



The Dental Laboratory Association of the State of New York

presents:



SafeLink FDA Workshop

All materials, equipment and people used in the production of your product must conform to the FDA's regulations. The documentation alone for this program is intimidating. Not to mention, you must ensure your staff, vendors, and clients are fully informed of your system according to FDA regulations. This can be overwhelming for the lab owner to do without assistance.

Saturday – June 7th, 2008

*Turning Stone
Casino & Resort*

Workshop Material To Be Covered:

**FDA Compliance - Title 21, Chapter I, Subchapter H, Part 820 - QS/GMP's
Review requirements of a Quality System/Good Manufacturing Practices
(QS/GMP's);**

Prepare documentation for your customized QS/GMP System which includes:

*Are You a
Target For FDA?*



**All Dental Labs MUST
have an FDA Compliant
Quality System and
Good Manufacturing
Practices!**

- FDA Registration Requirements
- Quality Assurance Maintenance
- Standard Operating Procedures
- Measurable Worker Competency
- Purchasing Controls

- Traceability of Patient Contact Materials
- Non-Conforming Product (Internal or External Remakes)
- Corrective and Prevention Action Procedures
- Records Retention and Document Control

**Attendees will receive a CD so they can customize
their own Quality Manual*

**Special Pricing for DLANY Members*

**CDT's attending workshop will receive 7 hrs.
Documented Scientific Credits for Regulatory Standards.*

CALL or CHECK ONLINE FOR PRICING!

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