

Supplementary material (Recuero et al.)

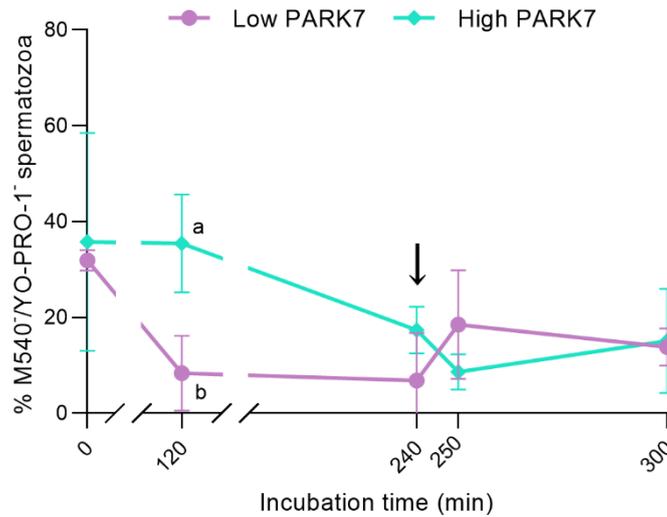


Figure S1. Percentage of viable spermatozoa with low lipid membrane disorder (M540-/YO-PRO-1-) during in vitro capacitation and progesterone-induced acrosome exocytosis (300 min). Black arrow indicates the time at which progesterone was added at a final concentration of 10 $\mu\text{g}/\text{mL}$ (240 min). Data are shown as mean \pm SD. Different letters indicate significant differences ($p < 0.05$) between samples containing high or low levels of PARK7 at 0 min of incubation.

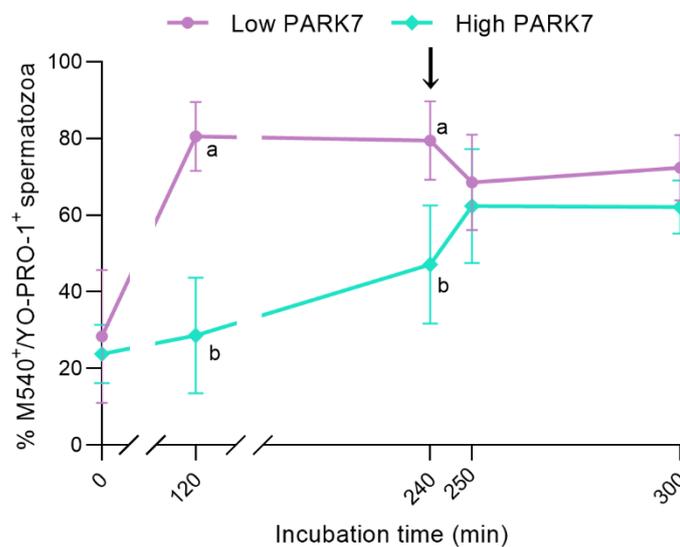


Figure S2. Percentage of non-viable spermatozoa with high lipid membrane disorder (M540+/YO-PRO-1+) during in vitro capacitation and progesterone-induced acrosome exocytosis (300 min). Black arrow indicates the time at which progesterone was added at a final concentration of 10 $\mu\text{g}/\text{mL}$ (240 min). Data are shown as mean \pm SD. Different letters indicate significant differences ($p < 0.05$) between samples containing high or low levels of PARK7 at 0 min of incubation.

