



## Quality Certificate

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### VWR® LAB MARKER

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

<b>North American Catalog No:</b>	52877-310
<b>Lot Number:</b>	105056
<b>Description:</b>	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, appearing to read "Ken Crossley", written over a light blue horizontal line.

Ken Crossley  
Manager  
Quality Assurance

**Date:** June 22, 2016