input type="checkbox"/> Encasement	<input type="checkbox"/> No <input type="checkbox"/> Yes	<input type="checkbox"/> Repair <input type="checkbox"/> Reconstruction <input type="checkbox"/> Ligation	<input type="checkbox"/> No <input type="checkbox"/> Yes
<input type="checkbox"/> Subclavian/innominate artery	<input type="checkbox"/> Left <input type="checkbox"/> Right <input type="checkbox"/> Bilateral	<input type="checkbox"/> Adherence <input type="checkbox"/> Encasement	<input type="checkbox"/> No <input type="checkbox"/> Yes	<input type="checkbox"/> Repair <input type="checkbox"/> Reconstruction <input type="checkbox"/> Ligation	<input type="checkbox"/> No <input type="checkbox"/> Yes
<input type="checkbox"/> Subclavian/innominate vein	<input type="checkbox"/> Left <input type="checkbox"/> Right <input type="checkbox"/> Bilateral	<input type="checkbox"/> Adherence <input type="checkbox"/> Encasement	<input type="checkbox"/> No <input type="checkbox"/> Yes	<input type="checkbox"/> Repair <input type="checkbox"/> Reconstruction <input type="checkbox"/> Ligation	<input type="checkbox"/> No <input type="checkbox"/> Yes
<input type="checkbox"/> Superior vena cava	<input type="checkbox"/> Left <input type="checkbox"/> Right <input type="checkbox"/> Bilateral	<input type="checkbox"/> Adherence <input type="checkbox"/> Encasement	<input type="checkbox"/> No <input type="checkbox"/> Yes	<input type="checkbox"/> Repair <input type="checkbox"/> Reconstruction <input type="checkbox"/> Ligation	<input type="checkbox"/> No <input type="checkbox"/> Yes
<input type="checkbox"/> Pulmonary artery	<input type="checkbox"/> Left <input type="checkbox"/> Right <input type="checkbox"/> Bilateral	<input type="checkbox"/> Adherence <input type="checkbox"/> Encasement	<input type="checkbox"/> No <input type="checkbox"/> Yes	<input type="checkbox"/> Repair <input type="checkbox"/> Reconstruction <input type="checkbox"/> Ligation	<input type="checkbox"/> No <input type="checkbox"/> Yes
<input type="checkbox"/> Thoracic Aorta		<input type="checkbox"/> Adherence <input type="checkbox"/> Encasement	<input type="checkbox"/> No <input type="checkbox"/> Yes	<input type="checkbox"/> Repair <input type="checkbox"/> Reconstruction <input type="checkbox"/> Ligation	<input type="checkbox"/> No <input type="checkbox"/> Yes
<input type="checkbox"/> Abdominal Aorta		<input type="checkbox"/> Adherence <input type="checkbox"/> Encasement	<input type="checkbox"/> No <input type="checkbox"/> Yes	<input type="checkbox"/> Repair <input type="checkbox"/> Reconstruction <input type="checkbox"/> Ligation	<input type="checkbox"/> No <input type="checkbox"/> Yes
<input type="checkbox"/> Portal vein		<input type="checkbox"/> Adherence <input type="checkbox"/> Encasement	<input type="checkbox"/> No <input type="checkbox"/> Yes <input type="checkbox"/>	<input type="checkbox"/> Repair <input type="checkbox"/> Reconstruction <input type="checkbox"/> Ligation	<input type="checkbox"/> No <input type="checkbox"/> Yes
<input type="checkbox"/> Celiac axis		<input type="checkbox"/> Adherence <input type="checkbox"/> Encasement	<input type="checkbox"/> No <input type="checkbox"/> Yes	<input type="checkbox"/> Repair <input type="checkbox"/> Reconstruction <input type="checkbox"/> Ligation	<input type="checkbox"/> No <input type="checkbox"/> Yes
<input type="checkbox"/> Superior mesenteric artery		<input type="checkbox"/> Adherence <input type="checkbox"/> Encasement	<input type="checkbox"/> No <input type="checkbox"/> Yes	<input type="checkbox"/> Repair <input type="checkbox"/> Reconstruction <input type="checkbox"/> Ligation	<input type="checkbox"/> No <input type="checkbox"/> Yes
<input type="checkbox"/> Renal artery	<input type="checkbox"/> Left <input type="checkbox"/> Right <input type="checkbox"/> Bilateral	<input type="checkbox"/> Adherence <input type="checkbox"/> Encasement	<input type="checkbox"/> No <input type="checkbox"/> Yes	<input type="checkbox"/> Repair <input type="checkbox"/> Reconstruction <input type="checkbox"/> Ligation <input type="checkbox"/> Nephrectomy	<input type="checkbox"/> No <input type="checkbox"/> Yes
<input type="checkbox"/> Renal vein	<input type="checkbox"/> Left <input type="checkbox"/> Right <input type="checkbox"/> Bilateral	<input type="checkbox"/> Adherence <input type="checkbox"/> Encasement	<input type="checkbox"/> No <input type="checkbox"/> Yes	<input type="checkbox"/> Repair <input type="checkbox"/> Reconstruction <input type="checkbox"/> Ligation <input type="checkbox"/> Nephrectomy	<input type="checkbox"/> No <input type="checkbox"/> Yes
<input type="checkbox"/> Inferior mesenteric artery		<input type="checkbox"/> Adherence <input type="checkbox"/> Encasement	<input type="checkbox"/> No <input type="checkbox"/> Yes	<input type="checkbox"/> Repair <input type="checkbox"/> Reconstruction <input type="checkbox"/> Ligation	<input type="checkbox"/> No <input type="checkbox"/> Yes
<input type="checkbox"/> Iliac artery	<input type="checkbox"/> Left <input type="checkbox"/> Right <input type="checkbox"/> Bilateral	<input type="checkbox"/> Adherence <input type="checkbox"/> Encasement	<input type="checkbox"/> No <input type="checkbox"/> Yes	<input type="checkbox"/> Repair <input type="checkbox"/> Reconstruction <input type="checkbox"/> Ligation	<input type="checkbox"/> No <input type="checkbox"/> Yes
<input type="checkbox"/> Iliac vein	<input type="checkbox"/> Left <input type="checkbox"/> Right <input type="checkbox"/> Bilateral	<input type="checkbox"/> Adherence <input type="checkbox"/> Encasement	<input type="checkbox"/> No <input type="checkbox"/> Yes	<input type="checkbox"/> Repair <input type="checkbox"/> Reconstruction <input type="checkbox"/> Ligation	<input type="checkbox"/> No <input type="checkbox"/> Yes
<input type="checkbox"/> Inferior vena cava		<input type="checkbox"/> Adherence <input type="checkbox"/> Encasement	<input type="checkbox"/> No <input type="checkbox"/> Yes	<input type="checkbox"/> Repair <input type="checkbox"/> Reconstruction <input type="checkbox"/> Ligation	<input type="checkbox"/> No <input type="checkbox"/> Yes
<input type="checkbox"/> Other: ...	<input type="checkbox"/> Left <input type="checkbox"/> Right <input type="checkbox"/> Bilateral	<input type="checkbox"/> Adherence <input type="checkbox"/> Encasement	<input type="checkbox"/> No <input type="checkbox"/> Yes	<input type="checkbox"/> Repair <input type="checkbox"/> Reconstruction <input type="checkbox"/> Ligation	<input type="checkbox"/> No <input type="checkbox"/> Yes

