

Noroviruses is the main causative agent of gastroenteritis in all age groups, particularly during winter. These viruses are the major cause of acute gastroenteritis (AG) in adults and the second among children. Symptoms are characterized with sudden vomiting (sometimes violent) with diarrhea, and sometimes with abdominal pain, nausea and fever. Incubation period is about 12 to 72 hours. According to the weak infective dose estimated at 10 to 100 viral particles, the potential for transmission from person to person is significant. Noroviruses of genogroups I and II have a substantial gene diversity, and are correspondingly further divided into genotypes.

RELIABLE FOODBORNE VIRUS DETECTION

Confidence comes from a simplified workflow

EQUIPMENT AND REAGENTS PROVIDED BY BIOMÉRIEUX

Recommended Equipment and Reagents			
PCR Real Time	414 056	GENE-UP® Thermocycler	
Lysis & Extraction	421 367	Electronic Pipette for eGENE-UP	
Lysis & Extraction	421 366	eGENE-UP® Stand	
Lysis & Extraction	200 292	NucliSENS® lysis buffer (2 mL) 48 tests	
Lysis & Extraction	280 134	NucliSENS® easyMAG® Lysis Buffer (4 x1 liter)	
Lysis & Extraction	280 130	NucliSENS® easyMAG® Extraction Buffer 1 (4 x1 liter)	
Lysis & Extraction	280 131	NucliSENS® easyMAG® Extraction Buffer 2 (4 x1 liter)	
Lysis & Extraction	280 132	NucliSENS® easyMAG® Extraction Buffer 3 (4 x1 liter)	
Lysis & Extraction	280 133	NucliSENS® easyMAG® Magnetic Silica (384 extractions)	

Process Control Kit

Process Control Kit	KMG (50 Samples)	Mengo Extraction Control		
Virus Kits For Detection			Quantification Standards	
PCR Kit	KADVC (48 reactions)	Adenoconsensus@ ceeramTOOLS	Not available	
PCR Kit	KADV (48 reactions)	Adenovirus@ ceeramTOOLS	KQADV	Adenovirus – Q Standard
PCR Kit	KASV (48 reactions)	Astrovirus@ ceeramTOOLS	Not available	
PCR Kit	KENV (48 reactions)	Enterovirus@ ceeramTOOLS	KQENV	Enterovirus – Q Standard
PCR Kit	KHAV (48 reactions)	HepatitisA@ ceeramTOOLS	KQHAV	Hepatitis A – Q Standard
PCR Kit	KHEV (48 reactions)	HepatitisE@ ceeramTOOLS	KQHEV	Hepatitis E – Q Standard
PCR Kit	KNVGIGII (48 reactions)	NoroGIGII@ ceeramTOOLS	See KQNVGI & KQNVGII	
PCR Kit	KNVGI (48 reactions)	NorovirusGI@ ceeramTOOLS	KQNVGI	Norovirus GI – Q Standard
PCR Kit	KNVGII (48 reactions)	NorovirusGII@ ceeramTOOLS	KQNVGII	Norovirus GII – Q Standard
PCR Kit	KRV (48 reactions)	Rotavirus@ ceeramTOOLS	KQRV	Rotavirus – Q Standard
PCR Kit	KSAV (48 reactions)	Sapovirus@ ceeramTOOLS	KQSAV	Sapovirus - Q Standard

Other Detection Kits Available - Parasites			Quantification Standards	
PCR Kit	KCRYPT	cryptosporidium@ ceeramTOOLS	Not available	
PCR Kit	KGIAR	giardia@ ceeramTOOLS	Not available	

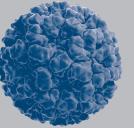
Full list of equipment, accessories and reagents upon request



DETECTION OF FOODBORNE VIRUSES

- Training according to OPTIMIZED bioMérieux protocols or
- Standardized work flow and kits complying to ISO15216
- Kits for extraction, detection, identification, quantification
- A large range of virus kits available
- LOD Method 250 genome copies/tested portion (shellfish 500)
- · High sensitivity & reproducibility

ROTAVIRUS - SAPOVIRUS - H

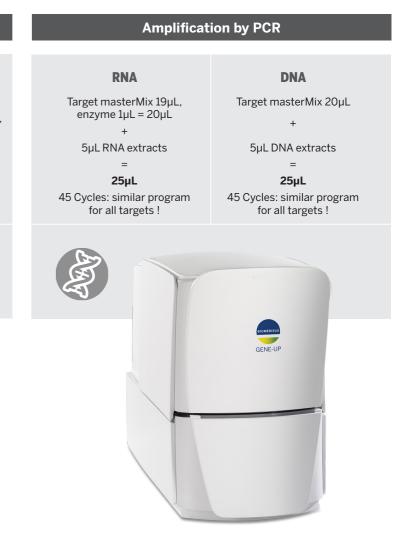


Request your practical training with our team of highly qualified Application Specialists to ease the implementation of a virus detection method in your own laboratory.