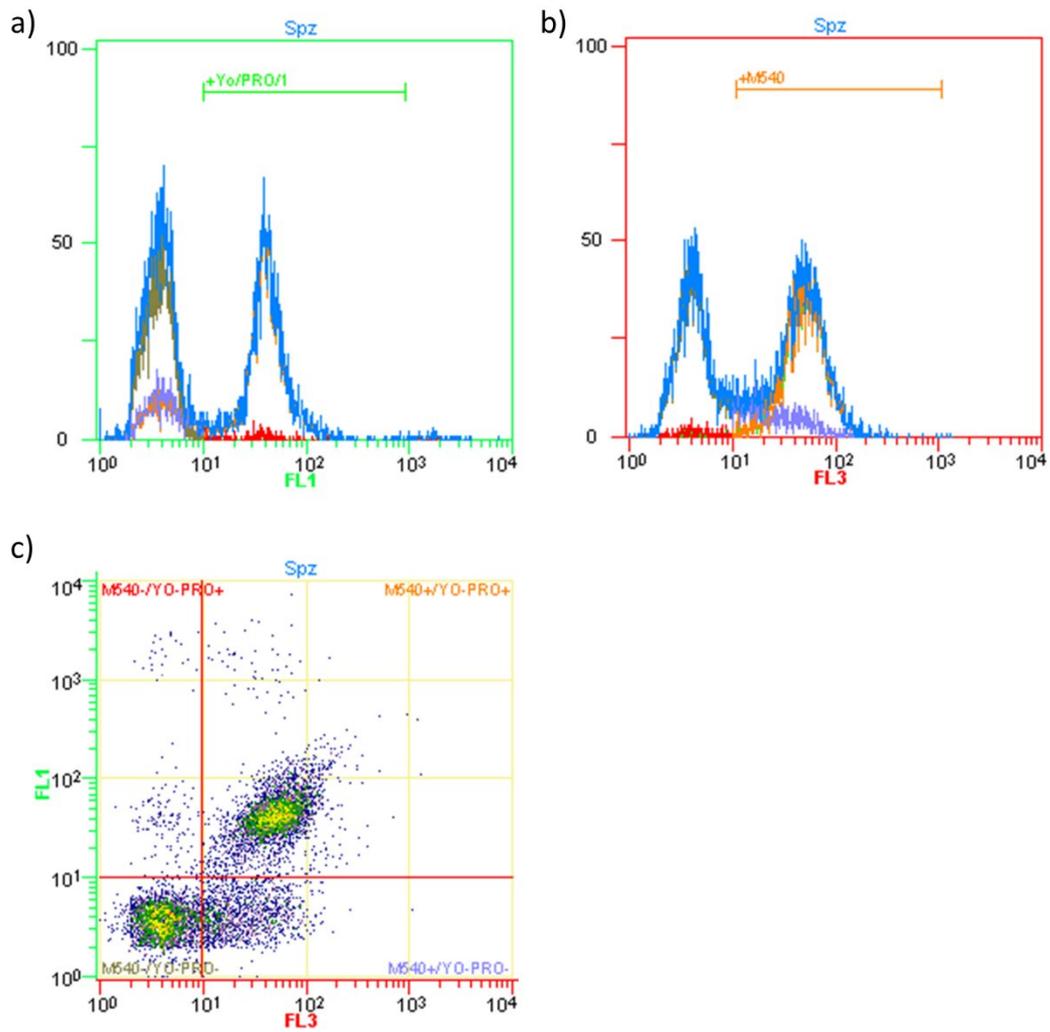


Figure S3. Representative flow cytometry YO-PRO (a) and M540 (b) histograms of sperm samples at the beginning of the in vitro capacitation experiment (0 min of incubation). (c) Dot plots of M540 and YO-PRO staining showing four distinct populations: M540-/YO-PRO⁺, M540+/YO-PRO⁻, M540-/YO-PRO⁻, and M540+/YO-PRO⁺.

