Medical Device Production Quality Assurance System
Certificate GB23/00000313

The management system of

Davis, Schottlander & Davis Ltd

Fifth Avenue Letchworth Hertfordshire SG6 2WD United Kingdom

has been assessed and certified as meeting the requirements of

Part II of The Medical Devices Regulations 2002, Annex V [as modified by Part 2 of Schedule 2A to The Medical Devices Regulations 2002]

For the following products

The Scope of Registration appears on page 2 of this certificate

This certificate is valid from 25 March 2025 until 17 August 2028 and remains valid subject to satisfactory surveillance audits.
Issue 4. Certified since 17 August 2023

L. Henderson

Authorised by
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Medical Device Production Quality Assurance System
Certificate GB23/00000313, continued

Davis, Schottlander & Davis Ltd

Part II of The Medical Devices Regulations 2002, Annex V [as modified by Part 2 of Schedule 2A to The Medical Devices Regulations 2002]

Issue 4

Temporary Crown & Bridge Materials:
Quicktemp 2, Quicktemp Cosmetic,
Quicktemp Cosmetic Glaze & Bond.
Crown & Bridge Materials: Ceramic Powder -
Matchmaker Powders MC, enliven MC

Restoratives & Cements:
Light cured composite filling material

Prosthetics:
Acrylic Resin Teeth - Enigma, Enigmalife,
Natura, & 4natur.

Prosthetics Materials:
Pegasus Pourable cold cure denture Base material.
Pegasus Plus repair acrylic.
Pegasus plus denture base
Acrylic Denture Base - Enigma High Base.

Where the above scope includes class IIb or class III medical device(s), a valid Type Examination Certificate according to Annex III [as modified by Part 2 of Schedule 2A to The MDR 2002] is a mandatory requirement for each device in addition to this certificate to place that device on the market

Certification is based on reports numbered GB/PC/07450
Previous certificate number: N/A
Change in between this certificate and previous one: removal of devices form the scope: Ultrafineglass composite device, Cobalt chromium alloy - Croform Excel, Bonding alloys non-precious Matchmate NP20