Poroma is a benign skin tumor exhibiting terminal sweat gland duct differentiation. The present study aimed to explore the potential role of gene fusions in the tumorigenesis of poromas. RNA sequencing and reverse transcription PCR identified highly recurrent YAP1-MAML2 and YAP1-NUTM1 fusions in poromas (92/104 lesions, 88.5%) and their rare malignant counterpart, porocarcinomas (7/11 lesions, 63.6%). A WWTR1-NUTM1 fusion was identified in a single lesion of poroma. Fluorescence in situ hybridization confirmed genomic rearrangements involving these genetic loci. Immunohistochemical staining could readily identify the YAP1 fusion products as nuclear expression of the N-terminal portion of YAP1 with a lack of the C-terminal portion. YAPI and WWTR1, also known as YAP and TAZ, respectively, encode paralogous transcriptional activators of TEAD, which are negatively regulated by the Hippo signaling pathway. The YAP1 and WWTR1 fusions strongly transactivated a TEAD reporter and promoted anchorage-independent growth, confirming their tumorigenic roles. Our results demonstrate the frequent presence of transforming YAP1 fusions in poromas and porocarcinomas and suggest YAP1/TEAD-dependent transcription as a candidate therapeutic target against porocarcinoma.

Introduction

Poroma is a skin adnexal tumor exhibiting features of the terminal sweat gland duct (1, 2). It typically presents as a solitary sessile nodule on the soles and palms but may occur in any area where sweat glands are present. Histologically, poroma consists of monomorphic small basaloid cells and contains occasional ductal structures. Most commonly, the tumor is broadly connected to the epidermis and grows down into the dermis, forming broad anastomosing bands. Some lesions exhibit a spectrum of architectural variations that allows them to be classified into histological subtypes (Supplemental Figure 1) (2–6). Hidroacanthoma simplex represents an intraepidermal variant, whereas a dermal duct tumor is primarily confined to the dermis. Poroid hidradenoma plex represents an intraepidermal variant, whereas a dermal duct tumor is primarily confined to the dermis. Poroid hidradenoma paciniform is associated with benign poromas, a finding implying malignant transformation (2, 9, 11). Furthermore, porocarcinomas arising from longstanding poromas have been reported (9, 12). Apart from these, many porocarcinomas do not have a benign precursor component and thus may develop de novo (11, 13). A recent sequencing analysis of 5 porocarcinomas identified recurrent mutations in TP53, RB1, EGFR, and HRAS, but simultaneously revealed the molecular diversity of this rare type of tumor (8).

Given that mutations in known oncogenes and tumor suppressor genes are unlikely to underlie the tumorigenesis of a majority of poromas, we used RNA sequencing to explore the possible roles of gene fusions. Our analysis identified highly recurrent YAP1 fusions in poromas and porocarcinomas and demonstrated their transforming activity.

Results and Discussion

To probe the possible tumorigenic roles of gene fusions in poroma, we subjected 12 formalin-fixed, paraffin-embedded poroma specimens to RNA sequencing. The analysis identified YAP1-MAML2 and the reciprocal MAML2-YAP1 fusions in 4 lesions, YAP1-NUTM1 fusions in 3 lesions, and a WWTR1-NUTM1 fusion in 1 lesion (Supplemental Table 1). To expand on this finding, we examined
Interestingly, there appear to be associations between specific YAP1 fusion types and clinicopathological features. Hidroacanthoma simplex predominantly expressed the YAP1 exon 1-MAML2 fusion (4/5 lesions). YAP1-NUTM1 fusions were enriched in dermal duct tumors (2/2 lesions), poroid hidradenomas (4/6 lesions), porocarcinomas (6/11 lesions), poromas in the head and neck region (15/27 lesions), and poromas with necrotic foci (10/15 lesions). Because the number of each histological variant is rather limited in this study due to their rarity, further investigation based on a larger case series is desirable. However, it is notable that YAP1-NUTM1 fusions are enriched in porocarcinomas and poromas with primarily dermal localization, implying the link between the specific fusions and growth patterns. Examination of other skin tumors, including 24 squamous cell carcinomas, 32 basal cell carcinomas, 5 cutaneous adenocarcinomas, 9 Merkel cell carcinomas, and 27 seborrheic keratoses, did not identify any of the recurrent YAP1 fusions. These observations indicate that YAP1 fusions are highly recurrent and specific to poromas and porocarcinomas among skin neoplasms.

