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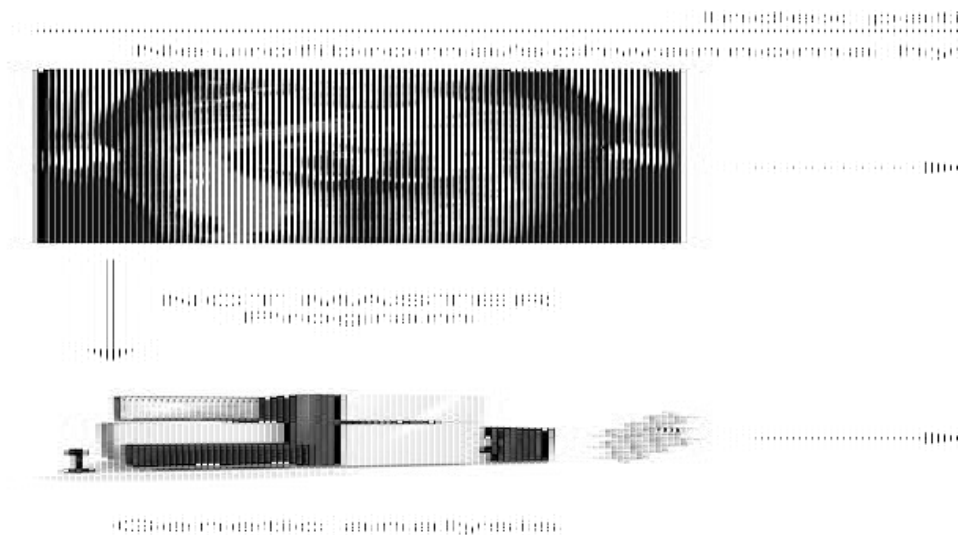
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Targetable ERBB2 Mutations Identified in Neurofibroma/Schwannoma Hybrid Nerve Sheath Tumors

Running Title: ERBB2 Mutations in Neurofibroma/Schwannoma Hybrid Tumors

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Abstract

Background: Neurofibroma/schwannoma hybrid nerve sheath tumors (N/S HNSTs) are neoplasms associated with larger nerves that occur sporadically and in the context of schwannomatosis or neurofibromatosis type 2 or 1. Clinical management of N/S HNST is challenging, especially for large tumors, and established systemic treatments are lacking.

Methods: We used next-generation sequencing and array-based DNA methylation profiling to determine the clinically actionable genomic and epigenomic landscapes of N/S HNST.

Results: Whole-exome sequencing within a precision oncology program identified an activating mutation (p.Asp769Tyr) in the catalytic domain of the ERBB2 receptor tyrosine kinase in a patient with schwannomatosis-associated N/S HNST, and targeted treatment with the small-molecule ERBB inhibitor lapatinib led to prolonged clinical benefit and a lasting radiographic and metabolic response. Analysis of a multicenter validation cohort revealed recurrent ERBB2 mutations (p.Leu755Ser, p.Asp769Tyr, p.Val777Leu) in N/S HNSTs occurring in patients who met diagnostic criteria for sporadic schwannomatosis (3 of 7 patients), but not in N/S HNSTs arising in the context of neurofibromatosis (6 patients) or outside a tumor syndrome (1 patient), and showed that ERBB2-mutant N/S HNSTs cluster in a distinct subgroup of peripheral nerve sheath tumors based on genome-wide DNA methylation patterns.

Conclusion: These findings uncover a key biological feature of N/S HNST that may have important diagnostic and therapeutic implications.

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Introduction

Neurofibroma/schwannoma hybrid nerve sheath tumors (N/S HNSTs) are peripheral nerve sheath tumors that are recognized as a distinct entity by the 2016 update of the World Health Organization (WHO) Classification of Tumours of the Central Nervous System (1). While schwannomas are solely composed of Schwann cells, neurofibromas are heterogeneous tumors consisting of Schwann cells, axonal processes, perineurial/perineurial-like cells, fibroblasts, and mast cells (2). N/S HNSTs are composed of schwannoma-like nodules within a neurofibroma-like tumor (3), and more than 60% of cases are associated with an underlying tumor predisposition syndrome. In particular, 26%, 17%, and 9% of N/S HNST patients are afflicted with neurofibromatosis type 2 (NF2), schwannomatosis, and neurofibromatosis type 1 (NF1), respectively (4); 71% of patients with schwannomatosis develop N/S HNST (4); and 26% of NF2 patients undergoing resection of peripheral nerve sheath tumors have N/S HNST (5). Thus, N/S HNST may occur in up to 1/10,000 persons per year, assuming a birth incidence for schwannomatosis of 1/40,000 (6).

