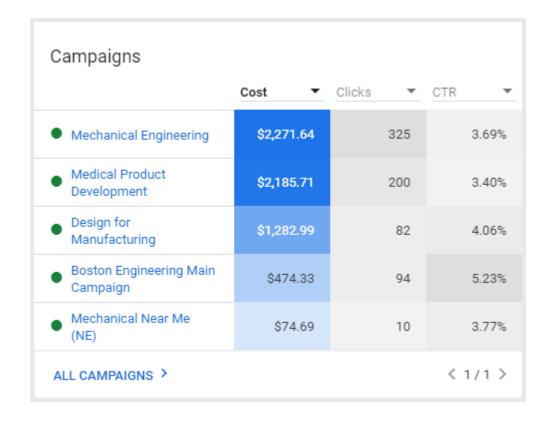
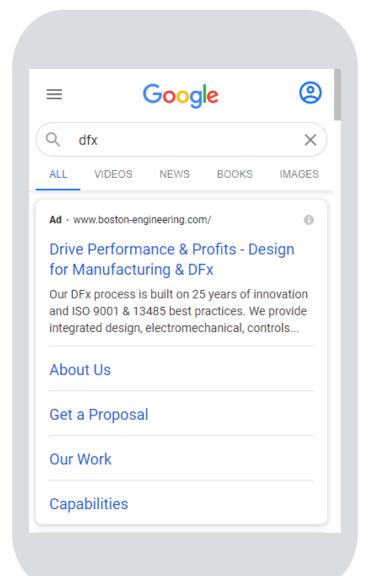
DIGITAL ADVERTISING: GOOGLE AND LINKEDIN ADS

Alex Wallace
B2B Marketing and Demand Generation
www.wallace.marketing

GOOGLE SEARCH ADS

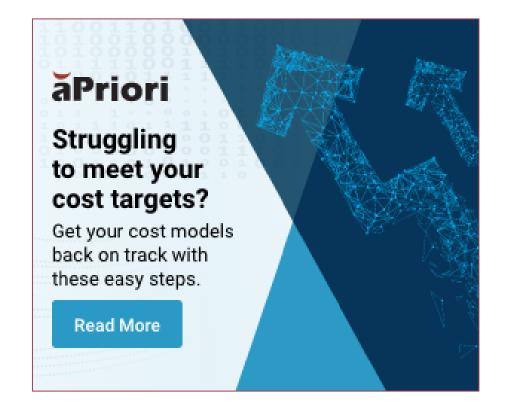




GOOGLE DISPLAY ADS



How to get your cost targets back on track with these easy steps Read More





LINKEDIN SINGLE IMAGE AD

This LinkedIn Ads campaign generated leads via gated content – including a \$1.7M sales opportunity.





Rethink Post-market Surveillance (PMS) for Medical Devices

How Smart, Connected Medical Devices Can Deliver Critical Insights

Six years after receiving FDA clearance, a medical device company's investment in a new therapeutic area is finally paying dividends. But a new patient injury report could send the company reeling if it can't identify the root cause of the device failure quickly.

The financial impact for product recalls and related corrections can be significant. The medical device industry spends \$2.5-\$5 billion annually to address non-routine quality events in the US – such as major observations, recalls, warning letters, consent decrees, warranties, and lawsuits – according to McKinsey & Company¹.

The consulting firm also reports the cost of a major medical device recall or other single non-routine quality event can reach \$600 million. The incident review and remediation process itself can take months or years to complete due to manual processes such as gathering old device data.

So, even if a device did not cause an adverse event, the company's sales and its reputation could still be negatively affected during this protracted process.

Eliminate Performance Blind Spots

The lack of visibility into device field performance has a significant impact beyond patient safety and risk management. Failure to truly understand how customers are using a device represents a missed opportunity to make user-driven enhancements and to identify untapped market needs.

To address this challenge, medical device companies are incorporating secure Industrial