Considering that YAPI and MAML2 are on the opposite strands of adjacent genetic loci on chromosome 11p, the frequent coexpression of YAPI-MAML2 and MAML2-YAPI fusion transcripts is thought to be the consequence of intrachromosomal inversions (Supplemental Figure 3). Since NUTM1 and WWTR1 are on chromosome 15q14 and 3q25, respectively, the YAPI-NUTM1 and WWTR1-NUTM1 fusions are expected to result from interchromosomal translocations. Fluorescence in situ hybridization analysis revealed the YAPI locus rearrange-
portion expression in 100 poromas (96.2%) and 7 porocarcinomas (63.6%), including all the lesions with the YAP1 fusions (Figure 2 and Figure 3). The discrepant expression of the N- and C-terminal portions of YAP1 is consistent with the presence of YAP1-MAML2 or YAP1-NUTM1 fusion products. At the same time, it suggests the absence of MAML2-YAP1 fusion products. Seven poromas showed the discordant expression of N- and C-terminal regions of YAP1 in the absence of YAP1 fusion transcripts. We expect that these lesions might also have different types of YAP1 fusions, but the limitation in sample quality due to the use of formalin-fixed paraffin-embedded specimens precluded further investigation of other fusions in these lesions. The lesion with the WWTR1-NUTM1 fusion was also diffusely positive for the YAP1 N-terminus antibody, likely reflecting the known cross-reactivity of the antibody with the N-terminal portion of WWTR1 (16). Expression of NUTM1, also known as NUT, was observed in 25 poromas (24.0%) and 6 porocarcinomas (54.5%), including all lesions with the YAP1-NUTM1 fusions.

YAP1 immunohistochemistry on other skin tumors, including 23 squamous cell carcinomas, 24 basal cell carcinomas, 5 cutaneous adnexal carcinomas, 9 Merkel cell carcinomas, and 26 seborrheic keratoses, showed common expression of YAP1, in agreement with previous reports (17, 18). However, in contrast to poromas and porocarcinomas, the expression patterns of the N- and C-terminal regions of YAP1 were consistently identical in the respective tumors, consistent with the WT protein expression. Additionally, NUTM1 expression was never observed in skin tumors other than poromas and porocarcinomas. Thus, our immunohistochemical analysis could readily distinguish YAP1-MAML2 and YAP1-NUTM1 fusion products from the WT YAP1 based on the discordant expression of the N- and C-terminal regions of YAP1.

To examine the expression of fusion protein products in clinical specimens, we performed immunohistochemical staining using 2 antibodies against YAPI. The analysis revealed the diffuse nuclear portion expression in 100 poromas (96.2%) and 7 porocarcinomas (63.6%), including all the lesions with the YAPI fusions (Figure 2 and Figure 3). The discrepant expression of the N- and C-terminal portions of YAPI is consistent with the presence of YAPI-MAML2 or YAPI-NUTM1 fusion products. At the same time, it suggests the absence of MAML2-YAPI fusion products. Seven poromas showed the discordant expression of N- and C-terminal regions of YAPI in the absence of YAPI fusion transcripts. We expect that these lesions might also have different types of YAPI fusions, but the limitation in sample quality due to the use of formalin-fixed paraffin-embedded specimens precluded further investigation of other fusions in these lesions. The lesion with the WWTR1-NUTM1 fusion was also diffusely positive for the YAPI N-terminus antibody, likely reflecting the known cross-reactivity of the antibody with the N-terminal portion of WWTR1 (16). Expression of NUTM1, also known as NUT, was observed in 25 poromas (24.0%) and 6 porocarcinomas (54.5%), including all lesions with the YAPI-NUTM1 fusions.}

