**Annexure 6**

**FORMAT A**

**Consent for audit of their manufacturing site/offices**

(To be signed by full time Director / CEO / MD of the company /Partners of the firm/Proprietor (as the case may be) duly depicting the Name & designation and submitted on official stationery of the applicant along- with the authorization to do so)

**1.**      Whereas, the applicant namely *(name of manufacturer with address)* has submitted an application under Production Linked Incentive Scheme (PLI) for domestic manufacturing of critical Key Starting Materials (KSMs)/ Drug Intermediates (DIs)/Active Pharmaceutical Ingredients (APIs) to Department of Pharmaceuticals (DoP), Government of India seeking incentives for the application pertaining to manufacturing…………(Eligible Product) at……………(location(s)).

**2.**      Now, therefore, the applicant or its agencies or its consultants engaged with the process of manufacturing of eligible products shall allow the PMA or any other authority as designated by DoP for verification of facility and documents submitted for the approval of application and disbursement of incentives under PLI Scheme.

*Date  (Name & designation with address)*

Full time Director / CEO / MD of the company/ Partner/ Proprietor of the firm