Label drawing & Label Location Diagram

Model: µ TASWako i30

Left Side













FCC ID X2IUTASWAKOI30 IC ID 8779A-UTASWAKOI30

This device complies with Part 15 of the FCC Rules. Operation is subject to the following two conditions: (1) this device may not cause harmful interference, and (2) this device must accept any interference received, including interference that may cause undesired operation.

This Class B digital apparatus complies with Canadian ICES-003. Cet appareil numérique de la classe B est conforme à la norme NMB-003 du Canada.

Wako Pure Chemical Industries, Ltd. 1-2, Doshomachi 3-Chome, Chuo-ku, Osaka 540-8605, Japan

IMMUNO ANALYZER

MODEL μTASWako i30 SERIAL No. MANUFACTURED

FFTPA

Complies with FDA performance standards for laser products except for deviations pursuant to Laser Notice No.50, dated (June 24, 2007). Manufacturer Wako Pure Chemical Industries, Ltd. 1-2, Doshomachi 3-Chome, Chuo-ku, Osaka 540-8805, Japan Fluorometer for Clinical Use

IMMUNO ANALYZER

μTASWako i30

100-240V~ 4-1.7A 50-60Hz

FOR IN VITRO DIAGNOSTIC USE



LISTED
LABORATORY EQUIPMENT
19NS

SN

405N101413