\* Please tick the involved side. If Bilateral, please specify surgical details per side clearly as free text

\*\* Type of vessel involvement: <50% of circumference of the vessel = Adherence ; 50-100% of circumference of the vessel = Encasement.

(Brisse HJ et al., Guidelines for Imaging & Staging of Neuroblastic Tumors: Consensus Report from the INRG Project. Radiology. 2011;261:243-57)

FIGURE 2. (Continued).

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International Neuroblastoma Surgical Report Form (INSRF)

ORGAN INVOLVEMENT*	Type*** (SURGICAL FINDING)	SURGICAL TREATMENT of organ involved	Macroscopic residual tumor
<input type="checkbox"/> Skull base	<input type="checkbox"/> Adherence <input type="checkbox"/> Infiltration	<input type="checkbox"/> No organ resection <input type="checkbox"/> Partial organ resection: .....	<input type="checkbox"/> No <input type="checkbox"/> Yes
<input type="checkbox"/> Thyroid	<input type="checkbox"/> Adherence <input type="checkbox"/> Infiltration	<input type="checkbox"/> No organ resection <input type="checkbox"/> Partial organ resection	<input type="checkbox"/> No <input type="checkbox"/> Yes
<input type="checkbox"/> Trachea <input type="checkbox"/> Main bronchus: <input type="checkbox"/> Right* <input type="checkbox"/> Left*	<input type="checkbox"/> Adherence <input type="checkbox"/> Infiltration	<input type="checkbox"/> No organ resection <input type="checkbox"/> Partial organ resection	<input type="checkbox"/> No <input type="checkbox"/> Yes
<input type="checkbox"/> Lung: <input type="checkbox"/> Right* <input type="checkbox"/> Left* <input type="checkbox"/> Bilateral*	<input type="checkbox"/> Adherence <input type="checkbox"/> Infiltration	<input type="checkbox"/> No organ resection <input type="checkbox"/> Partial organ resection: .....	<input type="checkbox"/> No <input type="checkbox"/> Yes
<input type="checkbox"/> Heart	<input type="checkbox"/> Adherence <input type="checkbox"/> Infiltration	<input type="checkbox"/> No organ resection <input type="checkbox"/> Partial organ resection: .....	<input type="checkbox"/> No <input type="checkbox"/> Yes
<input type="checkbox"/> Pericardium	<input type="checkbox"/> Adherence <input type="checkbox"/> Infiltration	<input type="checkbox"/> No organ resection <input type="checkbox"/> Partial organ resection: .....	<input type="checkbox"/> No <input type="checkbox"/> Yes
<input type="checkbox"/> Diaphragm: <input type="checkbox"/> Right* <input type="checkbox"/> Left* <input type="checkbox"/> Bilateral*	<input type="checkbox"/> Adherence <input type="checkbox"/> Infiltration	<input type="checkbox"/> No organ resection <input type="checkbox"/> Partial organ resection	<input type="checkbox"/> No <input type="checkbox"/> Yes
<input type="checkbox"/> Kidney: <input type="checkbox"/> Right* <input type="checkbox"/> Left* <input type="checkbox"/> Bilateral*	<input type="checkbox"/> Adherence <input type="checkbox"/> Infiltration	<input type="checkbox"/> No organ resection <input type="checkbox"/> Partial organ resection: .....	<input type="checkbox"/> No <input type="checkbox"/> Yes
<input type="checkbox"/> Liver, specify segment(s) **: .....	<input type="checkbox"/> Adherence <input type="checkbox"/> Infiltration	<input type="checkbox"/> No organ resection <input type="checkbox"/> Partial organ resection: .....	<input type="checkbox"/> No <input type="checkbox"/> Yes
