

California Endoscopy is an ISO 13485 company certified in medical device repair

No, we are not audited by the FDA, they only register scope manufacturing, not scope repair

We are certified by an independent 3rd party company (Hospital = JCAHO)

3rd party company is audited and certified by ANAB

Standards used are written by 156 countries, the US being one of them

Standards and regulations are almost identical to FDA 21 CFR820, which is the quality system regulations used by the FDA to certify the process

We are required to comply with FDA 510K regulations which certifies the device

We are audited by the 3rd party company every six months

 CALIFORNIA ENDOSCOPY

