

DATA REPORT

Somatic mosaicism of *EPAS1* mutations in the syndrome of paraganglioma and somatostatinoma associated with polycythemiaChunzhang Yang^{1,5}, Christopher S Hong¹, Josef T Prchal², Melina T Balint³, Karel Pacak^{4,5} and Zhengping Zhuang^{1,5}

We recently described a novel, non-inherited syndrome of tumor-specific mutations of hypoxia-inducible factor 2 α , encoded by *EPAS1*, leading to formation of multiple paragangliomas and somatostatinomas in the setting of congenital polycythemia. Although we had suspected that somatic mosaicism of *EPAS1* mutations was the underlying cause of tumorigenesis, we could not validate this theory in our initial findings. In this report, we developed a sensitive, peptide nucleic acid sequencing assay to uncover the presence of *EPAS1* mutations in blood and other somatic tissues of the two patients who were described in the initial characterization of this syndrome. As such, the current study demonstrates that the underlying pathogenesis of the syndrome of multiple paraganglioma and somatostatinoma formation with congenital polycythemia is somatic mosaicism of *EPAS1* mutations.

Human Genome Variation (2015) 2, 15053; doi:10.1038/hgv.2015.53; published online 10 December 2015

We previously reported a new syndrome, characterized by tumor-specific gain-of-function mutations of hypoxia-inducible factor 2 α (encoded by *EPAS1*), leading to the development of multiple paragangliomas and duodenal somatostatinomas in the setting of congenital secondary polycythemia.^{1–3} We found that all *EPAS1* mutations occurred near the primary hydroxylation site of hypoxia-inducible factor 2 α , resulting in augmented *EPAS1* transcriptional activity leading to increased erythropoietin (EPO) and polycythemia. We had hypothesized that a somatic *EPAS1* mutation may have occurred early in embryogenesis, leading to tissue-restricted somatic mosaicism (such as in neural crest cells) as this mutation was not detected in blood, nail, urine, buccal mucosa or skin DNA. This conclusion of ruling out germ-line mutations was further strengthened by the fact that the *EPAS1* mutation was not detected in the parents and other relatives of the affected patients. As such, the presence of *EPAS1* mutations in the tissues with mosaicism was postulated to render affected cells susceptible to tumorigenesis and retinal abnormalities,⁴ leading to simultaneous manifestation of separate and functionally distinct tumors and other manifestations. In a separate study, we did not detect such mutations in normal tissues from four additional patients.² Furthermore, more recently *EPAS1* mutations were found in DNA from blood leukocytes and buccal mucosa in one of two patients affected with paraganglioma, somatostatinoma and polycythemia, suggesting this syndrome may be secondary to somatic mosaicism that has variable tissue distribution.⁵ We also suspected that at least in some tissues a proportion of *EPAS1*-mutated non-tumor cells existed but was too minute to be detected by routine Sanger sequencing. To further investigate this possibility, we developed a peptide nucleic acid (PNA, PNA Bio, Thousand Oaks, CA, USA) assay designed to hybridize with the wild-type *EPAS1* sequence at codons 528–532 (Supplementary

Appendix)—the sites of tumor mutations in our initially reported patients. PNAs have been previously used as an effective PCR clamp to detect somatic mosaicisms of *GNAS* mutations in blood DNA of patients with McCune–Albright syndrome.⁶ Using a combined PNA and PCR assay, we detected mutations in the blood DNA of two patients, identical to those in the tumors of the patients we had described in our initial report (Figure 1a and Supplementary Figure S1 in the Supplementary Appendix).¹ Additional testing of DNA extracted from hair, nail and saliva from patient 1 revealed the same mutation present in the tumor tissue (Supplementary Figure S2) but not in the same tissues of patient 2. We quantified the degree of heterozygosity present in the blood of our patients through TA cloning, detecting 10 mutants out of 90 clones in patient 1 and 1 mutant out of 72 clones in patient 2. These data suggested that 22.2% of circulating cells in patient 1 and 2.8% of circulating cells in patient 2 harbored *EPAS1* mutations, respectively. Using serially titrated combined PNA and PCR assay experiments, we demonstrated that the PNA could detect as low as 2.5% of *EPAS1*-mutated cells (Figure 1b). Detailed materials and methods may be found in the Supplementary Appendix.