Although there were some differences in the prevalence of respective fusion types, YAPI fusions were frequently detected in both poromas and porocarcinomas. Therefore, we wanted to find out if any genetic alterations were involved in the malignant transformation of poroid neoplasms. To address this issue, we analyzed 23 poromas and 9 porocarcinomas using the NCC oncpanel test, a next-generation sequencing panel targeting 114 cancer-associated genes (15). The analysis identified a total of 49 protein-altering mutations (Supplemental Table 4 and Supplemental Table 5). Porocarcinomas tended to harbor a larger number of mutations than poromas (average 3.0 vs. 0.96 mutations/lesion). However, only 3 genes were recurrently mutated in porocarcinomas: KRAS, SETD2, and TP53. Each of these genes was mutated in only 2 of the lesions and SETD2 and TP53 mutations were also detected in poromas. These results indicate that mutations in major cancer-related genes cannot be used to distinguish benign and malignant poroid neoplasms.

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YAPI immunohistochemistry on other skin tumors, including 23 squamous cell carcinomas, 24 basal cell carcinomas, 5 cutaneous adnexal carcinomas, 9 Merkel cell carcinomas, and 26 seborrheic keratoses, showed common expression of YAPI, in agreement with previous reports (17, 18). However, in contrast to poromas and porocarcinomas, the expression patterns of the N- and C-terminal regions of YAPI were consistently identical in the respective tumors, consistent with the WT protein expression. Additionally, NUTM1 expression was never observed in skin tumors other than poromas and porocarcinomas. Thus, our immunohistochemical analysis could readily distinguish YAPI-MAML2 and YAPI-NUTM1 fusion products from the WT YAPI based on the discordant expression of the N- and C-terminal regions of YAPI. Additionally, NUTM1 expression was never observed in skin tumors other than poromas and porocarcinomas.
NUTM1-derived portion of YAP1-NUTM1 fusions also constitutes other tumor-related fusion proteins, including BRD4-NUTM1 in NUT midline carcinoma and CIC-NUTM1 in central nervous system Ewing sarcoma family tumor (28–30), and the recruitment of CBP and p300 is required for the oncogenic property of the BRD4-NUTM1 fusion (29). Thus, all the YAP1 and WWTR1 fusions detected in poromas and porocarcinomas harbor the N-terminal TEAD-binding domain of YAP1 or WWTR1 (31), and the MAML2- or NUTM1-derived regions that interact with transcriptional coactivators, CBP and p300 (27, 29).

Based on the common presence of domains to interact with transcription factors and transcriptional cofactors across these fusions, we postulated that these fusions function as a potent transcriptional activator of TEAD. In contrast, MAML2-YAP1 fusions lack these domains.

To test the functional significance of the YAP1 fusions, we first examined the intracellular localization of the YAP1 fusion products in HEK293T cells. Normally, active Hippo pathway signaling leads to the cytoplasmic retention and degradation of YAP1, inhibiting the formation of a complex with TEAD transcription factors (19, 20). WT YAP1 and a constitutively active mutant, YAP1S127A, were distributed in both the nucleus and cytoplasm, with a predominant localization in the cytoplasm (Figure 4B). In contrast, YAP1-MAML2 and YAP1-NUTM1 fusions exhibited exclusively nuclear localization with speckled and homogeneous staining patterns, respectively, suggesting that the 3′-fusion partners contribute to the nuclear localization of these fusions through distinct mechanisms. Next, we performed a reporter assay to test the role of the fusions in TEAD-dependent transcription. As expected, YAP1-MAML2 and YAP1-NUTM1 fusions strongly upregulated the expression of a TEAD reporter, and the reporter expression levels were even higher than those with 2 YAP1 mutants that are resistant to negative regu-
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**Figure 4. Functional significance of YAP1 and WWTR1 fusions.** (A) Structures of the putative fusion gene products expressed in poromas and porocarcinomas. Arrows indicate fusion breakpoints. Notice that YAP1-MAML2, YAP1-NUTM1, and WWTR1-NUTM1 fusion gene products harbor a TEAD-binding domain derived from YAP1 or WWTR1 and a p300-binding domain derived from MAML2 or NUTM1. (B) Intracellular localization of YAP1 fusion gene products. The FLAG-tagged proteins were detected by immunofluorescence staining with nuclear staining using DAPI. Scale bars: 10 μm. (C) TEAD luciferase reporter assays on HEK293T cells transfected with doxycycline-inducible vectors. YAP1 S127A and YAP1 S127A represent constitutively active mutants. The luciferase activity of EGFP-transfected cells in the absence of doxycycline was set at 1 to indicate relative luciferase activities. Data represent mean of triplicate measurements ± SD. (D) Soft agar colony formation assay of NIH3T3 cells expressing the fusion transgenes. NIH3T3 cells were transduced with doxycycline-inducible retrovirus vectors to express the respective transgenes and subjected to a soft agar colony formation assay in the presence of doxycycline. None of the clones formed colonies in the absence of doxycycline. Scale bar: 200 μm.

