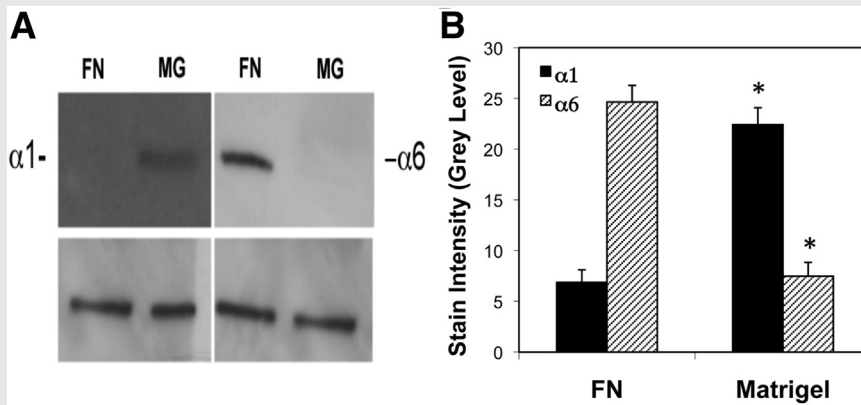


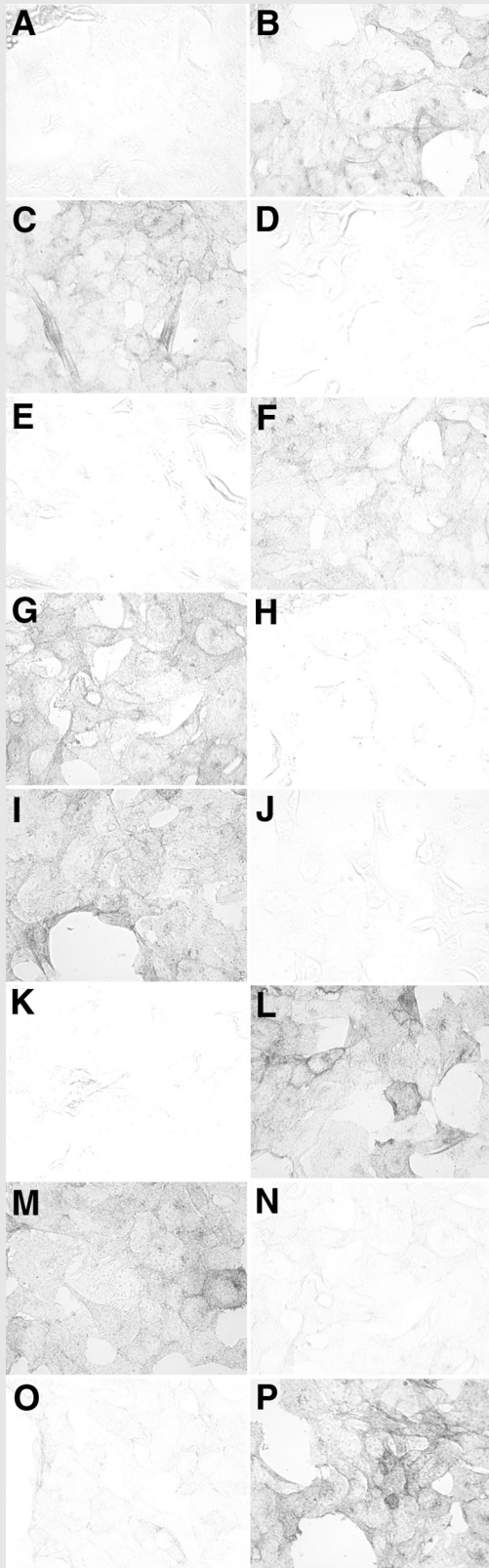
SUPPLEMENTAL FIGURE 1



Detection of integrin switching. (A) HTR-8/SVneo cytotrophoblast cells were cultured for 24 hours on fibronectin (FN) or Matrigel (MG) and analyzed by Western blotting of integrin subunits, as previously described (57). Each lane contained 30 μ g of protein extract and was labeled with antibodies against the indicated proteins (upper panels), and were stripped and reprobbed for β -actin (lower panels). (B) Integrin expression was measured by quantitative immunocytochemistry of $\alpha 1$ and $\alpha 6$ in cells cultured on either FN or MG. * $P < .05$ vs. FN. Both analytical approaches reveal integrin switching induced by Matrigel.

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SUPPLEMENTAL FIGURE 2



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SUPPLEMENTAL FIGURE 2 Continued

Expression of the $\alpha 1$ and $\alpha 6$ integrin subunits was determined as in Figure 2B. Examples of immunocytochemical staining of $\alpha 1$ (A, C, E, G, I, K, M, O) and $\alpha 6$ (B, D, F, H, J, L, N, P) in cells cultured with vehicle (A, B), 350 ng/mL sildenafil (C, D), combinations of sildenafil with L-NAME (E, F) or D-NAME (G, H), cGMP analogue (I, J), sildenafil with cGMP inhibitor (K, L), SNAP (M, N), or SNAP with cGMP inhibitor (O, P).

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