

行政院國家科學委員會專題研究計畫 期中進度報告

雙截切連續事件資料之分析(1/2)

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執行單位：國立臺灣大學公共衛生學院公共衛生學系

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處理方式：本計畫可公開查詢

中 華 民 國 92 年 6 月 2 日

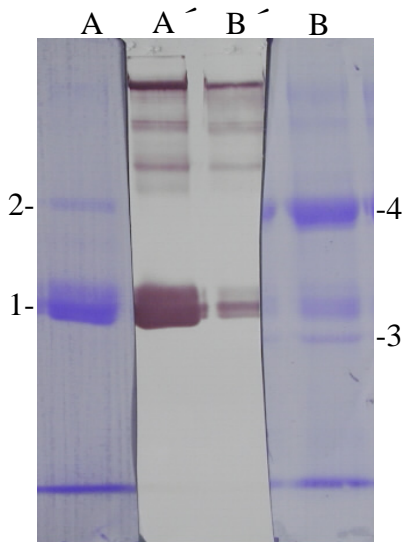


Figure 1. Western blot indicating the specific proteins recognized by rabbit antiserum against crude progesterone receptor proteins. A & B different fractional membrane proteins purified from progesterone-conjugated affinity column were separated by SDS-PAGE. A' & B' respectively corresponding to A & B, were further processed western blot with rabbit antiserum. The major apparent protein (app. 720 kDa, protein 1) were revealed by rabbit antiserum combined with peroxidase-conjugated goat anti-rabbit IgG. However, protein 2 (127 kDa), 3 (64 kDa) and 4 (125 kDa) are other major proteins that didn't recognized by rabbit antiserum.

Figure 2. Immunoreactivity of antibodies from rabbit antiserum against pig crude progesterone receptor proteins using immunofluorescence (A). Intense fluorescence had been observed over the whole acrosomal region. Control rabbit serum did not react with sperm. The conformation of the sperm with intact plasma membrane is made visible under a light microscope. Fluorescence was observed when 1st antibody was omitted.

Table 1. Effect of antibodies from rabbit antiserum against crude progesterone receptor proteins on sperm motility (means ± SD).

	Control	Male 1	Male 2	Female 1	Female 2
Activated serum	80±10 ^a	40±23 ^b	29±18 ^b	2±3 ^c	15±7 ^c
Inactivated serum*	82±8	65±15	80±17	75±13	69±17

***Inactivated serum:** serum were heat-treated at 56°C for 30 min to inactivate complement.

^{a,b,c} different superscripts are significantly different ($p < 0.05$ to $p < 0.001$) and same superscripts are non-significant ($p > 0.05$).

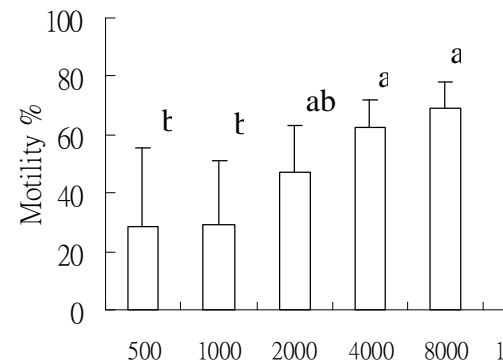


Figure 3. Effects of dilution of rabbit antiserum against crude progesterone receptor proteins (female 1) on sperm motility in the presence of guinea pig complement (55ng/mL). Bars represent mean ± SD. ^{a,b} different superscripts are significantly different ($p < 0.05$) and same superscripts are non-significant ($p > 0.05$).

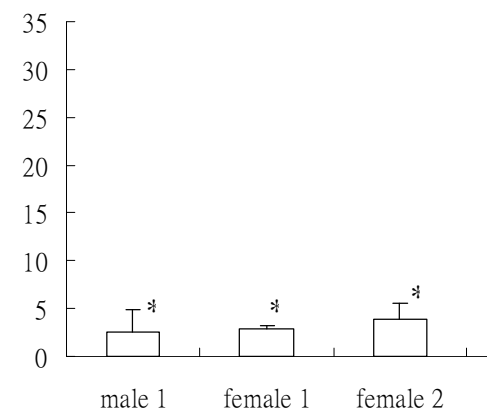
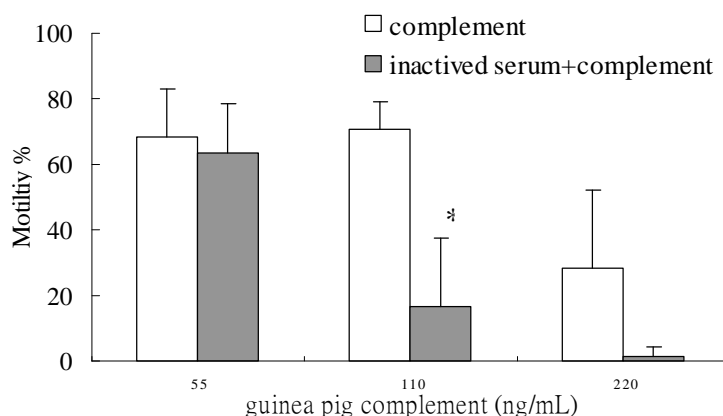


Figure 4. Effect of inactivated rabbit anti-PRs antiserum (female 1, 1:10) combined with guinea pig complement on sperm motility. Bars represent mean \pm SD. * $p < 0.05$ compared with the control, which had the same complement concentration as the treatment but without inactivated antiserum.

Figure 5. Effect of different inactivated rabbit antiserum (female 2 & 3) against crude PR proteins in a sperm motility binding assay. Sperm were preincubated with rabbit antiserum for 1 h, then co-incubated with crude PR proteins for 1 h. After incubation, 3000 motile sperm were added to medium containing salted piglet oocytes. Bars represent mean \pm SD. * $p < 0.05$ compared with the control.