HIF2A gain-of-function mutations detected in duodenal gangliocytic paraganglioma

Dear Editor,

Somatic hypoxia-inducible factor 2 alpha (*HIF2A*) mutations are responsible for a newly discovered syndrome of multiple paragangliomas (PGL) and duodenal somatostatinomas associated with polycythemia (Pacak–Zhuang syndrome) (Zhuang et al. 2012, Pacak et al. 2013, Toyoda et al. 2014). In this syndrome, found exclusively in females, somatostatinomas are located in the 2nd portion of the duodenum.

Gangliocytic PGL (GPGL), a mixed neuroectodermal–endodermal tumor, is a rare and unique type of PGL that is almost exclusively located in the 2nd portion of the duodenum, much like *HIF2A*-related somatostatinomas. Both GPGLs and *HIF2A*-related somatostatinomas produce and/or secrete somatostatin. Clinically, GPGL patients often present with gastrointestinal bleeding, iron deficiency anemia, abdominal pain, and weight loss. Despite their clinical presentation, signs and symptoms of catecholamine excess are not present, because these tumors do not produce catecholamines due to the absence of tyrosine hydroxylase, a rate-limiting enzyme in catecholamine biosynthesis. Until now, no familial cases of GPGL have been described.

Gain-of-function somatic *HIF2A* mutations in patients (with this syndrome or without it) are located close to the oxygen-dependent degradation domain (ODD) of HIF2 α (Zhuang et al. 2012, Comino-Mendez et al. 2013, Pacak et al. 2013, Toledo et al. 2013). The mutated HIF2 α protein prevents proper hydroxylation by prolyl hydroxylases, resulting in abnormally low binding to the von Hippel–Lindau (VHL) protein and enhances HIF2 α stabilization and activation. Consequently, the HIF2 α downstream genes, including the cancer signaling pathway genes, are upregulated, contributing to the pathogenesis of cancer. This notion was nicely presented in a study of Toledo et al., where HIF2 α mutants in HEK293, renal cell carcinoma 786-0, or rat pheochromocytoma PC12 cell lines, showed increased stability, resistance to

VHL-mediated degradation, target induction, and reduced chromaffin cell differentiation (Toledo et al. 2013). Moreover, mice injected with mutant *HIF2A* tumor cells developed tumors with a shorter latency than tumors in mice injected with wildtype (WT) *HIF2A* tumor cells. As concluded by the authors, these results further support a direct oncogenic role for somatic gain-of-function *HIF2A* mutations in human neoplasia and strengthen the link between hypoxic pathways and PGL.

We tested whether duodenal GPGLs share a similar pathogenic mechanism with PGLs associated with somatic *HIF2A* mutations. Ten GPGL tissues (Fig. 1A–D) were screened for somatic *HIF2A* mutations; patient 1 was also tested for a potential germline *HIF2A* mutation, which included blood leukocytes and gallbladder tissue. Genomic DNA was extracted from paraffin-embedded tissue and white blood cells using a DNeasy Blood & Tissue Kit (Qiagen). Two female patients were found to have pathogenic somatic *HIF2A* mutations in their GPGLs. No *HIF2A* mutation was found in the blood or gallbladder tissue of patient 1. In patient 1 (a 54-year-old woman), a periampullar GPGL (Fig. 1E) was found to have a C $\rightarrow$ T substitution at base 1556 in exon 12 of *HIF2A*. In patient 2 (a 46-year-old woman), a periampullar GPGL (Fig. 1E) exhibited a C $\rightarrow$ T substitution at base 1630 of the same exon. The mutations resulted in amino acid substitutions in HIF2 α protein to T519M and P544S, respectively. Alignment of multiple HIF2 α peptide sequences indicated that both amino acids are located in the proximal region of the primary hydroxylation site of HIF2 α protein and are evolutionary conserved across different species (Fig. 1F). Both mutations were closely located to the oxygen-sensing domain of HIF2 α , which affects its protein ubiquitination in the presence of oxygen.

To understand the functional impact of either mutation, we obtained a pcDNA3-HA-HIF2 α plasmid containing human *HIF2A* coding sequence (Addgene plasmid 18950) and introduced T519M and P544S mutations through Quikchange Lightning Site-directed

Figure 1

Histopathological features of GPGLs from patient 1 (A) and patient 2 (B). In both tumors, the neoplastic cells were arranged in solid nests, trabeculae, and fascicles with areas of epithelioid (arrow A), ganglion-like (arrow B), and spindle (arrow C) cells. Hematoxylin & eosin stain, magnification $\times 20$. Positive immunohistochemical stains for chromogranin A (C) and somatostatin (D) in the tumor from patient 1, magnification $\times 20$. Both GPGLs were negative for tyrosine hydroxylase (ImmunoStar) immunohistochemistry (data not shown). (E) Genomic DNA sequencing of *HIF2A* exon 12 nucleotides 1551–1561. Heterozygous T519M (patient 1) and P544S (patient 2) somatic mutations were identified in both GPGLs. No *HIF2A* mutations were found in the blood or gallbladder tissue of patient 1. (F) Alignment of amino acid sequence of HIF2 α residues 1625–1635 in humans, mice, rats, ox, chickens, and frogs.

Mutagenesis Kit (Agilent). The plasmids were introduced into HEK293 cells with Lipofectamine 2000 (Invitrogen) for detection of HIF2 α ubiquitination and measurement of protein half-lives.

