

### **Specifications for Interferon Gamma Release kits:**

1. Diagnostic IGRA kit should be capable of measuring the respective analytes in serum/plasma (as appropriate) of patients.
2. IGRA kit should be approved for in-vitro diagnostics (IVD) and have appropriate certifications (CE/US-FDA/CDSCO/BIS).
3. The kit should have detachable strips, so that 8/12 wells can be detached and run in batches of 8/12.
4. The seller can be asked for the proof of utility of diagnostic kits in reputed diagnostic laboratories in India.
5. The seller need to provide documentation as desired by the user department to assess the rate reasonability.
6. The seller will need to provide per plate cost for the respective analytes (96 wells).
7. The expiry of kits supplied must be at least 9 months from date of receipt at the user department.
8. The user department can change the quantity mentioned in the tender, depending on the departmental needs.
9. The kit should have separate collection tubes for unstimulated, mitogen stimulated and Tb antigen stimulated tubes.

