USP8 mutation in Cushing’s disease

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Pituitary corticotroph adenomas, also referred to as Cushing’s disease (CD), secrete large amounts of adrenocorticotropic hormone (ACTH), resulting in excess glucocorticoids and hypercortisolism [1]. The diagnosis of hypercortisolism is complex and sometimes difficult because its clinical features overlap with other common diseases. Currently, 65-90% of patients achieve complete or partial remission after initial transsphenoidal surgery. However, surgical clues are sometimes hindered by failure to locate the tumor. Moreover, it is particularly difficult to effectively treat patients with residual cortisol excess due to incomplete removal of tumor tissues or recurrent tumors. The genetic pathogenesis of CD remains obscure, hampering the development of novel diagnostic and therapeutic strategies for this disease.

Fortunately, two recent studies demonstrated that the ubiquitin-specific protease 8 (USP8) gene is frequently mutated in corticotroph adenomas [2,3]. One of these studies identified USP8 mutation in 6 of 17 corticotroph adenomas [2], while the other showed that USP8 was mutated in 75 of 120 corticotroph adenomas [3]. The later study also reported that USP8 mutation was undetected in other pituitary adenoma types, including somatotroph adenomas and prolactin-secreting adenomas. Mutations of guanine nucleotide-binding protein subunit (GNAS) and the Aryl Hydrocarbon receptor interacting protein (AIP) gene are preferentially associated with somatotroph and prolactin-secreting adenomas [4]. These facts support the notion that the genetic origin of pituitary adenomas is heterogeneous and corticotroph adenoma is pathogenically distinct. Moreover, USP8 mutations are rarely found in cancers (1%) based on the COMSINC and TCGA database. Collectively, USP8 mutation is a common and specific genetic alteration in corticotroph adenoma, providing the novel insight into the molecular pathogenesis of this disease. Testing the status of USP8 mutation may be a useful strategy for the diagnosis of CD.

All identified USP8 mutations are located within or adjacent to the 14-3-3 binding motif (RSYSS) of the USP8 protein. USP8 is a deubiquitinase (DUB) and can be phosphorylated in its RSYSS motif, leading to association with 14-3-3 protein and subsequent impairment of its DUB activity [5]. Three identified dominant USP8 mutants failed to bind to 14-3-3 protein and displayed elevated DUB activity through testing the epidermal growth factor receptor (EGFR) as substrate. Furthermore, these USP8 mutations facilitate proteolytic cleavage of USP8 into two fragments, one of which processes higher DUB activity compared to full-length USP8.

In response to EGF, EGFR is activated to trigger multiple downstream signaling pathways including MAPK, and subsequently undergoes polyubiquitination for lysosomal degradation to switch down EGFR signaling. The identified USP8 mutants increase EGFR deubiquitination to inhibit EGF-induced EGFR down-regulation, leading to augmented and more sustained EGFR signaling (Figure 1). This is supported by this fact that corticotroph adenomas harboring USP8 mutation display a higher incidence of EGFR expression and protein abundance as well as phosphorylated Erk1/2.

USP8 mutations contribute to ACTH overproduction in CD. USP8-mutated corticotroph adenomas had higher ACTH production and expressed more POMC (encoding the precursor of ACTH) mRNA than those with wild type USP8. USP8 knockdown using shRNA in primary corticotroph adenoma cells effectively reduced ACTH production. Ectopic expression of USP8 mutants or cleaved USP8 in murine corticotroph cell line induced higher POMC promoter activity and transcription than wild-type USP8. Mechanistically, USP8 mutation

Figure 1: USP8 mutation contributes to ACTH overproduction. USP8 deubiquitinates EGFR and protects it from lysosomal degradation. USP8 mutants lose the binding to 14-3-3 protein, resulting in the increased DUB activity and more sustained EGFR-Erk1/2 signaling. Erk1/2 activation promotes POMC transcription probably through Nur77.
potentiates EGFR-Erk1/2 signaling, probably resulting in the activation of Nur77, a key transcription activator of the POMC gene (Figure 1) [6]. Among numerous USP8 substrates, EGFR appears to be the pivotal mediator for ACTH overproduction in USP8-mutated CD. In the absence of EGFR, USP8 was unable to augment the POMC promoter activity in luciferase reporter assay, and gefitinib, an EGFR inhibitor, significantly impaired the ACTH production in primary corticotroph adenoma cells [3,7]. Besides EGFR, USP8 may regulate other receptor tyrosine kinases and additional pathways which are expressed in corticotroph cells and control ACTH production and secretion. The identification of these USP8 targets will be helpful in understanding the mechanism of USP8 action in CD and discovering novel therapeutic targets. Although the proliferation action of USP8 mutant is similar to that of wild-type USP8 in corticotroph-derived cell line, it is unclear whether this was the case in vivo [2]. USP8 mutation is expected to have multiple effects other than ACTH hypersecretion and may be important for corticotroph tumorigenesis, probably through accelerating the cell cycle inhibitor p27(Kip1) degradation, resulting from constitutive activation of EGFR-Erk1/2. Mice deficient in p27(Kip1) have been reported to develop ACTH-secreting pituitary adenomas. Mice with the desired USP8 mutation in the 14-3-3 binding motif are urgent to be developed to dissect the exact roles of USP8 mutation in the pathogenesis of corticotroph adenoma.

Surgery alone is not effective in treatment of most CD cases. Optimum treatment of patients with the recurrent tumor and/or residual tumor tissues requires the development of new drugs. Inhibiting the DUB activity of USP8 seems to be a promising therapeutic strategy for USP8-mutated cases. Further efforts are needed to develop the specific USP8 inhibitor for treatment of CD. Considering that EGFR plays a critical role in mediating USP8 action in corticotroph adenomas, treatment of Gefitinib seems to be a safe and effective in patients harboring USP8-mutated adenoma.

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