Treatment of N/S HNST is based on surgical resection. However, due to the association of neurofibromas, schwannomas, and N/S HNSTs with larger nerves as well as the number and extent of manifestations typically seen in cases of NF1-associated plexiform nerve sheath tumors (7), local control is sometimes difficult to achieve. Thus, despite generally being considered benign due to WHO grade I histology in both tumor components, N/S HNSTs can be associated with substantial morbidity, such as motor disability, neuropathic pain, or disfigurement (6). In patients who are not surgical candidates, palliative radiotherapy may be considered depending on the localization and extent of the tumor(s). In contrast, although clinical trials have been conducted in subgroups of schwannomas and neurofibromas (8, 9),

there is no established systemic treatment for N/S HNSTs, and molecular drug targets are lacking. Thus, there is an urgent need for novel treatment options to improve clinical outcomes in N/S HNST patients, especially those with unresectable symptomatic tumors.

We here report successful targeted treatment of an index patient with schwannomatosis-associated N/S HNST harboring an activating ERBB2 mutation within a genomics-guided precision oncology program, and describe for the first time that a substantial proportion of N/S HNSTs are associated with recurrent, pharmacologically tractable ERBB2 mutations.

Results

A 30-year-old woman presented with pain in her right leg in 2004. Magnetic resonance imaging (MRI) showed a large tumor mass comprising multiple nodules located around peripheral nerves in the lower right body quadrant, including lumbar and sacral spinal nerve roots, the right lumbosacral plexus, and the right femoral and sciatic nerves. Additional nodules were present in the abdomen and the left leg. Partial resection of nodules in the ventral compartment of the upper right thigh and the right gluteal region was performed in 2005, and a diagnosis of N/S HNST was established. There were no clinical signs of NF2, such as vestibular schwannomas or dermatologic abnormalities, and the patient fulfilled the clinical criteria of sporadic schwannomatosis (10) with no family history suggestive of a hereditary tumor predisposition syndrome. Over time, the tumor slowly progressed (Supplementary Fig. 1), and partial resection of a large tumor mass around the right lateral femoral cutaneous nerve was performed in 2011. Hematoxylin and eosin (HE) staining showed an encapsulated tumor with compact, cell-dense areas as well as areas with scattered, spindle-shaped cells with filament fragments. Focally, a slight lymphoid infiltrate was detectable in perivascular regions. The tumor cells displayed strong expression of S100 and slight expression of reticulin in a punctuated pattern around balloon-shaped swirls. The proliferation rate was low (average, 1%; range, 0-3%) (Fig. 1A).

Due to progressive pareses and intense neuropathic pain and because no further established therapy was available, the patient was enrolled in the NCT MASTER (Molecularly Aided Stratification for Tumor Eradication Research) program, a nationwide registry trial for genomics-guided stratification of younger adults with advanced-stage cancer across all histologies and patients with rare tumors who have exhausted standard treatment options (11). Whole-exome sequencing of tumor tissue

and matched peripheral blood mononuclear cells showed a “quiet” genomic profile with only two somatic missense mutations and no DNA copy number alterations. Of particular clinical interest was a heterozygous *ERBB2* p.Asp769Tyr mutation (variant allele frequency, 16%), which was confirmed by Sanger sequencing both in the initial specimen and a subsequent biopsy from a different tumor nodule, and whose expression was verified by RNA sequencing. Analysis of germline sequences (Supplementary Table 1) identified no pathogenic variants in established tumor predisposition genes, including *NF1*, *NF2*, *SMARCB1*, and *LZTR1*.

ERBB2 p.Asp769Tyr has been identified as a rare mutation in a range of carcinomas, in particular breast invasive ductal carcinoma (<http://www.cbioportal.org>), and studies in MCF10A human mammary epithelial cells have shown that this variant causes constitutive activation of the *ERBB2* receptor tyrosine kinase (RTK) and confers sensitivity to *ERBB2*-directed antibodies and small-molecule tyrosine kinase inhibitors (12). Given these findings and the lack of therapeutic alternatives, treatment with 1,500 mg lapatinib daily was initiated in October 2015 and has since then been well tolerated. Prior to lapatinib treatment, the patient had bilateral paresis of foot and toe elevation (2/5 on the right and 4/5 on the left side according to the British Medical Research Council Scale) with corresponding gait impairment. There was numbness of the right foot and lower leg as well as the medial part of the right thigh. Severe neuropathic pain (5-6/10 on a numeric rating scale) was present at all times despite treatment with gabapentin and nortriptyline and caused frequent insomnia. After 6 months of therapy, pareses had improved to 4+/5 on the right and 4+/5 on the left side and gait was no longer impaired. Furthermore, neuropathic pain had improved (1-2/10 on a visual analog scale) and was only present on 1-2 days per week for brief periods, and sleep was rarely affected. In contrast, the numbness of the right leg remained unchanged. As of October 2019, the patient remains on

