

K911640 OLYMPUS PK TP(TM) SYSTEM, MODIFICATIONJul 24, 1991
107 days to decisionK911640 · Product code: **GMT** · Microbiology
Source: <https://www.510kdatabase.net/k911640/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Antigens, Ha, Treponema Pallidum (GMT)
Date received	Apr 8, 1991
Decision date	Jul 24, 1991
Days to decision	107 days
Third-party review	No

APPLICANT

Company	Olympus Corp.
Location	Mchenry, IL, US
Contact	LAURA STORMS-TYLER
Website	https://www.olympus-global.com
510(k) history	142 submissions · 140 cleared · 1978-1995

Olympus Corp. is a Japanese optics and imaging manufacturer founded in 1919. The company operates from McHenry, US, and holds approximately 70 percent of the global endoscope market. Olympus has received FDA 510(k) clearances from total submissions between 1978 and 1995. The company's cleared devices span gastroenterology, urology, obstetrics, gynecology, and chemistry specialties. This regulatory record reflects the company's historical focus on endoscopic surgical instruments and related technologies. Notable cleared device categories include resection electrodes, fiber...
