Soft Tissue Sarcoma

Soft code: STS

Procedure: Intralesional resection

Marginal resection
Wide resection
Radical resection
Other:
Not specified

Tumor type (based on WHO classification):

	ors

Atypical lipomatous tumor

Well-differentiated liposarcoma

Liposarcoma, NOS

Dedifferentiated liposarcoma

Myxoid/round cell liposarcoma

Pleomorphic liposarcoma

Myofibroblastic Tumors

Dermatofibrosarcoma protuberans

Fibrosarcomatous dermatofibrosarcoma protuberans

Pigmented dermatofibrosarcoma protuberans

Solitary fibrous tumor, malignant

Inflammatory myofibroblastic tumor

Low-grade myofibroblastic sarcoma

Adult fibrosarcoma

Myxofibrosarcoma (formerly myxoid malignant fibrous histiocytoma [myxoid MFHI])

Low-grade fibromyxoid sarcoma

Sclerosing epithelioid fibrosarcoma

Fibrohistiocytic Tumors

Giant cell tumor of soft tissues

Muscle Tumors

Leiomyosarcoma (excluding skin)

(Perivascular) Tumors

Malignant glomus tumor

Muscle Tumors

Embryonal rhabdomyosarcoma (including botryoid, anaplastic)

Alveolar rhabdomyosarcoma (including solid, anaplastic)

Pleomorphic rhabdomyosarcoma

Spindle cell/sclerosing rhabdomyosarcoma

Vascular Tumors of Soft Tissue

Retiform hemangioendothelioma

Pseudomyogenic (epitihelioid sarcoma-like) hemangioendothelioma

Epithelioid hemangioendothelioma

Angiosarcoma of soft tissue

Chondro-osseus Tumors

Extraskeletal osteosarcoma

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Gastrointestinal Stromal Tumors

Gastrointestinal stromal tumor, malignant

Nerve Sheath Tumors

Malignant peripheral nerve sheath tumor

Epithelioid malignant peripheral nerve sheath tumor

Malignant Triton tumor

Malignant granular cell tumor

Tumors of Uncertain Differentiation

Ossifying fibromyxoid tumor, malignant

Stromal sarcoma, NOS

Myoepithelial carcinoma

Phosphaturic mesenchymal tumor, malignant

Synovial sarcoma, NOS

Synovial sarcoma, spindle cell

Synovial sarcoma, biphasic

Epithelioid sarcoma

Alveolar soft part sarcoma

Clear cell sarcoma of soft tissue

Extraskeletal myxoid chondrosarcoma

Extraskeletal Ewing sarcoma

Desmoplastic small round cell tumor

Extrarenal rhabdoid tumor

Perivascular epithelioid cell tumor (PEComa), NOS

Intimal sarcoma

Undifferentiated/Unclassified Sarcomas

Undifferentiated spindle cell sarcoma

Undifferentiated pleomorphic sarcoma

Undifferentiated round cell sarcoma

Undifferentiated epithelioid sarcoma

Undifferentiated sarcoma, NOS

Tumor site:	
Tumor size	

__ x__ x__ cm

Localization

- Superficial
 - o Dermal
 - Subcutaneous (suprafacial)
- Deep
 - o Fascial
 - o Subfascial
 - o Intramuscular
 - o Mediastinal
 - o Intra-abdominal
 - o Retroperitoneal
 - Head and neck
 - Other (specify)_____

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Indeterminate	
Mitotic rate/10 HPF	
Percent Necrosis Not identified Present, percent:	
Grade (based on FNCLCC) 1 2 3 Ungraded sarcoma Cannot be determined	
 Margin status/Procedure Positive margin (tumor touching ink), site(s): Marginal resection (ink on pseudocapsule), site and distance to close Wide resection (layer of normal tissue surrounding tumor), site and Cannot be assessed 	
 Lymph-vascular invasion Present Not identified Indeterminate 	
Post-treatment effect percentage (based on combined gross and microsco Percent present: Not identified Indeterminate	opic examination)
Staging, TMN categories (AJCC, 8th ed.)	
TX-Primary tumor cannot be assessed T0-No evidence of primary tumor T1-Tumor ≤5 cm in greatest dimension T2-Tumor >5 cm and less than or equal to 10 cm in greatest dimension T3-Tumor >10 cm and less than or equal to 15 cm in greatest dimension	
N0-No regional lymph nodes or unknown lymph node status N1-Regional lymph node metastasis	
M0-No distant metastasis or unknown distant metastasis M1-Distant metastasis	