lapatinib, and the improvements of motor strength and neuropathic pain, which had begun as early as 4 weeks after initiation of therapy, are lasting. Interestingly, a 3-week interruption of lapatinib due to paronychia of the right hallux caused an increase in neuropathic pain, which subsided after reintroduction of therapy, indicating continued efficacy of ERBB2 blockade. Clinical improvement was paralleled by a reduction in size of some lesions (Fig. 1B, Supplementary Fig. 2A, B, C) as well as a lasting metabolic response on sequential fluorodeoxyglucose-positron emission tomography (FDG-PET)/MRI studies (Fig. 1C, Supplementary Fig. 2B). Together, these results uncover mutant ERBB2 as a novel driver of N/S HNST that provides an immediate therapeutic opportunity.

To investigate the frequency of mutations in ERBB2 or other RTKs in N/S HNST, we collected 18 additional tumors from 14 additional patients (Supplementary Table 2). This cohort included 7 patients that met diagnostic criteria for sporadic schwannomatosis, 2 of which had unusual clinical characteristics in addition to peripheral nerve sheath tumors (patient 3: marfanoid habitus, scoliosis, bilateral acetabular protrusions, mitral valve prolapse, dilated ascending aorta, history of retrognathia and juvenile glaucoma; patient 4: generalized epileptic seizures, dermal neurofibromas), 6 patients with NF2, and 1 patient without a tumor predisposition syndrome (Table 1).

Employing targeted next-generation sequencing, we detected ERBB2 mutations (p.Leu755Ser, p.Asp769Tyr, p.Val777Leu) in 3 of 7 (43%) schwannomatosis patients, whereas no ERBB2 mutations were found in NF2 or sporadic N/S HNST cases, resulting in a mutation frequency of 27% (4 of 15 patients) in the entire cohort including the index patient (Table 1). These kinase domain mutations had previously been found in breast cancer and characterized as activating and sensitive to pan-ERBB inhibitors such as neratinib (12). Of the 6 patients with

NF2, 1 (17%) had tumors that harbored a heterozygous *RET* p.Tyr791Phe mutation (Table 1), which has been identified in various cancers and characterized as activating, although its cancer-driving role has been disputed (13). Histologic analysis revealed no phenotypic differences compared to samples without RTK alterations (Supplementary Fig. 3).

To delineate potential additional steps of tumorigenesis and corroborate the clinical diagnoses, we analyzed the sequencing data and genome-wide DNA methylation profiles of all tumors for mutations in *NF1*, *NF2*, *SMARCB1*, or *LZTR1* and for loss of heterozygosity of chromosome 22q, where *NF2* is located, respectively (Table 2), and observed that ERBB2 mutations were mutually exclusive with alterations of *NF2*, *SMARCB1*, or *LZTR1* and loss of heterozygosity of chromosome 22q. No *NF1* mutation was detected in the study cohort (Table 2). Of note, both *RET*-mutant tumors from a patient who had been diagnosed with NF2 lacked *NF2* alterations, whereas the mutational profiles of all other tumors were consistent with the respective clinical diagnoses. Collectively, these results identify activating ERBB2 mutations as recurrent events in N/S HNST arising in schwannomatosis patients, and suggest that this tumor entity may be driven by aberrant ERBB2 signaling in a substantial proportion of cases.

DNA methylation profiling has emerged as a powerful tool to distinguish biologically distinct tumor entities (14), and a recent analysis of peripheral nerve sheath tumors revealed several clinically important methylation subgroups (15). DNA methylation-based classification of the index tumor yielded high classifier scores for the subgroups “melanocytic schwannoma” and “regular schwannoma”, indicating a potential relationship with these entities. Unsupervised hierarchical clustering of the DNA methylation profiles of our cohort of 19 N/S HNSTs and 80 benign Schwann cell tumors from previously described methylation categories (15) identified a distinct

subcluster of tumors (Fig. 2, Subcluster A) that contained the majority of N/S HNSTs associated with sporadic schwannomatosis (7 of 9 tumors), including all ERBB2-mutant cases, whereas most N/S HNSTs associated with NF2 (5 of 7 tumors), which lack activating ERBB2 mutations, lay outside this cluster (Fig. 2, Cluster B).

