Quality Certificate

VWR® LAB MARKER

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

North American Catalog No:	52877-310
Lot Number:	100039
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:

Ken Crossley
Manager
Quality Assurance

Date: May 05, 2016