



Hycor Biomedical, Inc.
7272 Chapman Avenue
Garden Grove, California 92841

800 382 2527 telephone
714 933 3222 facsimile
www.hycorbiomedical.com

Certification of Product Quality

Shipment To:	ANICAM CARGO INC., 1770 NW 96 TH AVE., DORAL, FL 33172				
Shipper No.	486693.1				
	Product Description	List/Part No.	Lot No.	Quantity	Expiration Date
	KOVA PETTER, 500	87135E	147339	1	Not Applicable
	KOVA PETTER, 500	87135E	147755	91	Not Applicable
	KOVA DECANTING RACK	87136	N/A	6	Not Applicable
	KOVA TUBE, 500	87137E	148049	59	Not Applicable

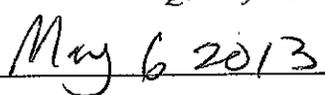
HYCOR BIOMEDICAL INC., a manufacturer of medical devices and diagnostic products licensed by both the United States Food and Drug Administration and the State of California, certifies that this product has been manufactured, tested and controlled according to the requirements of the United States Food Drug and Cosmetic Act, the Canadian Medical Devices Regulations, SOR/92-282, the In-Vitro Diagnostic Devices Directive (IVDD) and the Quality System Regulation.

All raw materials have been tested or inspected to Hycor's specifications to assure fitness for use. The products listed above have also been sampled, tested, packaged and stored according to Hycor's specifications.

Human serum source materials used in Hycor's products are tested for the presence of the antibody to the human immunodeficiency virus (HIV 1 and 2), for hepatitis B surface antigen (HBsAg) and for anti-hepatitis C (HCV), and found to be negative. Human serum source material is potentially infectious; therefore appropriate biohazard precautions should be taken.

Animal source materials used in Hycor's products are obtained exclusively from donor animals maintained under veterinary supervision and found free of contagious diseases.



Quality Certification


Date



7272 Chapman Avenue
Garden Grove
California
USA
92841

Telephone: 800-382-2527
Fax: 714-933-3222

Certificate of Origin

Hycor Biomedical Inc. Urinalysis (KOVA/Disposable Plastics/Controls) Products

We, Hycor Biomedical Inc. of 7272 Chapman Avenue, Garden Grove, California, 92841, USA, hereby declare that the following product(s): **Urinalysis (KOVA/Disposable Plastics/Controls) products** are manufactured in California in the United States of America according to Quality System Regulations (QSR). Hycor Biomedical Inc. holds certificates of registration of Quality System to I.S. EN ISO 13485: 2003, ISO 9001:2008 and is registered as a medical device manufacturer in accordance with the Canadian Medical Device Regulation.

Signed on behalf of Hycor Biomedical Inc.

Date: _____

5/6/13

Quality Management