Through an immunoprecipitation assay, we found that the ubiquitination of HIF2 α was reduced in either the T519M or P544S HIF2 α variant compared with WT protein, with a reduction of 35.16 and 61.76%, respectively (Fig. 2A). We confirmed the changes in protein stability of the mutant HIF2 α via a cycloheximide (CHX) chase assay. WT HIF2 α protein exhibited a half-life of 7.60 min, suggesting rapid turnover of HIF2 α protein under normoxic conditions. By contrast, T519M and P544S variants were more stable, with half-lives extended to 67.21 and 81.09 min, respectively, when compared with WT protein with a half-life of 7.6 min (Fig. 2B and C). Furthermore, the expression of canonical hypoxia-related genes in tumors was measured through quantitative real-time PCR (Fig. 2D). We identified the upregulation of genes downstream of HIF2 α , including *EDN1*, *EPO*, *GLUT1*, *GNA14*, *LDHA*, and *VEGFA* measured through a real-time PCR assay on a ViiA 7 real-time PCR system (Applied Biosystems).

Until now, most GPGLs were considered sporadic, although some of them are found rarely in association with neurofibromatosis type 1. This is the first report

that shows that somatic gain-of-function *HIF2A* mutations are present in 20% of GPGLs in the present series. The mutations appear to be located in the hot spot of the oxygen-sensing domain of HIF2 α , resulting in increased HIF2 α stabilization and impaired ubiquitination and degradation, as also described in studies by Dahia *et al.* (2005), Toledo *et al.* (2013), and Comino-Mendez *et al.* (2013). Increased half-life of HIF2 α and enhanced activity trigger its downstream regulated genes and thus, the upregulation of the HIF signaling pathway, classifying these tumors as cluster 1. The evidence of *HIF2A* oncogenicity of PGLs was previously demonstrated by Dahia *et al.* (2005). The authors showed that *HIF2A* mutations conferred growth advantage (e.g., by resistance to VHL-mediated degradation, increased tumor cell stability, and reduced PC12 cell differentiation) in a mouse subcutaneously injected with various mutant *HIF2A* tumor cells.

The developmental origin of GPGL is currently unclear, but it is believed that this tumor derives from a defective sheet of intraembryonic endoderm progenitor cells (primordial gut; epithelial cells in GPGL), rests that have recruited progenitor cells of neuroectoderm, nerves/Schwann and ganglion cells (spindle cells and ganglion-like cells, respectively in GPGL), and smooth muscle (Perrone *et al.* 1985, Witzigmann *et al.*

Figure 2

(A) Immunoprecipitation of HIF2α ubiquitination. Recombinant HIF2α protein was immunoprecipitated with an antibody against HA-tag. Protein ubiquitination was probed with a polyclonal antibody against ubiquitin. (B) CHX chase assay measurement of HIF2α protein stability. (C) Quantification of HIF2α protein stability on the basis of data shown in B. (D) Quantitative mRNA expression measurement of four hypoxia-related genes in tumor specimen from patient 1: *EDN1*, *EPO*, *GLUT1*, *GNA14*, *LDHA*, and *VEGFA* – performed by quantitative PCR assay. A normal adrenal medulla (NM) specimen was used as a control.

2002). GPGLs are often positive for somatostatin, and both tumors are initially derived from endoderm, which is capable of recruiting neuroendocrine cells. Since somatic *HIF2A* mutations are found in both tumor types, we hypothesize that these mutations with HIF signaling pathway upregulation may affect common precursor cells and their differentiation to somatostatin-secreting endocrine cells. Therefore, some PGLs, somatostatinomas, and GPGLs share similarities in their pathogenic mechanisms.

Amino acid substitutions in the ODD domain have been linked to abnormalities in oxygen-dependent hydroxylation of HIF2α, as well as abnormal accumulation of HIF2α protein under a normal oxygen level. Indeed, similar to previous findings related to HIF2α stabilization and degradation, both mutations resulted in significant increased stability (half-lives extended to 67.21 and 81.09 min for T519M or P544S, respectively) and decreased protein ubiquitination (reduction of 35.16 and 61.76% for T519M or P544S, respectively). The results confirm that HIF2α mutant protein found in both patients escaped from the degradation mechanism and likely induced a pseudohypoxic phenotype, as well as oncogenic gene transcription, through the canonical hypoxia pathway.

Moreover, mutated *HIF2A* overexpression could contribute to a more immature phenotype of GPGL since *HIF2A* knockdown in neuroblastomas, a tumor with

similar developmental, biochemical, and localization characteristics as a PGL, promotes a sympathetic neuronal differentiation (Pietras et al. 2009). By contrast, high normoxic and hypoxic levels of HIF2α protein in neuroblastoma cells are associated with immature recruitment of endothelial cells as well as a neural crest-like phenotype, all which resemble GPGL as an immature tumor (Pietras et al. 2009).

Whether *HIF2A* mutations may be associated with the presence of other (neuro)endocrine tumors or health-related abnormalities, especially in tumors that are found in the 2nd portion of the duodenum apart from those associated with *MEN1*, *NF1*, and *HIF2A*-related somatostatinomas, is currently unclear. If a *HIF2A* mutation is found, patients may be considered to have personalized clinical and therapeutic management with a HIF2α-targeted drug as for other hereditary neuroendocrine tumors undergoing various genetic screening, including next-generation sequencing (Neumann & Eng 2009).

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Declaration of interest

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