## Progress in Biochemistry and Biophysic

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## 肺腺癌A<sub>549</sub>/DDP细胞周期变化及其多药耐药性

Change of Cell Cycle and Resistance of  $A_{549}$  Cells to Cisplatin

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中文关键词: 多药耐药性 胞浆内游离Ca<sup>2+</sup> 细胞周期 肺腺癌细胞A<sub>540</sub>

英文关键词: <u>mul</u>tidrug resistance <u>intracellular free Ca<sup>2+</sup> cell cycle</u> <u>lung adenocarcinoma cancer A<sub>549</sub> cells</u>

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中文摘要:

用Fura-2/AM标记药物敏感的肺腺癌细胞A<sub>549</sub>和抗顺铂药物的肺腺癌细胞A<sub>549</sub>/DDP两种细胞胞内游离Ca<sup>2+</sup>,用碘化丙锭(PI)标记细胞DNA,检测其胞内Ca<sup>2+</sup>的 变化及两种细胞增殖能力和细胞周期. 实验结果表明,抗药性细胞株A<sub>549</sub>/DDP胞浆内游离Ca<sup>2+</sup>的浓度仅为药物敏感细胞株A<sub>549</sub>的1/3左右,同时前者的细胞增殖能力 较后者明显增强,而且细胞周期也明显缩短. 当用BAPTA-AM和EGTA或A $_{23187}$ 和Thapsi gargi n处理细胞以降低或升高其胞内自由Ca $^2$ +浓度时可改变细胞的生长周期, 二者也呈现明显差别. 这些结果表明,对顺铂产生耐药性的人肺腺癌A<sub>549</sub>/DDP细胞胞内Ca<sup>2+</sup>浓度的降低,可能影响细胞的增殖,缩短细胞的生长周期,特别是影响 起决定作用的G1期,从而有利于肿瘤细胞多药耐药特性的维持.

## 英文摘要:

The change of intracellular free  $\mathrm{Ca}^{2^+}$  in  $\mathrm{A}_{549}$  cells sensitive and  $\mathrm{A}_{549}/\mathrm{DDP}$  cells resistant to the cis-dichlorodiammine platinum (cisplat in) were measured by Fura-2/AM, the proliferation ability and cell cycle were measured by propidium iodide(PI) labeling cellular nuclear DNA. The results indicated that the concentration of intracellular free calcium of the sensitive  $A_{549}$  cells was 2 times higher than that of the re sistant  $A_{549}/DDP$  cells; the proliferation ability of the latter increased significantly than that of the former, the cell cycle also shortene d. The proliferation ability and cell cycle of the two cell lines also clearly showed difference by decreasing or increasing their intracellu lar free calcium concentration if the cells were treated with BAPTA-AM or EGTA and  ${\sf A}_{23187}$  or Thapsigargin. All of the results demonstrated that t the concentration decrease of intracellular free calcium in the  $A_{549}/DDP$  cells resistant to cisplatin may affect the cellular proliferation n, shorten the cellular cycle, which would be helpful to remain the multidrug resistance characteristics of  $A_{549}/DDP$  cells by specially modul ating the cellular decisive G1 of cell cycle.

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