<input type="checkbox"/> Gallbladder	<input type="checkbox"/> Adherence <input type="checkbox"/> Infiltration	<input type="checkbox"/> No organ resection <input type="checkbox"/> Partial organ resection: .....	<input type="checkbox"/> No <input type="checkbox"/> Yes
<input type="checkbox"/> Pancreas	<input type="checkbox"/> Adherence <input type="checkbox"/> Infiltration	<input type="checkbox"/> No organ resection <input type="checkbox"/> Partial organ resection: .....	<input type="checkbox"/> No <input type="checkbox"/> Yes
<input type="checkbox"/> Spleen	<input type="checkbox"/> Adherence <input type="checkbox"/> Infiltration	<input type="checkbox"/> No organ resection <input type="checkbox"/> Partial organ resection: .....	<input type="checkbox"/> No <input type="checkbox"/> Yes
<input type="checkbox"/> Mesentery	<input type="checkbox"/> Adherence <input type="checkbox"/> Infiltration	<input type="checkbox"/> No organ resection <input type="checkbox"/> Partial organ resection: .....	<input type="checkbox"/> No <input type="checkbox"/> Yes
<input type="checkbox"/> Bowel: specify <input type="checkbox"/> duodenum <input type="checkbox"/> small bowel <input type="checkbox"/> colorectum	<input type="checkbox"/> Adherence <input type="checkbox"/> Infiltration	<input type="checkbox"/> No organ resection <input type="checkbox"/> Partial organ resection: .....	<input type="checkbox"/> No <input type="checkbox"/> Yes
<input type="checkbox"/> Bladder	<input type="checkbox"/> Adherence <input type="checkbox"/> Infiltration	<input type="checkbox"/> No organ resection <input type="checkbox"/> Partial organ resection: .....	<input type="checkbox"/> No <input type="checkbox"/> Yes
<input type="checkbox"/> Ureter: <input type="checkbox"/> Right* <input type="checkbox"/> Left* <input type="checkbox"/> Bilateral*	<input type="checkbox"/> Adherence <input type="checkbox"/> Infiltration	<input type="checkbox"/> No organ resection <input type="checkbox"/> Partial organ resection: .....	<input type="checkbox"/> No <input type="checkbox"/> Yes
<input type="checkbox"/> Uterus	<input type="checkbox"/> Adherence <input type="checkbox"/> Infiltration	<input type="checkbox"/> No organ resection <input type="checkbox"/> Partial organ resection: .....	<input type="checkbox"/> No <input type="checkbox"/> Yes
<input type="checkbox"/> Ovary: <input type="checkbox"/> Right* <input type="checkbox"/> Left* <input type="checkbox"/> Bilateral*	<input type="checkbox"/> Adherence <input type="checkbox"/> Infiltration	<input type="checkbox"/> No organ resection <input type="checkbox"/> Partial resection of organ: .....	<input type="checkbox"/> No <input type="checkbox"/> Yes
<input type="checkbox"/> Costovertebral junction T9-12	<input type="checkbox"/> Intraspinal extension	<input type="checkbox"/> Partial resection through foramen <input type="checkbox"/> Complete resection up to level of foramen	<input type="checkbox"/> No <input type="checkbox"/> Yes
<input type="checkbox"/> Other level spinal foramen: Specify level: .....	<input type="checkbox"/> Intraspinal extension	<input type="checkbox"/> Partial resection through foramen <input type="checkbox"/> Complete resection up to level of foramen	<input type="checkbox"/> No <input type="checkbox"/> Yes
<input type="checkbox"/> Other: ..... Please specify:	<input type="checkbox"/> Adherence <input type="checkbox"/> Infiltration	<input type="checkbox"/> No organ resection <input type="checkbox"/> Partial organ resection: .....	<input type="checkbox"/> No <input type="checkbox"/> Yes

