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The motor activity of the mouse spermatozoa during the in-vitro capacitation. We have examined the behavior of spermatozoa from two laboratory strains of mice, C57BL/6J and A/J, during the in-vitro capacitation by a modification of a previously described method (Williams et al., 1982). This modification involved changing the K⁺ concentration of the incubation medium from 0.1 to 0.2 M. In all experiments, spermatozoa from the A/J strain underwent significantly greater movement than those from the C57BL/6J strain. This difference in movement was not attributable to differences in motility or to differences in fertilization rates between the two strains, since the motile percentages of the spermatozoa from the C57BL/6J and A/J strains were similar. Spermatozoa from both strains reached a plateau in movement after a short incubation period, whereas those from the C57BL/6J strain reached this plateau by 60 min. However, the velocity of the spermatozoa from the C57BL/6J strain increased dramatically after they reached this plateau, whereas the velocity of those from the A/J strain decreased slightly. These results show that the motile activity of spermatozoa is heterogeneous among the laboratory mouse strains and that the motor activity of mouse spermatozoa is influenced by factors in addition to the medium used in the in-vitro capacitation.

Nano-scale deposition of chiral ligands for the oriented immobilization of functional antibodies. A method is presented for the oriented immobilization of functional antibodies, in particular murine IgG, on a glass surface through the use of short chiral ligands. The immobilized antibodies show improved specific binding to antigens in a competitive ELISA assay. Because the ligands are small (2d92ce491b