Next, we tested if YAPI and WWTR1 fusions also show transforming activity in epithelial cells by using an immortalized human dermal keratinocyte (HDK) cell line (33). A reporter assay was performed by using doxycycline-dependent inducible cell lines stably transfected with a TEAD reporter (Supplemental Figure 5A), because of the low transfection efficiency of the HDK cells. As

...tion by the Hippo pathway. In contrast, the MAML2-YAP1 fusion, MAML2, and NUTM1 did not significantly induce the TEAD reporter expression (Figure 4C).

We then established a series of doxycycline-dependent inducible NIH3T3 cell lines. Despite repeated attempts, we could not obtain sublines with high expression levels of YAP1-MAML2 fusions (Supplemental Figure 4). However, in a soft agar colony formation assay, the induction of YAP1-MAML2, YAP1-NUTM1, and WWTR1-NUTM1 expression resulted in anchorage-independent growth, similar to that seen in constitutively active YAP1 mutants (Figure 4D) (32). In contrast, the wild-type YAP1 and the MAML2-YAP1 fusion did not promote anchorage-independent growth. These observations indicate the role of YAP1 and WWTR1 fusions in the activation of TEAD transcription factors and their transforming activity. Although two-thirds of YAP1-MAML2 fusions were associated with the reciprocal MAML2-YAP1 fusions, functional assays showed the lack of their transforming activity, and immunohistochemical study failed to detect their protein products. Thus, MAML2-YAP1 fusions are likely nonfunctional by-products of YAP1-MAML2 fusions, resulting from intrachromosomal inversion.

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observed with NIH3T3 cells, expression of YAPI-MAML2, YAPI-NUTM1, and WWTR1-NUTM1 fusions strongly upregulated the expression of the TEAD reporter (Supplemental Figure 5B). A soft agar colony formation assay showed enhanced anchorage-independent growth upon the induction of YAPI-MAML2, YAPI-NUTM1, and WWTR1-NUTM1 (Supplemental Figure 5C). However, these colonies were smaller than those formed by cells expressing constitutively active YAPI and WWTR1 mutants. In addition, cells overexpressing WT YAPI and WWTR1 also formed small colonies. These findings indicate that YAPI and WWTR1 fusions activate TEAD transcription factors and also promote anchorage-independent growth in epithelial cells. However, they may exhibit somewhat different biological effects in different cell types.

The present study identified highly recurrent YAPI and WWTR1 fusions in poromas and porocarcinomas. The YAPI and WWTR1 fusions are potent transcriptional activators of TEAD and have transforming activity. The respective fusion types are likely associated with several clinicopathological features, including the enrichment of YAPI-NUTM1 fusions in porocarcinomas. Since YAPI fusions are specific to poromas and porocarcinomas among skin tumors, the detection of their protein products would have immediate diagnostic applications. Furthermore, the identification of recurrent YAPI fusions in porocarcinomas suggests YAPI/TEAD-dependent transcription as a potential therapeutic target against this rare malignancy.

Methods

Additional information is provided in Supplemental Methods.

Study approval. The study was approved by the Ethical Committee of National Cancer Center, Tokyo, Japan (approval 2017-078).

Author contributions

SS, TK, and TM contributed equally to this work. SS, TK, and TM designed the study. SS, ER, RO, HI, TM performed next-generation and Sanger sequencing. SS, SW, and TM performed histological, immunohistochemical, and FISH analyses. TK performed functional assays. KS, SA, KT, MI, YS, and NY provided tissue samples and clinical information. NG provided cDNA clones. SS wrote the manuscript with support from TK and TM.

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