\* Please tick the involved side. If Bilateral, please specify surgical details per side clearly as free text  
 \*\* Liver segmental anatomy according to Couinaud (Couinaud C. Liver Anatomy: Portal (and Suprahepatic) or Biliary Segmentation. Dig Surg 1999;16:459-67)  
 \*\*\* Type of involvement: In close contact with clear dissection plane: Adherence; No clear dissection plane: Infiltration

FIGURE 2. (Continued).

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International Neuroblastoma Surgical Report Form (INSRF)

INTRAOPERATIVE COMPLICATIONS	INTERVENTION	EXTRA SPECIFICATIONS
<b>Intraoperative hemorrhage &gt;30% blood volume</b> <input type="checkbox"/> NO <input type="checkbox"/> YES, please specify location/cause: .....	<input type="checkbox"/> Stable circulation after transfusion a/o fluids <input type="checkbox"/> Requiring inotropic support <input type="checkbox"/> Circulatory arrest – successful resuscitation <input type="checkbox"/> Perioperative death	
<b>Hypovolemia</b> <input type="checkbox"/> NO <input type="checkbox"/> YES, please specify location/cause: .....	<input type="checkbox"/> Stable circulation after transfusion a/o fluids <input type="checkbox"/> Requiring inotropic support <input type="checkbox"/> Circulatory arrest – successful resuscitation <input type="checkbox"/> Perioperative death	
<b>Fluid overload with need for diuretics and/or O2</b> <input type="checkbox"/> NO <input type="checkbox"/> YES, please specify cause:.....	<input type="checkbox"/> Requiring diuretics only <input type="checkbox"/> Requiring inotropic support <input type="checkbox"/> Circulatory arrest – successful resuscitation <input type="checkbox"/> Perioperative death	
<b>Pulmonary embolus</b> <input type="checkbox"/> NO <input type="checkbox"/> YES	<input type="checkbox"/> Stable circulation after transfusion a/o fluids <input type="checkbox"/> Requiring inotropic support <input type="checkbox"/> Circulatory arrest – successful resuscitation <input type="checkbox"/> Perioperative death	
<b>Vascular injury (→ see previous Table)</b> <input type="checkbox"/> NO <input type="checkbox"/> YES	/	
<b>Nerve injury</b> <input type="checkbox"/> NO <input type="checkbox"/> YES, please specify cause: .....	<input type="checkbox"/> No intervention <input type="checkbox"/> Primary repair <input type="checkbox"/> Reconstruction	Nerve(s) affected:
<b>Unplanned injury/removal of an organ other than the affected adrenal</b> <input type="checkbox"/> NO <input type="checkbox"/> YES, please specify cause: .....	<input type="checkbox"/> Partial nephrectomy: <input type="checkbox"/> Right / <input type="checkbox"/> Left <input type="checkbox"/> Total nephrectomy: <input type="checkbox"/> Right / <input type="checkbox"/> Left <input type="checkbox"/> Partial hepatectomy: segment:..... <input type="checkbox"/> Partial splenectomy <input type="checkbox"/> Total splenectomy <input type="checkbox"/> Partial colectomy <input type="checkbox"/> Partial colectomy with colostomy <input type="checkbox"/> Small bowel resection <input type="checkbox"/> Small bowel resection with enterostomy <input type="checkbox"/> Contralateral adrenal (non affected) <input type="checkbox"/> Partial pancreatectomy <input type="checkbox"/> Other: .....	
<b>Renal ischemia:</b> <input type="checkbox"/> NO <input type="checkbox"/> YES, please specify extent: <input type="checkbox"/> Partial, resolved before end of operation <input type="checkbox"/> Partial, persistent at the end of surgery <input type="checkbox"/> Total, resolved before end of operation <input type="checkbox"/> Total, persistent at the end of surgery	<input type="checkbox"/> No intervention <input type="checkbox"/> Renal vessel repair <input type="checkbox"/> Renal vessel reconstruction <input type="checkbox"/> Partial resection <input type="checkbox"/> Total resection	Reason: <input type="checkbox"/> Vascular injury <input type="checkbox"/> Vascular spasm <input type="checkbox"/> Hypovolemia <input type="checkbox"/> Other: .....
<b>Perioperative death</b> <input type="checkbox"/> NO <input type="checkbox"/> YES, please specify cause: <input type="checkbox"/> Bleeding <input type="checkbox"/> Other cause: ..... <input type="checkbox"/> Unknown cause	<input type="checkbox"/> Autopsy performed <input type="checkbox"/> YES <input type="checkbox"/> NO	Specify if known
<b>Other:</b>		