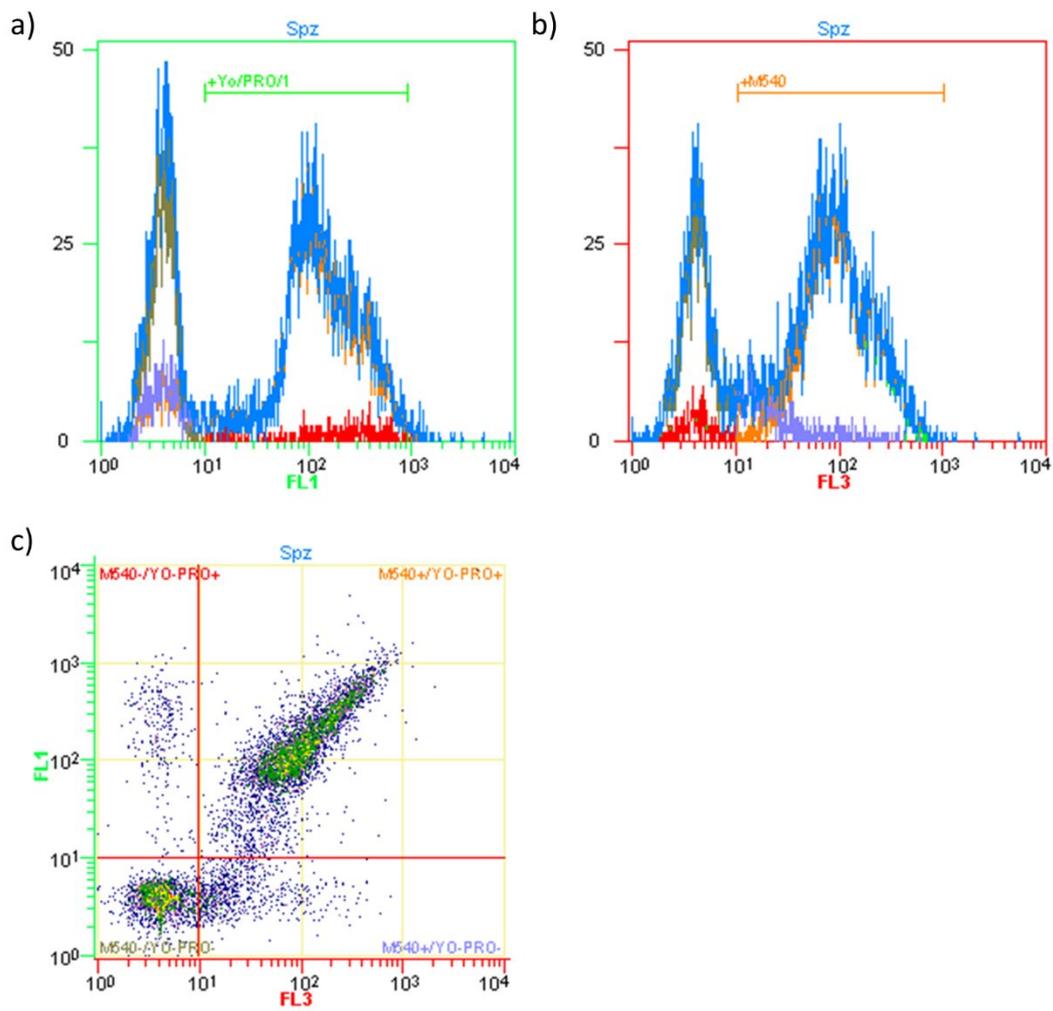


Figure S4. Representative flow cytometry YO-PRO (a) and M540 (b) histograms of sperm samples after 240 min of incubation in capacitation medium. (c) Dot plots of M540 and YO-PRO staining showing four distinct populations: M540-/YO-PRO⁺, M540+/YO-PRO⁻, M540-/YO-PRO⁻, and M540+/YO-PRO⁺.

