

[1]罗爽,王继东,何佳,等.apelin对大鼠卵巢颗粒增殖与凋亡的影响[J].第三军医大学学报,2013,35(16):1713-1716.

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## apelin对大鼠卵巢颗粒增殖与凋亡的影响(PDF) 分享到

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**Title:** Apelin promotes proliferation and suppresses apoptosis in rat ovarian granulosa cells *via* PI3K/Akt signaling pathway

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**关键词:** [ape1in](#); [APJ](#); [卵巢颗粒细胞](#); [PI3K/Akt](#)

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**摘要:** **目的** 探讨apelin对大鼠卵巢颗粒细胞增殖与凋亡的影响。 **方法** 用apelin  $10^{-8}$  mol/L培育SD大鼠卵巢原代颗粒细胞,小RNA干扰技术(siRNA)抑制颗粒细胞APJ表达,联合PI3K/Akt信号转导阻断剂LY294002和HIMO干预,MTT法及流式细胞术观察细胞增殖与凋亡情况;Western blot检测细胞凋亡相关信号蛋白表达。 **结果** MTT检测APJ-siRNA组、LY294002组、HIMO组细胞增殖率,与对照组比较差异均有统计学意义( $P<0.05$ );与对照组比较,APJ-siRNA组、LY294002组、HIMO组颗粒细胞Bad、Bax、Foxo3a蛋白表达明显上调( $P<0.05$ );与对照组比较,APJ-siRNA组、LY294002组、HIMO组Bc1-2蛋白表达明显下调( $P<0.05$ )。 **结论** ape1in通过APJ/PI3K/Akt信号通路促颗粒细胞增殖抗其凋亡。

**Abstract:** **Objective** To determine the effect of apelin on the proliferation and apoptosis in rat ovarian granulosa cells (GCs) *in vitro*. **Methods** SD rat ovarian GCs were isolated and primary cultured, and then cultured with apelin at a concentration of  $10^{-8}$  mol/L for 12, 24 and 48 h respectively, and then tested by MTT assay and flow cytometry (Annexin/PI). RNA interference (RNAi) was used to down-regulate the expression of APJ in GCs. PI3K/Akt inhibitor LY294002 and HIMO were used to determine the role of the signaling pathway. Western blotting was employed to determine the expression of cell apoptosis-related proteins. **Results** Apelin of  $10^{-8}$  mol/L promoted the proliferation and anti-apoptosis in GCs ( $P<0.05$ ). Apelin activated the phosphorylation of Akt, while APJ-

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siRNA, inhibitor LY294002 and HIMO abolished apelin-induced activation of Akt. Western blotting showed that apelin enhanced the expression of Bcl-2 but suppressed the expression of Bax, Bad, and Foxo3a ( $P<0.05$ ). The effects were blocked by APJ-siRNA, inhibitor LY294002 and HIMO ( $P<0.05$ ). Conclusion Apelin promotes the proliferation and suppresses the apoptosis in GC *via* the PI3K/Akt signaling pathway.

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罗爽, 王继东, 何佳, 等. apelin对大鼠卵巢颗粒增殖与凋亡的影响[J]. 第三军医大学学报, 2013, 35(16): 1713-1716.

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