Discussion

We here report for the first time that a substantial proportion of N/S HNSTs harbor ERBB2 kinase domain mutations that have been shown in other tumor entities to induce constitutive ERBB2 signaling, promote oncogenesis, and confer sensitivity to pharmacologic ERBB2 inhibition (12, 16). Furthermore, N/S HNSTs that had occurred in a patient with NF2 harbored a RET p.Tyr791Phe variant, which has been described as activating mutation in familial medullary thyroid carcinoma (17). Together, these findings indicate that N/S HNSTs are frequently driven by aberrant RTK signaling, and consistent with the hypothesis of a distinct molecular pathogenesis, we observed that ERBB2-mutant N/S HNSTs cluster in a separate subgroup of peripheral nerve sheath tumors based on genome-wide DNA methylation analysis (Fig. 2). The notion that these tumors might define a separate subgroup of N/S HNSTs is further supported by the mutual exclusivity of activating ERBB2 variants and mutations known to drive tumorigenesis in patients with schwannomatosis, NF2, and NF1 (Table 2).

In addition to providing new insights into underlying pathogenetic mechanisms, our findings may broaden the therapeutic armamentarium against N/S HNSTs. Pharmacologic treatment options for schwannomas or neurofibromas have been explored previously in the context of NF2 and NF1. In patients with NF2-associated vestibular schwannomas, both the dual EGFR/ERBB2 inhibitor lapatinib and the anti-VEGF antibody bevacizumab induced transient responses in a subset of tumors; in patients with NF1-associated plexiform neurofibromas, the MEK inhibitor selumetinib caused a decrease in tumor volume in 70% of cases (8, 9, 18, 19). The rationale to investigate the activity of lapatinib and selumetinib, respectively, was that loss of the Merlin tumor suppressor encoded by *NF2* is frequently associated with activation of EGFR or ERBB2 signaling (9), and that inactivation of the RAS GTPase-activating

protein encoded by *NF1* stimulates the RAS-MEK-ERK pathway (18). Our finding of recurrent ERBB2 kinase domain mutations now also suggests a targeted therapeutic strategy for N/S HNSTs occurring in non-NF patients. Importantly, the efficacy of ERBB2 kinase inhibition in ERBB2-mutant cancers varies across tumor entities and mutation types, with the greatest activity seen in breast, cervical, and biliary cancers and in tumors that contain missense variants in the catalytic domain (16). The prolonged response to lapatinib we observed in a patient with ERBB2-mutant N/S HNST is encouraging and supports prospective mutational profiling in N/S HNST patients with unresectable tumors as well as a molecularly guided clinical trial of ERBB2 inhibitors such as lapatinib, afatinib, or neratinib to establish mutant ERBB2 as valid therapeutic target in this disease and to identify modifiers of response.

Our findings may also have diagnostic implications as they indicate that the heterogeneous group of N/S HNSTs, which is currently defined based on histologic criteria alone (1), may be deciphered further using molecular analyses. ERBB2-mutant N/S HNSTs were exclusively associated with schwannomatosis, and future diagnostic algorithms may therefore incorporate the presence of an ERBB2 mutation as supportive or even core criterion (6, 10). Such additional parameters are urgently needed given that the differential diagnosis of tumor syndromes can be challenging due to overlap between entities, diverse mutations causing common phenotypes, and continuously evolving clinical criteria.

Finally, our results add to an emerging body of evidence indicating that oncogenic driver mutations can also be found in benign conditions and in premalignant lesions (20), and it will be interesting to study the impact of mutant ERBB2 signaling on the course of disease in N/S HNST patients and to identify individuals who may be candidates for preemptive therapeutic strategies.

Methods

Tumor Samples

For whole-exome and RNA sequencing of the index case, a fresh-frozen tumor specimen and matched peripheral blood mononuclear cells were collected. Samples were pseudonymized, and tumor histology and cellularity were assessed at the Institute of Pathology, Heidelberg University Hospital, prior to further processing.

For targeted next-generation sequencing, sections of 18 formalin-fixed paraffin-embedded (FFPE) tumors from 14 patients were provided by the Institute of Pathology and Neuropathology, University Hospital Tübingen, and the Institute of Neurology, University Hospital Frankfurt. These tumors were evaluated by a board-certified neuropathologist at the Institute of Pathology and Neuropathology, University Hospital Tübingen (JS), and original diagnoses were independently confirmed by another board-certified neuropathologist at the Institute of Neurology, University Hospital Frankfurt (PNH). Tumor tissue was microdissected to achieve a tumor cell content of at least 15%.

Study Approval

Tissue samples were used in accordance with the regulations of the respective tissue bank and after approval by the ethics committees of Heidelberg University (Ethikkommission of the Medical Faculty Heidelberg, Heidelberg, Germany) and Frankfurt University (Ethik-Kommission, University Hospital Frankfurt, Goethe University, Frankfurt, Germany) as well as the Neuroncology Biobank Board Tübingen (Tübingen, Germany) (protocol numbers S-206/2011, SNO-05-2017, and ZNO32, respectively). Written informed consent was obtained prior to the study. Research was conducted in accordance with the Declaration of Helsinki.

