

## Solutions to [Test Your Knowledge: Kidney Transplantation and the HIV](#)

### 1E. A&B only

Ritonavir and other protease inhibitors (PI) can significantly increase calcineurin inhibitor levels. Ritonavir is a potent inhibitor of the hepatic microsomal enzyme CYP (450) 3A4, of which cyclosporine and tacrolimus are substrates. Conversely, efavirenz strongly induces the CYP 3A4 system. As a result, serum cyclosporine and tacrolimus levels may decrease with efavirenz therapy. Careful immunosuppression monitoring is recommended with initiation or dose changes of concomitant efavirenz therapy in transplant recipients.

### 2. False.

As [Chandran and colleagues](#) describe in their AJKD article, the suppression of viral replication does not seem sufficient to prevent all HIV-mediated immunologic injury.

### 3C. Higher.

In the landmark article by [Peter Stock and colleagues](#), early and aggressive rejection was surprisingly higher in HIV-infected recipients despite low CD4 T-cell counts and control of viral replication.

### 4D. Has been explored as a possible option in HIV endemic areas.

[Muller and colleagues](#) describe in a letter to the editor in the *New England Journal of Medicine* four HIV-infected recipients who received kidney transplant from two HIV-infected deceased donors. These recipients had successful outcomes after the first year with regards to allograft function, HIV viral load, and CD4 cell count. To their credit, performing such a procedure took courage and saved four recipients from certain death in an area where dialysis therapies were not available. Uncertain is the possibility of co-HIV infection or viral mutation.