Device Schedule:

Intended Purpose as per the Instructions for Use:

The **cobas**® **liat** CT/NG 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) and *Neisseria gonorrhoeae* (NG) 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	10030933190	IVR 0503	Class C near-patient test	761333603004A2