The data presented in this report demonstrate that the syndrome of paraganglioma and somatostatinoma associated with polycythemia occurs secondary to somatic mosaicism of *EPAS1* mutations, affecting cells in different tissues. Although we initially hypothesized that polycythemia predominantly occurred secondary to increased erythropoietin (EPO) production by *EPAS1*-mutated tumor cells, we now acknowledge that it is more likely either *EPAS1*-mutated mosaic non-tumor neural crest cells in the adrenal medulla, gastrointestinal tract or other tissues (a non-EPO producing tissue type) or *EPAS1*-mutated mosaic cells in tissues naturally producing EPO are responsible for polycythemia, especially before tumors develop. However, we currently cannot

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Received 22 July 2015; revised 27 August 2015; accepted 17 September 2015

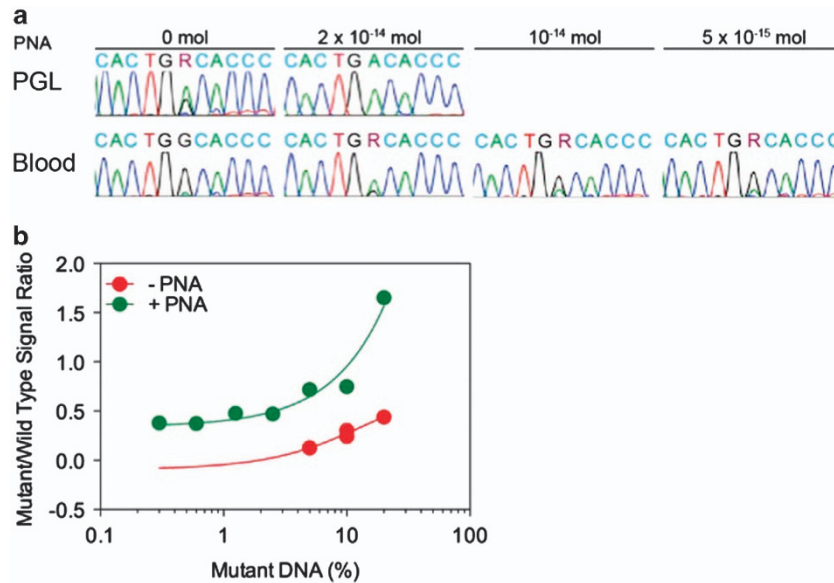


Figure 1. *EPAS1* sequencing of tumor and blood. (a) shows standard Sanger sequencing results, demonstrating a c.G1588A (p.A530T) mutation in DNA derived from paraganglioma (PGL; upper left sequence) but not blood leukocytes (lower left sequence) from patient 1. Addition of 2×10^{-14} mol peptide nucleic acid (PNA) resulted in amplification of the mutant allele signal in PGL DNA and uncovered the mutation in blood DNA. The mutant allele signal in blood DNA was still detectable using 50% and 75% diluted concentrations of the PNA. (b) illustrates results from serially titrated PCR reactions of TA clone-derived wild type and *EPAS1*-mutated plasmids with and without the PNA. Addition of the PNA allowed for detection of as low as 1.25% mutant plasmids, correlating with 2.5% mutant cells.

directly test this hypothesis because of lack of access to normal adrenal, renal and hepatic tissues. Given our two patients developed polycythemia shortly after birth, long before their tumor diagnoses, it is also plausible to suspect that *EPAS1* mutations may have involved hematopoietic precursor cells. Further testing of bone marrow-derived hematopoietic stem cells may answer this question.

In summary, we have demonstrated the presence of somatic mosaicism of *EPAS1* mutation-bearing cells in our initially described syndromic patients. We propose that *EPAS1* mutations occur early in embryogenesis, affecting hematopoietic and other tissue precursors and giving rise to early onset polycythemia, as well as affecting neuroendocrine cells, thus predisposing patients to paraganglioma and somatostatinoma formation later in life. Importantly, somatic mosaicism of *EPAS1* mutations should be considered in patients presenting with congenital polycythemia for whom conventional testing cannot detect *EPAS1* mutations. Application of the PNA assay described in this correspondence may be of clinical utility for detecting *EPAS1* mutations in these patients. In addition, further work may uncover *EPAS1* mutations in other tissues of affected patients, potentially predisposing to additional tumor types.

HGV DATABASE

The relevant data from this Data Report are hosted at the *Human Genome Variation Database* at <http://dx.doi.org/10.6084/m9.figshare.hgv.762>, <http://dx.doi.org/10.6084/m9.figshare.hgv.765>.

ACKNOWLEDGEMENTS

This study was funded by the Intramural Research Program of the National Institutes of Health (Bethesda, MD, USA).

COMPETING INTERESTS

The authors declare no conflict of interest.

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Supplementary Information for this article can be found on the *Human Genome Variation* website (<http://www.nature.com/hgv>).