Whole-Exome and RNA Sequencing

DNA and RNA from the tumor specimen and DNA from the blood sample were isolated using the AllPrep DNA/RNA/Protein Mini Kit (Qiagen), followed by quality control and quantification using a Qubit 2.0 Fluorometer (Life Technologies), a 2200 TapeStation system (Agilent), and a 2100 Bioanalyzer system (Agilent).

Exome capturing was performed using SureSelect Human All Exon V5 in-solution capture reagents (Agilent). Briefly, 1.5 µg genomic DNA were fragmented to 150-200-base pair (bp; paired-end) insert size with a Covaris S2 device, and 250 ng of Illumina adapter-containing libraries were hybridized with exome baits at 65°C for 16 hours. Paired-end sequencing (101 bp) was carried out with a HiSeq 2500 instrument (Illumina) in rapid mode. RNA sequencing libraries were prepared using the TruSeq RNA Sample Preparation Kit v2 (Illumina). Briefly, mRNA was purified from 1 µg total RNA using oligo(dT) beads, poly(A)⁺ RNA was fragmented to 150 bp and converted into cDNA, and cDNA fragments were end-repaired, adenylated on the 3' end, adapter-ligated, and amplified with 12 cycles of PCR. The final libraries were validated using a Qubit 2.0 Fluorometer (Life Technologies) and a Bioanalyzer 2100 system (Agilent). Paired-end sequencing (2 x 101 bp) was carried out with a HiSeq 2500 instrument (Illumina) in rapid mode.

Exome sequencing reads were mapped to the 1000 Genomes Phase 2 assembly of the human reference genome (NCBI build 37.1) using BWA (version 0.6.2) with default parameters and maximum insert size set to 1,000 bp (21). BAM files were sorted with SAMtools (version 0.1.19) (22), and duplicates were marked with Picard tools (version 1.90). Average target coverage was 180x for the tumor and 160x for the control. For the detection of single-nucleotide variants (SNVs), we applied an in-house analysis pipeline based on SAMtools mpileup and bcftools with parameter adjustments to allow for calling of somatic variants with heuristic filtering

as previously described (23-25). After annotation with RefSeq (version September 2013) using ANNOVAR (26), somatic, non-silent coding variants of high confidence were selected. Short insertions and deletions (indels) were identified using Platypus (version 0.5.2; parameters genIndels=1, genSNPs=0, ploidy=2, nIndividuals=2) by providing the tumor and control BAM files (27). To be of high confidence, somatic calls (control genotype 0/0) were required to either have the Platypus filter flag PASS or pass custom filters allowing for low variant frequency using a scoring scheme. Candidates with the badReads flag, alleleBias or strandBias if the variant allele frequency was less than 10%, or more than 2 of the remaining flags were discarded. Additionally, combinations of Platypus non-PASS filter flags, bad quality values, low genotype quality, very low variant counts in the tumor, and presence of variant reads in the control were not tolerated. Indels were annotated with ANNOVAR, and somatic high-confidence indels falling into a coding sequence or splice site were extracted. Copy number variants were analyzed by read depth plots and an in-house pipeline using the VarScan2 copynumber and copyCaller modules (28). Regions were filtered for unmappable genomic stretches, merged by requiring at least 70 markers per called copy number event, and annotated with RefSeq genes using BEDTools (29).

RNA sequencing reads were mapped with STAR (version 2.3.0e) (30). For building the index, the 1000 Genomes reference sequence with GENCODE version 17 transcript annotations were used. For alignment, the following parameters were used: alignIntronMax 500000, alignMatesGapMax 500000, outSAMunmapped Within, outFilterMultimapNmax 1, outFilterMismatchNmax 3, outFilterMismatchNoverLmax 0.3, sjdbOverhang 50, chimSegmentMin 15, chimScoreMin 1, chimScoreJunctionNonGTAG 0, chimJunctionOverhangMin 15. The output was converted to sorted BAM files with SAMtools, and duplicates were marked with Picard tools (version 1.90).

Expression levels were determined per gene and sample as reads per kilobase (kb) of exon model per million mapped reads, and RefSeq was used as gene model. For each gene, overlapping annotated exons from all transcript variants were merged into non-redundant exon units with a custom Perl script. Nonduplicate reads with mapping quality above zero were counted for all exon units with coverageBed from the BEDtools package. Read counts were summarized per gene and divided by the combined length of its exon units (in kb), and the total number of reads (in millions) was counted by coverageBed. SNVs and indels were annotated with RNA information by generating a pileup of the DNA variant position in the RNA BAM file with SAMtools.

