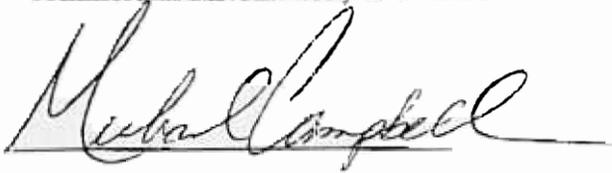


Premarket Notification  
510(k) Statement

I certify that in my capacity as Regulatory Affairs and Quality Assurance Manager of Olympus America Inc., I will make available all information included in this premarket notification on safety and effectiveness within 30 days of request by any person if the device described in the premarket notification is determined to be substantially equivalent. The information I agree to make available will be a duplicate of the premarket notification submission, including any adverse safety and effectiveness information, but excluding all patient identifiers, and trade secret and confidential commercial information, as defined in 21 CFR 20.61.

A handwritten signature in cursive script, appearing to read "Michael Campbell", written over a horizontal line.

Name: Michael Campbell

Date: January 31, 2003