**Were there any intra-operative DEVIATIONS from the Preoperative plan discussed at Multidisciplinary Tumor Board (MTB)?**

NO /  YES → Please specify: .....

**Did this lead to intra- or post-operative complications?**  NO /  YES → Please specify: .....

FIGURE 2. (Continued).

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International Neuroblastoma Surgical Report Form (INSRF)

POSTOPERATIVE COMPLICATIONS <i>up till Postoperative Day 30</i>	CLAVIEN-DINDO CLASSIFICATION (see next page)	SPECIFY
Postoperative bleeding: <input type="checkbox"/> NO / <input type="checkbox"/> YES, specify cause:	<input type="checkbox"/> I <input type="checkbox"/> II <input type="checkbox"/> III-a <input type="checkbox"/> III-b <input type="checkbox"/> IV-a <input type="checkbox"/> IV-b <input type="checkbox"/> V	
Hypovolemia requiring inotropic support: <input type="checkbox"/> NO / <input type="checkbox"/> YES	<input type="checkbox"/> I <input type="checkbox"/> II <input type="checkbox"/> III-a <input type="checkbox"/> III-b <input type="checkbox"/> IV-a <input type="checkbox"/> IV-b <input type="checkbox"/> V	
Fluid overload requiring diuretics +/- O2: <input type="checkbox"/> NO / <input type="checkbox"/> YES	<input type="checkbox"/> I <input type="checkbox"/> II <input type="checkbox"/> III-a <input type="checkbox"/> III-b <input type="checkbox"/> IV-a <input type="checkbox"/> IV-b <input type="checkbox"/> V	
Systemic inflammatory response syndrome (SIRS) requiring intensive care > 72h: <input type="checkbox"/> NO / <input type="checkbox"/> YES	<input type="checkbox"/> I <input type="checkbox"/> II <input type="checkbox"/> III-a <input type="checkbox"/> III-b <input type="checkbox"/> IV-a <input type="checkbox"/> IV-b <input type="checkbox"/> V	
Stroke: <input type="checkbox"/> NO / <input type="checkbox"/> YES	<input type="checkbox"/> I <input type="checkbox"/> II <input type="checkbox"/> III-a <input type="checkbox"/> III-b <input type="checkbox"/> IV-a <input type="checkbox"/> IV-b <input type="checkbox"/> V	
Vascular spinal cord injury: <input type="checkbox"/> NO / <input type="checkbox"/> YES	<input type="checkbox"/> I <input type="checkbox"/> II <input type="checkbox"/> III-a <input type="checkbox"/> III-b <input type="checkbox"/> IV-a <input type="checkbox"/> IV-b <input type="checkbox"/> V	
Renal atrophy (ischemia): <input type="checkbox"/> Right / <input type="checkbox"/> Left <input type="checkbox"/> NO <input type="checkbox"/> YES, specify: <input type="checkbox"/> Partial ischemia <input type="checkbox"/> Total ischemia	<input type="checkbox"/> I <input type="checkbox"/> II <input type="checkbox"/> III-a <input type="checkbox"/> III-b <input type="checkbox"/> IV-a <input type="checkbox"/> IV-b <input type="checkbox"/> V	Diagnosis: <input type="checkbox"/> US <input type="checkbox"/> CT <input type="checkbox"/> Nuclear scan
Renal dysfunction: <input type="checkbox"/> NO / <input type="checkbox"/> YES, specify:	<input type="checkbox"/> I <input type="checkbox"/> II <input type="checkbox"/> III-a <input type="checkbox"/> III-b <input type="checkbox"/> IV-a <input type="checkbox"/> IV-b <input type="checkbox"/> V	
Bladder dysfunction: <input type="checkbox"/> NO / <input type="checkbox"/> YES, specify:	<input type="checkbox"/> I <input type="checkbox"/> II <input type="checkbox"/> III-a <input type="checkbox"/> III-b <input type="checkbox"/> IV-a <input type="checkbox"/> IV-b <input type="checkbox"/> V	
Hepatic dysfunction: <input type="checkbox"/> NO / <input type="checkbox"/> YES, specify:	<input type="checkbox"/> I <input type="checkbox"/> II <input type="checkbox"/> III-a <input type="checkbox"/> III-b <input type="checkbox"/> IV-a <input type="checkbox"/> IV-b <input type="checkbox"/> V	
Pulmonary dysfunction: <input type="checkbox"/> NO / <input type="checkbox"/> YES, specify:	<input type="checkbox"/> I <input type="checkbox"/> II <input type="checkbox"/> III-a <input type="checkbox"/> III-b <input type="checkbox"/> IV-a <input type="checkbox"/> IV-b <input type="checkbox"/> V	
Postop stenosis of vascular structures : <input type="checkbox"/> NO / <input type="checkbox"/> YES:	<input type="checkbox"/> I <input type="checkbox"/> II <input type="checkbox"/> III-a <input type="checkbox"/> III-b <input type="checkbox"/> IV-a <input type="checkbox"/> IV-b <input type="checkbox"/> V	