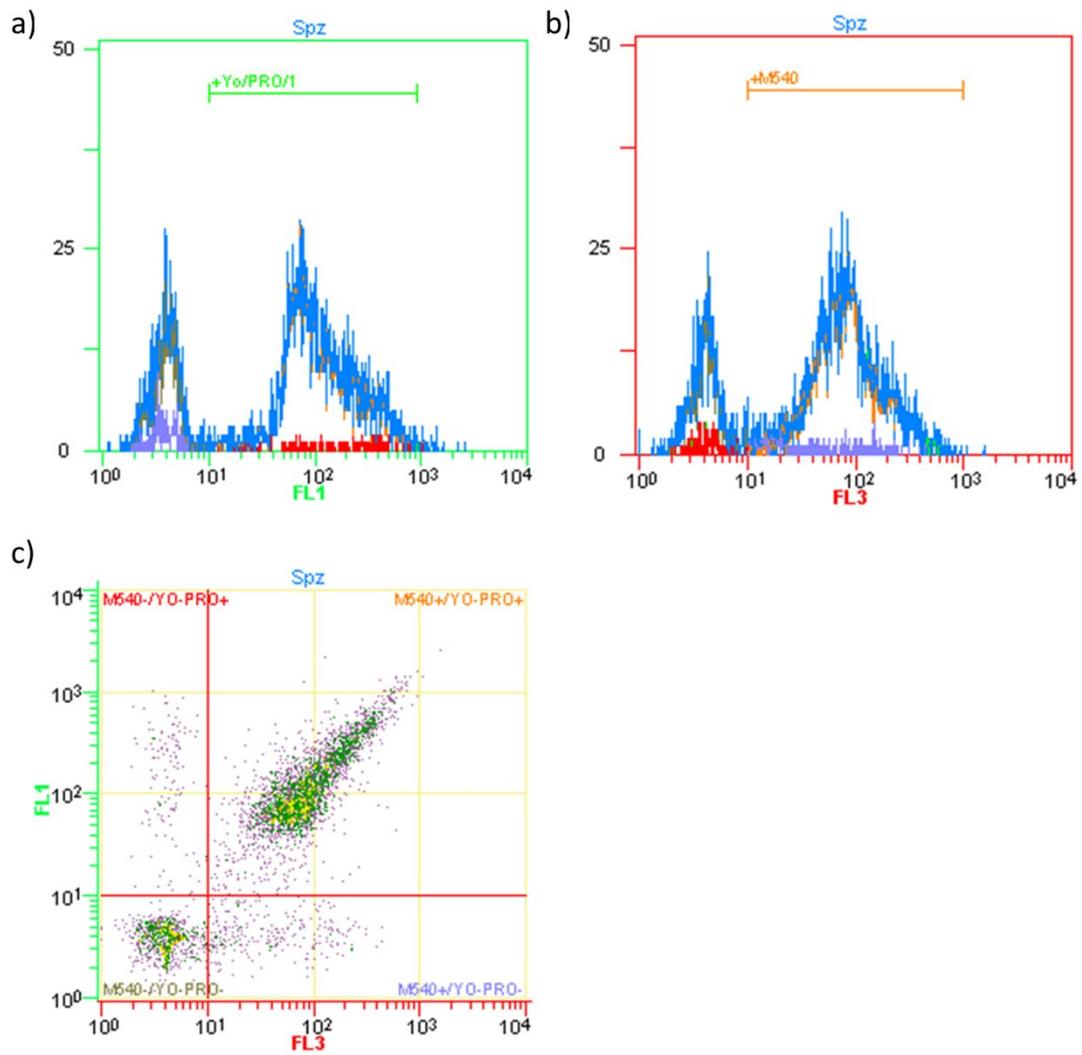


Figure S5: Representative flow cytometry YO-PRO (a) and M540 (b) histograms of sperm samples after 300 min of incubation in capacitation medium. (c) Dot plots of M540 and YO-PRO staining showing four distinct populations: M540-/YO-PRO⁺, M540+/YO-PRO⁺, M540-/YO-PRO⁻, and M540+/YO-PRO⁻.

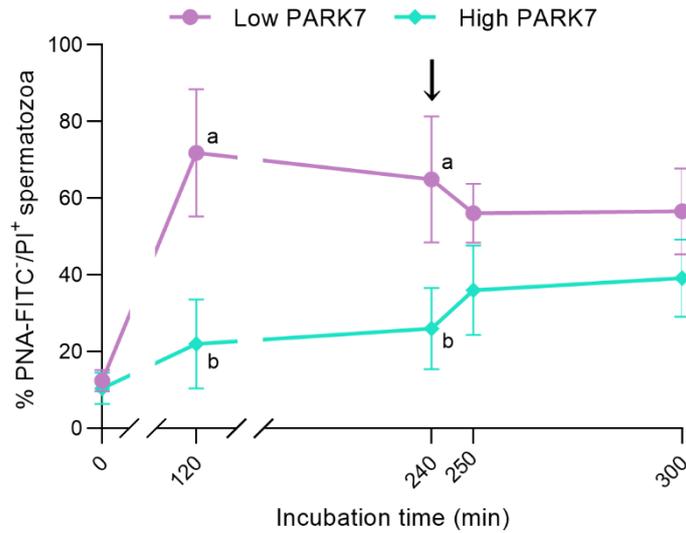


Figure S6: Percentage of non-viable spermatozoa with an acrosome that could not be fully intact (PNA-FITC⁺/PI⁺) during in vitro capacitation and progesterone-induced acrosome exocytosis (300 min). Black arrow indicates the time at which progesterone was added at a final concentration of 10 $\mu\text{g}/\text{mL}$ (240 min). Data are shown as mean \pm SD. Different letters indicate significant differences ($p < 0.05$) between sperm samples with high or low PARK7 levels at 0 min of incubation.

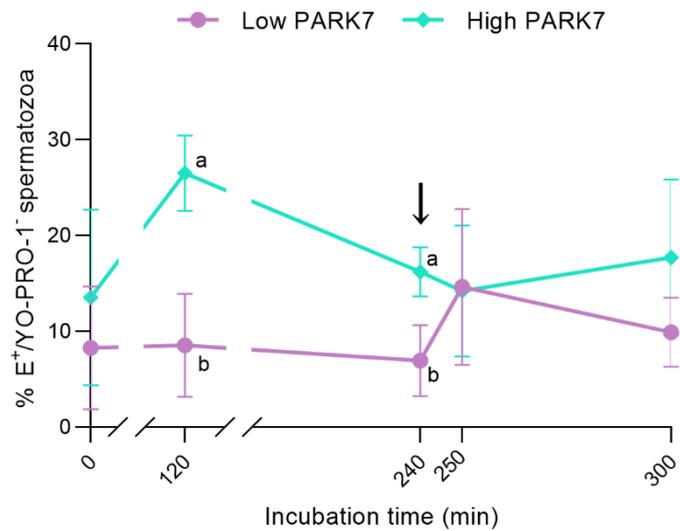


Figure S7: Percentage of viable sperm with high intracellular O_2^- levels (E⁺/YO-PRO-1⁻) during in vitro capacitation and progesterone-induced acrosome exocytosis (300 min). Black arrow indicates the time at which progesterone was added at a final concentration of 10 $\mu\text{g}/\text{mL}$ (240 min). Data are shown as mean \pm SD. Different letters indicate significant differences ($p < 0.05$) between sperm samples with high or low PARK7 levels at 0 min of incubation.