PERITONEAL FIBRINOLYSIS DURING PNEUMOPERITONEUM AND LAPAROSCOPIC SURGERY

AKADEMISK AVHANDLING

Som för avläggande av medicine doktorsexamen vid Göteborgs Universitet kommer att offentligt försvaras I Bakfickan, Servicehuset, Sahlgrenska Universitetssjukhuset-Östra, Göteborg torsdagen 6 december 2007 13.00

av

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leg läkare

Avhandlingen baseras på följande delarbeten:

I. Peritoneal response to pneumoperitoneum and laparoscopic surgery.
   Bergström M, Ivarsson ML, Holmdahl L,

II. CO₂ promotes plasminogen activator inhibitor type 1 expression in human mesothelial cells
   Bergström M, Falk P, Holmdahl L,

III. Effect of acidosis on expression of mesothelial cell plasminogen activator inhibitor type-1
    Bergstrom M, Falk P, Holmdahl L,
    Surg Endosc 2006;20(9):1448-52

IV. Peritoneal and systemic pH during Pneumoperitoneum with CO₂ and helium in a pig model
    Bergstrom M, Falk P, Park PO, Holmdahl L,

V. Clinical evaluation of peritoneal acidification and fibrinolytic response during laparoscopy, a randomized parallel group study comparing Helium and Carbon dioxide
    Bergstrom M, Falk P, Park PO, Bengtsson JP, Haglind E, Holmdahl L,
    in manuscript
PERITONEAL FIBRINOLYSIS DURING PNEUMOPERITONEUM AND LAPAROSCOPIC SURGERY

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BACKGROUND

Laparoscopic surgery is believed to induce less postoperative adhesion formation compared with open procedures, but information regarding biological impact of a laparoscopic approach is limited.

MATERIAL & METHODS

Peritoneal response to laparoscopic surgery was assessed in human peritoneal tissue in two clinical trials (paper I & V). Human mesothelial cell response to CO₂ was assessed in cell culturing media (paper II & III). Measurements of peritoneal pH was evaluated in an animal model (paper IV), and then performed in patients (paper V). Assays of the key fibrinolytic enzymes t-PA and PAI-1 were done at protein (paper I-III & IV) and mRNA levels (paper II & III).

RESULTS

The initial clinical study showed a similar decrease in peritoneal t-PA activity during both open and laparoscopic cholecystectomy. However, there was a higher initial peritoneal PAI-1 concentration in the laparoscopic group, which might be attributable to prior exposure to CO₂. The in vitro studies showed an up regulation of PAI-1 mRNA production in cells exposed to CO₂ or acidic conditions, and that acidification could be caused by CO₂. In vivo studies of peritoneal pH showed an immediate decrease to 6.5 during insufflation, reproducing the CO₂ effect during surgery. He did not affect peritoneal pH. When initial peritoneal exposure to CO₂ was controlled for, peritoneal levels of t-PA decreased and PAI-1 increased during laparoscopic surgery regardless of gas used. These findings are consistent with observations done in open surgery and indicate that the effect is related to the surgical trauma. However, the t-PA activity was better preserved using CO₂ suggesting that use of CO₂ might have less adverse effect on peritoneal fibrinolysis than He.

CONCLUSION

Peritoneal fibrinolytic response during laparoscopic surgery is similar to open surgery, but CO₂ elicits specific biological effects. Exposure of peritoneum and mesothelium to CO₂ leads to a local and systemic acidosis and seems to have direct effects on key fibrinolytic enzymes. The systemic acidosis is manageable through controlled ventilation. The clinical implication of the effect on peritoneal fibrinolysis is unclear, but the observations done in humans are consistent with a reduced propensity for adhesion formation.

Key words: Laparoscopy, Peritoneum, Fibrinolysis, Mesothelial cells, Carbon dioxide, Helium
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