Infection: <input type="checkbox"/> NO <input type="checkbox"/> YES : <input type="checkbox"/> Fever >48h postop eci, AB administ <input type="checkbox"/> Pneumonia <input type="checkbox"/> Urinary tract <input type="checkbox"/> Wound <input type="checkbox"/> Intravenous access line <input type="checkbox"/> Other, please specify:	<input type="checkbox"/> I <input type="checkbox"/> II <input type="checkbox"/> III-a <input type="checkbox"/> III-b <input type="checkbox"/> IV-a <input type="checkbox"/> IV-b <input type="checkbox"/> V	Specify on postop day...
Adrenal insufficiency : <input type="checkbox"/> NO / <input type="checkbox"/> YES, specify	<input type="checkbox"/> I <input type="checkbox"/> II <input type="checkbox"/> III-a <input type="checkbox"/> III-b <input type="checkbox"/> IV-a <input type="checkbox"/> IV-b <input type="checkbox"/> V	
Intestinal obstruction > 7 days <input type="checkbox"/> NO <input type="checkbox"/> YES, specify: <input type="checkbox"/> Paralytic Ileus <input type="checkbox"/> Mechanical Obstruction	<input type="checkbox"/> I <input type="checkbox"/> II <input type="checkbox"/> III-a <input type="checkbox"/> III-b <input type="checkbox"/> IV-a <input type="checkbox"/> IV-b <input type="checkbox"/> V	
Wound dehiscence <input type="checkbox"/> NO <input type="checkbox"/> YES, specify location & postop day	<input type="checkbox"/> I <input type="checkbox"/> II <input type="checkbox"/> III-a <input type="checkbox"/> III-b <input type="checkbox"/> IV-a <input type="checkbox"/> IV-b <input type="checkbox"/> V	
Chylous leak – Chest : <input type="checkbox"/> NO / <input type="checkbox"/> YES	<input type="checkbox"/> I <input type="checkbox"/> II <input type="checkbox"/> III-a <input type="checkbox"/> III-b <input type="checkbox"/> IV-a <input type="checkbox"/> IV-b <input type="checkbox"/> V	
Chylous leak – Abdomen: <input type="checkbox"/> NO / <input type="checkbox"/> YES	<input type="checkbox"/> I <input type="checkbox"/> II <input type="checkbox"/> III-a <input type="checkbox"/> III-b <input type="checkbox"/> IV-a <input type="checkbox"/> IV-b <input type="checkbox"/> V	
Unplanned ICU admission: <input type="checkbox"/> NO <input type="checkbox"/> YES, specify	<input type="checkbox"/> I <input type="checkbox"/> II <input type="checkbox"/> III-a <input type="checkbox"/> III-b <input type="checkbox"/> IV-a <input type="checkbox"/> IV-b <input type="checkbox"/> V	Postop day:: Cause: Treatment:
Unplanned return to operating theater <input type="checkbox"/> NO <input type="checkbox"/> YES, specify	<input type="checkbox"/> I <input type="checkbox"/> II <input type="checkbox"/> III-a <input type="checkbox"/> III-b <input type="checkbox"/> IV-a <input type="checkbox"/> IV-b <input type="checkbox"/> V	Postop day:: Cause: Treatment::
Diarrhea 30 days postop (without infectious cause) <input type="checkbox"/> NO <input type="checkbox"/> YES	<input type="checkbox"/> I <input type="checkbox"/> II <input type="checkbox"/> III-a <input type="checkbox"/> III-b <input type="checkbox"/> IV-a <input type="checkbox"/> IV-b <input type="checkbox"/> V	
Motor Nerve dysfunction: <input type="checkbox"/> NO <input type="checkbox"/> YES, please specify cause: .....	<input type="checkbox"/> I <input type="checkbox"/> II <input type="checkbox"/> III-a <input type="checkbox"/> III-b <input type="checkbox"/> IV-a <input type="checkbox"/> IV-b <input type="checkbox"/> V	Nerve(s) affected:
Sensory Nerve dysfunction: <input type="checkbox"/> NO <input type="checkbox"/> YES, please specify cause: .....	<input type="checkbox"/> I <input type="checkbox"/> II <input type="checkbox"/> III-a <input type="checkbox"/> III-b <input type="checkbox"/> IV-a <input type="checkbox"/> IV-b <input type="checkbox"/> V	Nerve(s) affected:
Postoperative death <input type="checkbox"/> NO <input type="checkbox"/> YES, specify cause: <input type="checkbox"/> Bleeding <input type="checkbox"/> Other: specify: <input type="checkbox"/> Unknown cause	<input type="checkbox"/> V	When? postop day ..... Autopsy performed?: Findings

