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spermatozoa: a comparison of  
their responses to the cholinergic

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receptor agonists acetylcholine and carbachol. The actions of acetylcholine (ACh) and carbachol (CCh) on isolated epididymal rat spermatozoa were investigated and compared to the responses of rat and human spermatozoa. The ability of ACh and CCh to elicit contractile activity in spermatozoa was examined by the digital imaging method of Bright et al. [1994]. Spermatozoa, in: J. H. Adelman, editor. Principles of androgen

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action, fourth edition, Lippincott, Philadelphia, pp. 351-356]. The actions of ACh and CCh on isolated rat spermatozoa were characterized by a response dose-response function and a rapid desensitization. Both ACh and CCh induced dose-dependent contractile responses in rat spermatozoa. However, the contractile responses evoked by ACh and CCh were less than that of rat spermatozoa incubated

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with the muscle relaxant, carbachol. The responses to ACh and CCh were inhibited by the muscarinic receptor antagonist atropine and the nicotinic receptor antagonist hexamethonium but not by the alpha-adrenergic receptor antagonist phentolamine. The contractile responses of rat spermatozoa to ACh and CCh were unaffected by the addition of cyclic AMP or cyclic GMP. In addition, the contractile

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responses of rat spermatozoa to ACh and CCh were not affected by the addition of protease inhibitors. However, the contractile response of rat spermatozoa to ACh was inhibited by the addition of the tyrosine kinase inhibitor genistein. On the other hand, the contractile response of rat spermatozoa to CCh was not affected by the addition of genistein. Incubation of rat spermatozoa with ATP resulted

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in a dose-dependent contractile response. This contractile response to ATP was unaffected by atropine, hexamethonium, phentolamine or genistein but was inhibited by the tyrosine kinase inhibitor, genistein. The responses of rat spermatozoa to ACh and CCh were inhibited by the addition of 20  $\mu\text{M}$

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