Hybrid Capture-Based Panel Sequencing

DNA was extracted using a Maxwell 16 Research System (Promega), followed by quantification using the QuBit 2.0 DNA High Sensitivity Kit (Thermo Fisher). Library preparation for the capture-based TruSight Oncology 500 panel (Illumina) was performed as described previously (31). DNA integrity was assessed using the Genomic DNA ScreenTape Analysis on a 4150 TapeStation System (Agilent). To fragment DNA to a length of 90-250 bp, 80 ng DNA were sheared for 50-78 seconds using an ME220 Focused-ultrasonicator (Covaris). Following target capture and purification steps, enriched libraries were amplified by 15 cycles of PCR and subsequently quality controlled using the KAPA SYBR Library Quantification Kit (Thermo Fisher) on a StepOnePlus qPCR system (Thermo Fisher). Libraries were sequenced on a NextSeq 500 instrument (Illumina) to a mean coverage of 1,096x using high-output cartridge and v2 chemistry. All assays were performed according to the manufacturers' protocols.

Processing of raw sequencing data and variant calling were carried out using the TruSight Oncology 500 Local App (version 1.3.0.39). Called variants were verified by visual inspection in the Integrative Genomics Viewer (32). Only variants with an allele frequency above 5% and a minimum coverage of greater than 100x were taken into account.

DNA Methylation Profiling and Unsupervised Hierarchical Clustering

Tumor DNA was analyzed using Infinium Human Methylation 450K BeadChip or Infinium MethylationEPIC BeadChip arrays (Illumina) at the DKFZ Genomics and Proteomics Core Facility. Data were processed with the R/Bioconductor package minfi (version 1.20). For unsupervised hierarchical clustering, the 25,000 probes with the highest median absolute deviation (MAD) across the beta values of 80 benign Schwann cell tumors (15) were selected. Samples were hierarchically clustered using Euclidean distance and Ward's linkage method. Filtering and genome-wide DNA copy number analyses were performed using the R/Bioconductor package conumee as described (33).

Immunohistochemistry

Immunohistochemical stainings were performed on FFPE whole-tissue slides using the following primary antibodies: S100 (polyclonal rabbit, 1:5000, Dako), Ki67 (monoclonal mouse, clone MIB-1, 1:200, Dako) and reticulin (Reticulum II Staining Kit, Ventana).

Sequencing Data Deposition

Sequencing data were deposited in the European Genome-phenome Archive under accession EGAS00001003776.

Author Contributions

Conception and design: MWR, PNH, JPS, DER, AS, SF. Development of methodology: MWR, PNH, MK, CH, BH, LG, CB, ES, WW, BB, AvD, JPS, DER, HG, AS, SF. Acquisition of data: MWR, PNH, MK, CH, VE, WW, DER, HG, AS, SF. Analysis and interpretation of data: MWR, PNH, MK, CH, BH, LG, JS, MUS, MT, GM, MW, V-FM, ES, WW, BB, AvD, MM, DER, HG, AS, SF. Writing and approval of the manuscript: MWR, AS, SF, and all other authors.

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Table 1. RTK mutations in the study cohort

Patient	Tumor No.	Tumor syndrome	Gene	Exon	Mutation (cDNA)	Mutation (protein)	COSMIC ID	Methylation Cluster
1	1	Schwannomatosis (index patient)	<i>ERBB2</i>	19	c.2305G>T	Asp769Tyr	COSM1251412	A
2	2	Schwannomatosis	<i>ERBB2</i>	19	c.2305G>T	Asp769Tyr	COSM1251412	A
3	3	Schwannomatosis	<i>ERBB2</i>	19	c.2264T>C	Leu755Ser	COSM14060	A
4	4	Schwannomatosis	<i>ERBB2</i>	20	c.2329G>T	Val777Leu	COSM14062	A
	5	Schwannomatosis	<i>ERBB2</i>	20	c.2329G>T	Val777Leu	COSM14062	A
	6	Schwannomatosis	<i>ERBB2</i>	20	c.2329G>T	Val777Leu	COSM14062	A
5	7	Schwannomatosis			None			A
	8	Schwannomatosis			None			A
6	9	Schwannomatosis			None			B
7	10	Schwannomatosis			None			A
8	11	NF2	<i>RET</i>	13	c.2372A>T	Tyr791Phe	COSM1159820	B
	12	NF2	<i>RET</i>	13	c.2372A>T	Tyr791Phe	COSM1159820	A
9	13	NF2			None			B
10	14	NF2			None			B
11	15	NF2			None			B
12	16	NF2			None			B
13	17	NF2			None			A
14	18	None			None			B
15	19	Schwannomatosis			None			B

COSMIC, Catalogue Of Somatic Mutations In Cancer (<https://cancer.sanger.ac.uk/cosmic>).