**Was there an unscheduled DELAY of postoperative chemotherapy regimen?**  NO /  YES → Delay in number of days: .....

**Was this delay of postoperative chemotherapy due to surgery?**  NO /  YES → Cause: .....

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FIGURE 2. (Continued).



## International Neuroblastoma Surgical Report Form (INSRF)

## Clavien-Dindo Classification\* of Surgical Complications

<b>Grade I:</b>	<b>Any deviation from the normal postoperative course without the need for pharmacological treatment or surgical, endoscopic and radiological interventions.</b> Acceptable therapeutic regimens are: drugs as antiemetics, antipyretics, analgesics, diuretics and electrolytes and physiotherapy. This grade also includes wound infections opened at the bedside.
<b>Grade II:</b>	Requiring pharmacological treatment with drugs other than such allowed for grade I complications. Blood transfusions and total parenteral nutrition are also included.
<b>Grade III:</b>	<b>Requiring surgical, endoscopic or radiological intervention</b>
<b>Grade III-a:</b>	Intervention not under general anesthesia
<b>Grade III-b:</b>	Intervention under general anesthesia
<b>Grade IV:</b>	<b>Life-threatening complication (including CNS complications)<sup>o</sup> requiring IC/ICU-management</b>
<b>Grade IV-a:</b>	Single organ dysfunction (including dialysis)
<b>Grade IV-b:</b>	Multi organ dysfunction
<b>Grade V:</b>	<b>Death of a patient</b>
<b>Suffix 'd':</b>	If the patient suffers from a complication at the time of discharge. The suffix 'd' (for ' <b>disability</b> ') is added to the respective grade of complication. This label indicates <b>the need for a follow-up</b> to fully evaluate the complication.

<sup>o</sup>brain hemorrhage, ischemic stroke, subarachnoidal bleeding, but excluding transient ischemic attacks (TIA); IC: intermediate care; ICU: intensive care unit

\*Dindo D, Demartines N, Clavien PA. Classification of surgical complications: a new proposal with evaluation in a cohort of 6336 patients and results of a survey. *Ann Surg*. 2004;240:205–213  
Clavien PA, Barkun J, de Oliveira ML et al. The Clavien-Dindo Classification of Surgical Complications. Five-Year Experience. *Ann Surg* 2009;250:187–196

FIGURE 2. (Continued).

surgical resection is an important part in the multimodal treatment protocols.<sup>17,18</sup>

In high-risk NBL and especially in case of distant metastatic disease, the role of surgical resection remains an unsolved issue even after more than 25 years of debate, with many authors favoring complete or gross total resection of the primary tumor<sup>10,11,19–24</sup> – whereas other authors question the role of aggressive surgical resection for high-risk NBL.<sup>9,11,12,25</sup> The reasons for these contradictory conclusions are linked to differences in protocols, especially indication, dose and field of adjuvant radiotherapy,<sup>26</sup> but also to the subjective appraisal of the completeness of resection, which relies predominantly on the surgeon's report. The assessment of the postoperative residue on imaging is not standardized and differently taken into account across the cooperative protocols.<sup>19,25,27</sup>

Postoperative irradiation has an important role in the multimodal treatment of high-risk NBL but is implicated in numerous late toxicities, including impairment in musculoskeletal growth, fertility, cardiopulmonary function and endocrinopathies, bladder dysfunction poor psychosocial health, and secondary malignancies.<sup>24</sup> The new SIOOPEN HR-NBL2 protocol will randomize patients with macroscopic residual disease to receive a radiation boost of 36 Gray on postoperative residue, in addition to the baseline irradiation field of 21 Gray (depending on the preoperative tumor volume)

