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## **ACCEPTED MANUSCRIPT**

Caveolin-1 promotes pituitary adenoma cells migration and invasion by regulating the

interaction between EGR1 and KLF5

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Abstract

Caveolin-1 (Cav-1) is a principal structural protein of caveolae. Cav-1 has been

implicated in cancer progression, but its precise functional roles in pituitary adenoma

cells remain largely unclear. In this study, we evidenced that the level of cav-1 was

elevated in the invasive pituitary adenoma. Cav-1 knockdown restrained the migration

and invasion of pituitary adenoma cells. In cav-1-depleting cells, the expression of

miR-145, miR-124 and miR-183 were up-regulated. Further investigation showed that

cav-1 knockdown inhibited the nuclear translocation of EGR1, reducing the

interaction between EGR1 and KLF5. The resulting free KLF5 promoted the

expression of miR-145, miR-124 and miR-183 by binding to their promoters, which

was blocked by EGR1. Luciferase reporter assay indicated that miR-145 targeted

FSCN1, miR-124 targeted PTTG1IP, and miR-183 targeted EZR in pituitary adenoma

cells, respectively. Knockdown of FSCN1, PTTG1IP or EZR suppressed the

migration and invasion of pituitary adenoma cells. In conclusion, our data suggested

that the elevated cav-1 promoted pituitary adenoma cells migration and invasion by

regulating the interaction between EGR1 and KLF5.

Keywords: Caveolin-1; EGR1; KLF5; microRNA; Migration and invasion; Pituitary

adenoma

Introduction

Pituitary adenomas are commonly benign monoclonal neoplasms, accounting for

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