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cell membrane. In contrast, IGFBP-4 expression was consistently reduced in response to uPAR expression. The importance of this effect was underscored by the finding that uPAR-mediated IGFBP-4 downregulation inhibited the IGFBP-4-mediated autocrine loop of growth, migration and invasion in these cells. In summary, these data reveal that uPAR is an important regulator of IGFBP-4 expression in breast carcinoma cells, and may therefore contribute 3da54e8ca3

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