



Quality Certificate

VWR® LAB MARKER

As per the manufacturer, the below given product meets the following criteria:

North American Catalog No:	52877-310
Lot Number:	139025
Description:	VWR LAB MARKER FINE BLK PK10

The product was manufactured in accordance with the current FDA Quality System Regulation 21 CFR Part 820, Medical Device Directive 93/42/EEC, Medical Device Quality Management System EN ISO 13485, and Canadian Medical Device Regulation SOR/98-282.

Quality Control Testing:

Representative production samples are collected and inspected in accordance with current applicable product specifications. Inspection records are reviewed and signed off by qualified personnel for product release.

Device Listing/Manufacturing Site Registration/Pre-market Notification:

Medical Devices are listed with FDA per 21CFR 807. Manufacturing sites are registered with FDA per 21CFR 807. The devices satisfy FDA pre-market notification requirements per 21 CFR 807.

Signed:

A handwritten signature in blue ink that reads "Linda K. Shine".

Linda K. Shine
Manager - Supplier Qualification
Global Compliance
VWR International LLC

Date: Jul 11, 2017