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rm4404 either with preblocking of fc receptors for 10 minutes using 0 25 μg of purified anti mouse cd16 32 fig a or without preblocking of fc receptors fig b note the decreased staining of v 7 tcr on cd3 negative cells when fc receptors are blocked note the flow cytometric data shown above may not necessarily have

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about tim bushnell phd tim bushnell holds a phd in biology from the rensselaer polytechnic institute he is a co founder of and didactic mind behind excyte the world s leading flow cytometry training company which organization boasts a veritable library of in the lab resources on sequencing microscopy and related topics in the life sciences

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designing a high dimensional flow cytometry experiment is principally not different from designing any other scientific experiment which tests a defined hypothesis once it is established that high dimensional flow cytometry is needed to test the hypothesis one can follow a simple flow chart fig 1 it is important to define the target cell

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