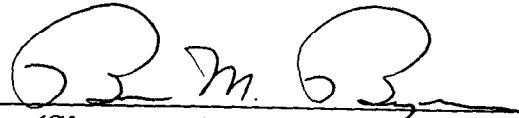


HemoCue, Inc.
510(k) Premarket Notification
HemoCue Low Hemoglobin System
July 21, 2000

510(k) STATEMENT
(As required by 21 CFR 807.93)

I certify that, in my capacity as President of HemoCue, 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.



(Signature)

Bruce M. Burgess

(Typed Name)

SEPTEMBER 20, 2000

(Dated)

510(k) Premarket Notification: K: _____