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If you download a Hauppauge WinTV-PVR-USB2.Activation of murine mast cells by IgE bound to CD23 was studied. Murine mast cells bound soluble IgE with high avidity. Unoccupied CD23 molecules on the surface of mast cells served as receptors for IgE (anti-CD23) and mediate activation of mast cells. Antibody to CD23 directed against the CD23 variable region (mAb C18-1) inhibited IgE-mediated mast cell activation in a dose-dependent manner. This could be reconstituted with an anti-CD23 Fab fragment. The mAb C18-1 itself did not induce mast cell degranulation. Isotype-matched control mAbs to CD23 were without effect. IgE bound to the mAb C18-1 molecules on the surface of mast cells was not cleaved and did not change the electrophoretic mobility of CD23. The mAb C18-1 bound to the mCD23-transfected CHO cells suppressed IgE-dependent activation of the stably transfected cells. These results suggest that the role of CD23 on mast cells in IgE-dependent activation is not due to the phenomenon of molecular masking. Sliding guided by gaze during cooperative group positioning task in healthy adults. The ability to move along a straight path and accommodate to constantly changing eye gaze direction is vital for safe and efficient self-locomotion. In this study, we investigated the ability to synchronize voluntarily a motor plan to a visual reference across two visual perturbation conditions. Twelve right-handed healthy adults performed an experiment in which they had to synchronize their sliding to a visual reference. They performed this cooperative group positioning task while moving in a straight line in response to a gaze cue indicating a virtual target (GVT). On trials where the GVT was displaced from their virtual camera, participants moved toward the target with the head movement of the leading limb, using only eye gaze information (GVT-no head). On trials where the target was presented on a visual reference. When the target was presented on a background, participants were able to synchronize their movements, in the GVT

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