

**DEPARTMENT OF GASTROENTEROLOGY
ALL INDIA INSTITUTE OF MEDICAL SCIENCES**

Reference: **31/Gastro./Proprietary/HEV RT Quantification Kit/NGB Diag./2016-17**

January 10, 2017

Subject: Purchase of HCV-Quantitation-FRT, HEV cDNA, RT Quantification Kit, Department of Gastro., AIIMS, New Delhi, on proprietary basis- Inviting comments thereon.

The Department of Gastroenterology has been procuring the above cited **HCV-Quantitation-FRT, HEV cDNA, RT Quantification Kit**, Cat No. NGB-PATH-HEV-Stand with mentioned volumes from M/s Ecoli s.r.o., on proprietary basis. The proposal submitted by M/s NGB Diagnostics, and PAC certifications are attached & uploaded on website.

The above documents are being uploaded for open information to submit objections, comments, if any, from any manufacturer regarding proprietary nature of the equipment/item within 15 days from the date of issue/uploading of the notification giving reference no. **31/Gastro./Proprietary/HEV RT Quantification Kit/NGB Diag./2016-17.**

The comments should be sent to Professor & Head, Department of Gastroenterology, Room No. 3111, Teaching Block, 3rd Floor, AIIMS, New Delhi on or before 28 January 2017 upto 5:00 p.m., failing which it will be presumed that any other vendor is having no comment to offer and case will be decided on merits.

Yours sincerely,

Dr. Anoop Saraya
Prof. & Acting Head

Encl:

1. Quotation
2. PAC certificate



NGB DIAGNOSTICS PVT LTD
PLOT NO: 120, 3rd Floor,
Hargovind Enclave,
DELHI-110092, INDIA
PH- 01142683435, +91-9717652828
Email: rakesh@ngbdx.com

Quotation

Buyer:

Dr. UMESH KAPIL,

Head, Dept of Gastroenterology

All India Institute of Medical Sciences New Delhi, India

Quotation No- 816-AIIMS-22112016

Dated: 22- Nov-2016

S.no.	Catalogue no.	Description	No of Reactions	Shelf Life	Price
01	CAT#NGB-PATH-HEV- Stand	HEV-Quantitation-FRT.HEV cDNA Real time Quantification Kit	96	09 Months	85108.20
VAT @ 13.5%					11489.61
Total Amount					96598.00

Amount Chargeable (in words): Ninety Six thousand five hundred ninety eight.

Company Details:

PAN no: AAECN4454E

VAT no: 07776929551

Bank Name: Yes Bank

Account Number: 023983800001219

Account Name: NGB Diagnostics Private Limited

IFSC code: YESB0000239

Account type: Current

Branch Address: GROUND FLOOR, 195 RAM VIHAR NEW DELHI 110095.



This is a computer generated statement hence no signature is required



Ecoli s.r.o.

Studenohorská 12
841 03 Bratislava 47
Slovak Republic

ecoli@ecoli.sk

Tel: +421 2 64 789 336
Fax: +421 2 64 789 040
Mobil: +421 903 160 701

Authorization Certificate

Date: 26th July, 2016

Reference no: AC-CE-406

To whomsoever it may concern


Subject: Distribution Authorization to M/s NGB Diagnostics Pvt Ltd

We E.coli s.r.o. having our registered office at *Studenohorská 12, 84103 Bratislava 47, Slovak republic*, do hereby authorize **M/S NGB Diagnostics Pvt Ltd** having there registered office at *Plot no: 120, 3rd Floor, Hargovind Enclave, New Delhi: 110092, India* for all our bacterial/viral marker Detection Kits based on PCR, Real Time PCR and Capillary electrophoresis and for all other reagents including nucleic acid extraction, with effect from 1st April 2016 till 31st March 2017, to submit bids, negotiate, Quote, and receive orders on behalf of our products.

Authorised Signatory


Ecoli s.r.o.
Studenohorská 12
841 03 Bratislava
tel: +421 2 64 789 336 fax: +421 2 64 789 040
obchodné oddelenie

Marek Kalnický

 EC declaration of conformity (certificate no. 4122.58.01/0) for in vitro diagnostic medical device according to the Directive 98/79/EC, ISO certificate – EN ISO 13485, EC declaration of conformity for in vitro diagnostic medical devices according to Annex I, List B of the Directive 98/79/EC (IVD Directive), CE certificate (no. 110040 QS/NB/c) for in vitro diagnostic medical devices, according to Annex II, List B of the Directive 98/79/EC



Ecoli s.r.o.
Studenčohorská 12
841 03 Bratislava 47
Slovak Republic

ecoli@ecoli.sk
Tel: +421 2 64 789 336
Fax: +421 2 64 789 040
Mobil: +421 903 160 701

Proprietary Certificate

Reference no: SER-GA-351
2016

Date: 21st Nov,

Kit Items and respective contents:

Genesi® HEV-Quantative RT PCR KIT-PATH-STAND FRT

This is to certify that:

Genesi® HEV-Quantative RT PCR KIT-PATH-STAND FRT Real Time detection kits are based on proprietary designed & validated regions. No other company in our knowledge has such HEV genome detection methods in a single test. The methods are thoroughly investigated for specificity, sensitivity and reproducibility. Detection of HEV genome with 100% homology with a broad range of HEV sequences based on a comprehensive bioinformatics analysis & is based on the amplification of a pathogen genome specific region using Proprietary short Oligos. The amplified product is detected by using fluorescent dyes. These dyes are linked to Proprietary oligonucleotide probes which bind specifically to the amplified product. Monitoring of fluorescence intensities during the real-time PCR allows the detection of accumulating product without re-opening the reaction tubes after the PCR run. Genesi® HEV-genotype-FRT PCR kit uses Proprietary "hot-start mastermix with modified Polymerase for high efficiency", which greatly reduces frequency of nonspecifically primed reactions.

Kit detection probe configuration.

Genesi® HEV-Quantative RT PCR KIT-PATH-STAND FRT For detection, FAM/Green and JOE/Yellow/HEX/Cy3 channels are needed.

The analytical sensitivity depends on the clinical sample volume and is 100 IU/ml (if the sample volume is 100 ul) or 10 IU (if the sample volume is 1000 ul).

Authorised Signatory


Ecoli ECOLI s.r.o.
Studenčohorská 12
841 03 Bratislava
tel: +421 2 64 789 336 fax: +421 2 64 789 040
e-mail: ecoli@ecoli.sk

Marek Kalnický

EC declaration of conformity (certificate no. 4122.58.01/0) for in vitro diagnostic medical device according to the Directive 98/79/EC, ISO certificate - EN ISO 13485, EC declaration of conformity for in vitro diagnostic medical devices according to Annex I, List B of the Directive 98/79/EC (IVD Directive), CE certificate (no. 110040 QS/NB/c) for in vitro diagnostic medical devices according to Annex II, List B of the Directive 98/79/EC



An ISO 9001-2008 Company

Date - 19.12.16

To
The Management
AIMS
Delhi

We Indus Medical Instruments Pvt Ltd, a reputed manufacturer of Gastrointestinal Endoscopic Devices and the only importer of SK-Ella products having our factory at Ramchandrapur, Kolkata hereby authorize A.K.Medica, GH 5/7, Flat no- 922, Paschim Vihar, New Delhi- 110087 for the territory of Delhi to quote, supply and collect payments from hospitals. This Authorization is valid till 31st March, 2017.

Thanking You

Yours Truly

Indus Medical Instruments Pvt Ltd