(unpublished data). The patient will be considered to have no macroscopic residue at the time of radiotherapy if, cumulatively:

- the postoperative magnetic resonance imaging (or computed tomography scan, if no magnetic resonance imaging available) shows no definite residual tumor and
- the postoperative metaiodobenzylguanidine scan shows no residual tumor and
- the surgical report mentions a complete or minimal residual resection (<5 cm<sup>3</sup> residual tumor remaining, according to international consensus (unpublished data) agreed upon during joint meetings with the surgeons and oncologists of the SIOOPEN, COG, and GPOH surgical committees (Fig. 2).

Postoperative treatment will; therefore, rely on the surgeon's estimation and image-based description of the residue.

A recent analysis of the COG A3973 data did not show a statistically significant difference in outcomes based on the extent of prophylactic lymph node irradiation, regardless of the degree of surgical resection.<sup>24</sup> Although awaiting the results of the COG phase 3 trial ANBL0532, lowering the volume of postoperative irradiation and adapting an eventual boost on postoperative residue may also become part of upcoming COG NBL trials.<sup>24</sup> (unpublished data, personal communication).

Adequate documentation by the surgeon of the extent of surgical resection and of the volume and localization of postoperative residue, will; therefore, become essential and will need to rely on uniform, structured reporting, guided by clear-cut, unequivocal definitions.<sup>12</sup>

Recent analysis of 220 patients in the COG A3973 study (evaluating the impact of extent of primary tumor resection on local progression and survival and the assessed concordance between clinical and central imaging review-based assessments of resection extent), revealed however an important discordance (37%) between the operating surgeon's assessment of the extent of resection and imaging-aided assessment.<sup>19</sup> This may be related to the pitfalls of narrative operative reports as their content is not standardized nor regulated and may therefore be of variable quality.<sup>28,29</sup> In cancer surgery, several authors have recently pointed out that narrative operative reports are seldom complete and may be of poor quality<sup>29–34</sup> suggesting the development and use of standardized operative reports, also known as synoptic operative reports (SORs). Electronic SORs have been developed and implemented with proven benefit and multiple studies proving a gain in time.<sup>35</sup> In addition, structured operative reporting may even be beneficial for surgical education.<sup>36</sup>

The first version of the INSRF presented here is not intended to be a SOR: it is at present a standardized surgical report form, conceived as a structured checklist for the uniform registration of important variables and relevant clinical information on the surgery of NBL and its immediate outcomes. The obligatory standardized registration of intra- and postoperative complications according to the Clavien-Dindo classification<sup>14,15</sup> will improve the quality of data and facilitate even more the international comparison of different surgical timings, approaches, extents of resection and their outcomes.<sup>19</sup>

As the INSRF incorporates standardized reporting of the preoperative post-chemotherapy IDRFs, it will also aid in the analysis of the role of pre- versus post-chemotherapy IDRFs and tumor volumes.<sup>4,7,8,12,37,38</sup>

Furthermore, the INSRF may be used as well for the reporting of surgical interventions of other neuroblastic tumors (ie. ganglioneuroblastoma, ganglioneuroma, . . .) – where there is also still controversy on the approach, extent, and timing of surgical treatment.<sup>8,37</sup>

After its use by the members of the core working party group, confirming the feasibility, user-friendliness, and completeness of the INSRF, next steps in this joint international collaboration will be the further implementation into clinical practice, by the members of the surgical committees of SIOPEN, COG, and GPOH, to collect feedback from the individual users and further study adherence to the form. Interested pediatric surgical oncologists from other continents are also kindly invited to join this project. Development of an electronic web-based INSRF is highly recommended to allow surgeons to fill out the form immediately after surgery, warranting adherence, and the quality of data. The inclusion of an automated reminder to the surgeon 30 days after the operation to complete the registration of postoperative complications will ensure, as it was demonstrated in adult SORs for cancer,<sup>39</sup> higher rates of essential data completeness, intra- and interobserver reliability and faster and more efficient data entry.

The INSRF is highly compatible with different treatment protocols for localized, intermediate, and high-risk NBL. Its systematic incorporation in the new upcoming international NBL protocols of all cooperative groups will allow better analysis and definition of the surgical strategy in NBL treatment and comparison of local control modalities between international and cooperative groups.

In conclusion, the INSRF is the first attempt towards a universal operative report form for the structured and uniform reporting of all NBL-related surgical procedures. By documenting important perioperative data and outcomes, the INSRF will facilitate the analysis of the surgical treatment of NBL.

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