SUPPLEMENTARY INFORMATION

TABLE S1 Determination of patient infected status from urogenital specimens from

Sample type	BD ProbeTec [™] CT/GC Q ^x	Hologic APTIMA Combo 2 [®] CT/NG	Abbott RealTi <i>m</i> e CT/NG	PIS
Women ^a	Urine/vaginal	Urine/vaginal		
	+/+	+/+	NA	Infected
	+/+	+/- or -/+	NA	Infected
	+/- or -/+	+/+	NA	Infected
	+/-	-/+	NA	Infected
	-/+	+/- or -/+	NA	Infected
	-/+	+/-	NA	Infected (urine) Not infected
	l or l	1	ΝΙΛ	(vaginal)
	+/- or -/+	-/-	NA	Not infected
	+/+	-/-	NA	Not infected
	-/-	+/+	NA	Not infected
	-/-	+/- or -/+	NA	Not infected
	-/-	-/-	NA	Not infected
Men ^b	Urine	Urine		
	+	+	+	Infected
	+	+	-	Infected
	+	-	+	Infected
	_	+	+	Infected
	-	-	+	Not infected
	-	+	_	Not infected
	+	-	-	Not infected
	-	_	-	Not infected

women and urine specimens from men

CT, Chlamydia trachomatis; GC, Neisseria gonorrhoeae; NG, Neisseria gonorrhoeae;

NA, not applicable; PIS, patient infected status.

^a PIS was defined as positive if \geq 1 result was positive in each assay; PIS was defined as negative by any other combination of results. PIS was considered indeterminate if results for \geq 1 sample type were invalid and the results of the remaining sample types were discordant.

Clinical Performance of the cobas® CT/NG Test

^{*b*} PIS was defined as positive if ≥ 2 of 3 test results were positive; PIS was defined as negative by any other combination of results. PIS was considered indeterminate if 1 test result was invalid or missing and the other 2 test results were discordant, or if ≥ 2 of 3 test results were invalid.