Table 2. NF1, NF2, SMARCB1 and LZTR1 mutations and chromosome 22q status in the study cohort

Patient	Tumor No.	Germline mutation	NF1 mutation (tumor)	NF2 mutation (tumor)	SMARCB1 mutation (tumor)	LZTR1 mutation (tumor)	Chromosome 22q status (tumor)
1	1	No NF1, NF2, SMARCB1, or LZTR1 mutation	None	None	None	None	Normal
2	2	NA	None	None	None	None	Normal
3	3	No NF1 or NF2 mutation	None	None	None	None	Normal
4	4	NA	None	None	None	None	Normal
	5	NA	None	None	None	None	Normal
	6	NA	None	None	None	None	Normal
5	7	No NF2 mutation	None	None	c.130_138del; p.Lys45_Tyr47 delLysArgTyr	None	Heterozygous deletion
	8	No NF2 mutation	None	c.955C>T; p.Gln319*	c.130_138del; p.Lys45_Tyr47 delLysArgTyr	None	Heterozygous deletion
6	9	No NF2 mutation	None	c.817_824del; p.Ile273fs*3	None	None	Heterozygous deletion
7	10	No NF2 mutation	None	c.623T>C; p.Leu208Pro	c.987_987del; p.Ser329fs*28	None	Heterozygous deletion
8	11	NA	None	None	c.1130G>A; p.Arg377His	None	Heterozygous deletion
	12	NA	None	None	None	None	Heterozygous deletion
9	13	No NF2 mutation	None	c.491delC; p.Gln165fs*9	None	None	Normal
10	14	NF2: c.1145-1149insT p.Leu384fs*42	None	c.1149delG; p.Leu384fs*42	None	c.614T>C; p.Ile205Thr	Heterozygous deletion
11	15	No NF2 mutation	None	c.561delA; p.Ile188fs*21	None	None	Normal
12	16	NF2: c.576C>G; p.Tyr192X	ND	ND	ND	ND	Heterozygous deletion
13	17	No NF2 mutation	None	c.1447-2A>C	None	None	Heterozygous deletion
14	18	NA	None	c.674delG p.spl?	c.1130G>A; p.Arg377His	None	Heterozygous deletion
15	19	No NF2 mutation	None	None	None	None	Heterozygous deletion

NA, not assessed. ND, not done.

