

Studying the interactions of COVID Treatment with the Developing Fetus

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A cross-section of developing placenta stained with Haematoxylin









In a healthy placenta, IgG antibodies are transferred from the mother to the developing child. There is evidence that exogenous IgG can cross the placental barrier just as well as endogenous IgG. These effects are dependent on gestational age and IgG concentration in the blood.





Lysossomal enzymes

Palmeira, Patricia et al. "IgG placental transfer in healthy and pathological pregnancies." *Clinical & developmental immunology* vol. 2012 (2012): 985646. doi:10.1155/2012/985646



Sections of placental tissue were co-cultured with monoclonal antibody treatments casirivimab and imdevimab for twenty-four hours at 2.4mg/mL to allow the antibodies to cross the syncytiotrophoblast cells







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The incubated placental tissue was fixed with formyl and embedded in paraffin. Then 3 micrometer cross-sections were cut on a microtome (left) and mounted on slides for staining using immunohistochemistry (right).





For our immunohistochemistry protocol, the primary antibodies are casirivimab and imdevimab, added before the tissue is fixed. The secondary antibody we use is a polyclonal rabbit anti-human conjugated to a peroxidase. This should produce a brown stain on the slides anywhere that the casirivimab and imdevimab have penetrated or attached to the tissue. Additionally, we add a haematoxylin stain to show cellular structures such as syncytiotrophoblasts and cytotrophoblasts



Image by Boster Bio



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Notice the different cell types and how they interact with the monoclonal IgG differently – the Cytotrophoblasts, Syncytiotrophoblasts, and the Mesenchyme.





Cytotrophoblasts do not appear to be permeable to antibodies, which is confusing because both Syncytiotrophoblasts and the Mesenchyme have antibodies present.





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Are there any questions?