S.No	RFP Document Reference(s) (section number/ page)	Original Clause	Clause add/delete/modify	Amended Clause
1	Attachment A to Annexure 7.1 – Technical specifications, Page 5, 2. Broad specifications of the equipment of the Composite Medical Kit. Point n	n. Medical KITs must approved by DCGI (Drug Controller General (India) and also must approved by either USFDA (United States - Food and Drug Administration) or EU (European Union).	Modify	n. Medical KITs must be approved/certified by DCGI (Drug Controller General (India)) or USFDA (United States - Food and Drug Administration) or EU (European Union).