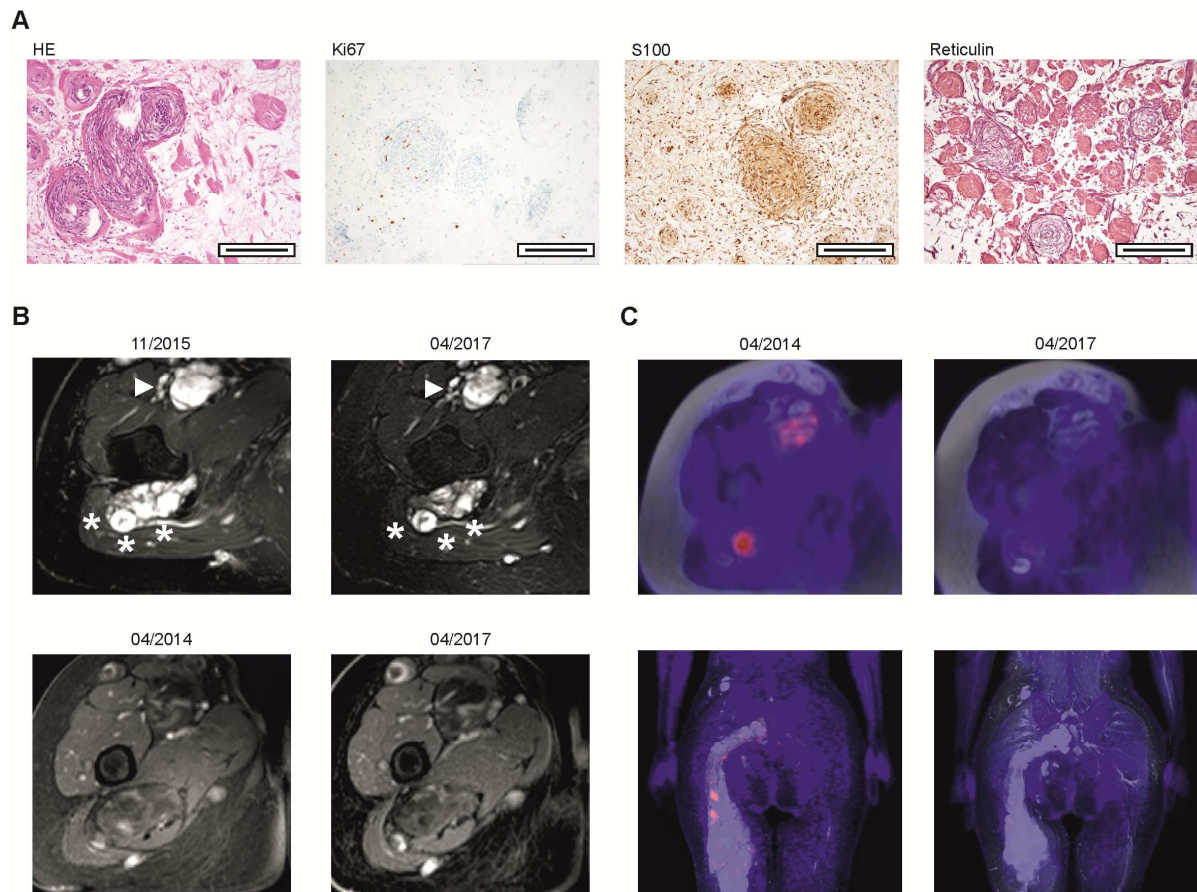


Figure 1. Histologic appearance, tumor burden, and assessment of response to lapatinib in a patient with N/S HNST.

A: HE, S100, reticulin, and Ki67 staining. Scale bar, 200 μm.

B: Axial MRI studies showing decreased tumor masses over time. Upper panel, T2-weighted and fat-suppressed images of hyperintense tumors along the right sciatic (asterisks) and femoral (arrowhead) nerves; lower panel, T1-weighted, contrast-enhanced, and fat-suppressed images of slightly and inhomogeneously contrast-enhancing tumors along the right sciatic and femoral nerves.

C: FDG-PET/MRI studies showing decreased tracer uptake over time. Upper row, axial images of tumors along the right sciatic and femoral nerves; lower row, coronal images of tumors along the right sciatic nerve.

Fig. 2

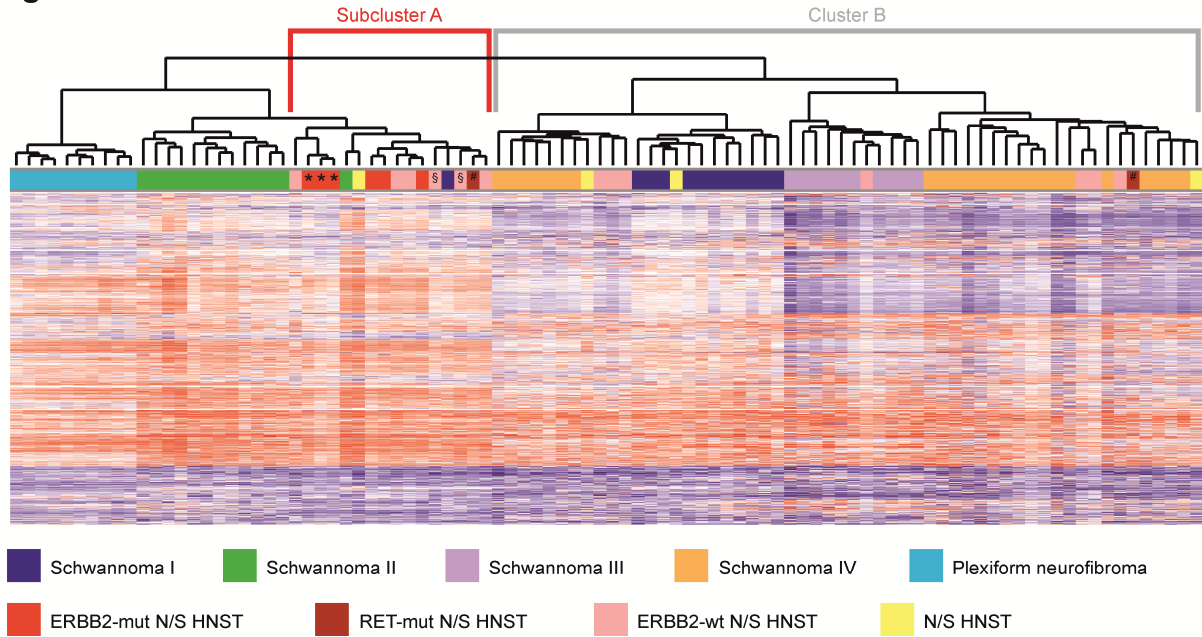


Figure 2. DNA methylation profiling of peripheral nerve sheath tumors.

Dendrogram showing the results of unsupervised hierarchical clustering of the DNA methylation profiles of N/S HNSTs and 80 benign Schwann cell tumors. “Schwannoma I-IV” refers to previously described methylation subgroups (15). DNA methylation levels (beta values) are represented as heatmap. mut, mutant; wt, wildtype. *, §, and # each identify tumors from the same patient.