Device Schedule:

Intended Purpose as per the Instructions for Use:

The **cobas® liat** CT/NG/MG nucleic acid test is an automated, qualitative in vitro nucleic acid diagnostic test that utilizes real-time polymerase chain reaction (PCR) for the direct detection of Chlamydia trachomatis (CT), Neisseria gonorrhoeae (NG), and Mycoplasma genitalium (MG) nucleic acid in male/female urine and vaginal swabs (clinician-collected and self-collected), all in **cobas®** PCR Media (Roche Molecular Systems, Inc.).

This test is intended for professional use in a clinical laboratory setting, Near Patient Testing, or point-of-care (POC) location as an aid in the diagnosis of urogenital infections in both symptomatic and asymptomatic individuals.

Device Name	**Model**	**Type** (Codes as per (EU) 2017/2185)	**Risk Classification**	**Basic UDI-DI**
cobas® liat CT/NG/MG	09449604190	IVR 0503	Class C near-patient test	7613336030029W