A COMPLETE WORKFLOW

Sample pre-treatment ISO or non ISO matrices **ISO** matrices Other matrices **Bivalve molluscan** Vegetables, leafs, Semi-dried tomatoes. **Soft fruits Bottled water Food surfaces** shellfish bulbs, stems RTE Food, date fruits... 2 grams of 1 liter 100 cm² 25 grams 25 grams **Upon request** digestive tissues 1 • Dissection (2 1 • Preparation 1 • Preparation of 1 • Filtration on 1 • Elution (swab Specific protocols g of digestive of samples samples (25 g membrane (10 with PBS, 10 µL are available for most glands (25 g in a sterile in a sterile filter μL of Mengovirus of Mengovirus) matrices. blender bag) bag) 2 • Elution of 2 • Lysis of virus 2 • Recovery of 2 • Elution (40 mL 2 • Elution (40 mL membrane and viruses (10 uL of Mengovirus. of TGEB. 1 mL of of TGEB. bottle (4 mL and 10 mL of TGEB. 2ml of pectinase. 10 µL Optional 1 mL of proteinase K) of Mengovirus) pectinase. 10 µL incubate, recoof Mengovirus) very of eluate. 3 • Clarification 3 • Lysis of virus second rinse of 3 • Clarification (retrieve eluate. bottle, secovery check pH, (retrieve eluate, of eluate) check pH. centrifuge) centrifuge) 3 · Concentration 4 • Incubation (40 (adjust pH mL supernatant 4 • Incubation (40 7, transfer in + 10 mL PEG/ mL supernatant concentration NaCl, incubate) + 10 mL PEG/ device, cen-NaCl, incubate) 5 • Concentration trifuge, retain 5 • Concentration (centrifuge, concentrate) (centrifuge, retain pellets) 4 • ILysis of virus retain pellets) 6 • Purification (chloroform 6 • Optional purifibutanol, cencation (chlorotrifuge, retain form butanol, aqueous phase) centrifuge, retain aqueous 7 • Lysis of virus phase) 7 • Lysis of virus



















Mengovirus Process Control for all Matrices - ISO Recommended







DNA/RNA Purification ISO recommended

Real Time PCR GENE-UP ceeramTOOLS Kits 1 - Detection 2 - Quantification







with EGENE-UP® & **NucliSENS®** reagents



EGENE-UP®

Your Ultimate Semi-Automated Device for Lysis & RNA/DNA Purification

BIOMÉRIEUX

eGENE-UP

The perfect add-on to GENE-UP® RT-PCR

Complete integrated workflow for foodborne virus detection with ceeramTools® kits

Safe, rapid & efficient recovery of purified nucleic acids

8 Samples

Cost, Time and Labour saving

Manual extraction systems are labour intensive, with low reproducibility, and typically demand a strict attention to detail. The EGENE-UP® semi-automated extraction platform speeds up the whole process, increases repeatability and reproducibility, whilst providing superior ergonomics for the user.



Automated pipetting

Reduced footprint





EGENE-UP®

Your next generation extraction platform

Performance

8 samples in less than 40 minutes can be processed with EGENE-UP®, an extraction performance equivalent to ISO15216 (Annex).

Conforms to the Standard Method

ISO15216: Microbiology of the food chain -- Horizontal method for determination of hepatitis A virus and norovirus using real-time RT-PCR -- Part 1: Method for quantification —Part 2: Method for qualitative detection Annex (informative): RNA extraction using NucliSens®

Space Saving

On your bench, limited space will be needed $175 \times 95 \times h160$ mm (max h200). Can even be placed in a hood when space is scarce.

Ref	Product	Content	
	Electronic	Pipette	
421367	Pipette for EGENE-UP®	Programming stand	
		Stand & Holder	
421366	EGENE-UP® Stand	2 x 96 grip tips 1250µl	
		10 48 well Deepwell plates	

Ref	Product	Content
200292	NucliSens® Lysis Buffer (Alternative to 280134)	48 x 2mL
280134	NucliSens® Lysis Buffer	(4 x) 1 L
280133	NucliSens® Magnetic Silica	24 x 1.2 mL = 28.8 mL
280130	NucliSens® Extraction Buffer 1	(4 x) 1 L
280131	NucliSens® Extraction Buffer 2	(4 x) 1 L
280132	NucliSens® Extraction Buffer 3	(4 x) 1 L

EASY with EGENE-UP

Prepare your microplate

Start your pipette, follow instructions

Nucleic Acids ready for the next step on the GENE-UP® RT-PCR thermal cycler

> Get ready for your next batch with a simple cleaning process

Reagent lay-out in a Deepwell Plate

Deepwell line 6 NucliSens®

Extraction Buffer 3 (2 mL)

Deepwell line 5 NucliSens®

Extraction Buffer 2 (2 mL)

Deepwell line 4 NucliSens®

Extraction Buffer 2 (2 mL)

Deepwell line 3 NucliSens®

Extraction Buffer 1 (2 mL)

Deepwell line 2 NucliSens®

Extraction Buffer 1 (2 mL)

Deepwell line 1

(for sample + Lysis buffer+ Silica)

