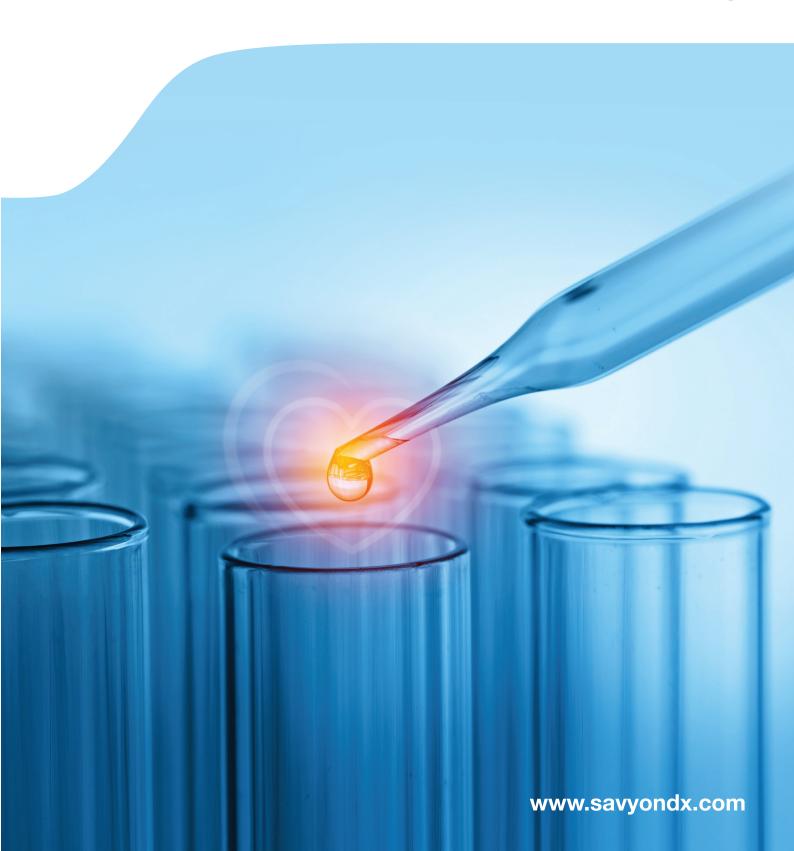
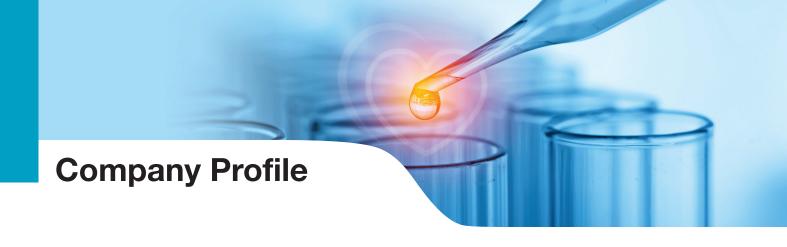


Product Catalog





Savyon Diagnostics develops, manufactures and markets high quality diagnostic kits and systems for the detection of Infectious Diseases for more than 35 years through a worldwide network of distributors.

Savyon Diagnostics offers a range of In Vitro Diagnostics kits for clinical laboratories, Point Of Care (POC) and Over-The-Counter kits allowing easy-to-use diagnostics in the comfort of one's home.

Savyon Diagnostics products are based upon various immunological and molecular biology techniques (ELISA, Lateral Flow, RT-PCR, MIF and others) and are developed by our experienced and skilled R&D team who maintain close relationships with international key opinion leaders and academic institutions. This collaboration expands Savyon's innovation in the field of in vitro diagnostics and enables the company to harness technologies and biomarkers that can be used in promoting diagnostics and in answering unmet clinical needs.

Savyon Diagnostics is accredited with the highest international quality standards of research, development and manufacture, including ISO 13485-2016. The company's products are all CE certified and those products sold in the USA, China, Mexico, Brazil and Australia are FDA, CFDA, ANVISA, COFERIS and TGA approved, respectively.

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DNA/RNA Extraction Pa	age 7
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ELISA

INTRODUCTION

Savyon Diagnostics offers a comprehensive line of Enzyme-Linked Immunosorbent Assays (ELISA) for clinical diagnostic use. Assays include our globally recognized line of infectious serology tests (The Sero Line), a wide range of direct antigen capture ELISA tests for enteric pathogens (The Copro Line), and our unique Haptoglobin typing ELISA for assessing the risk of cardiovascular complications in diabetic patients.

- Superior clinical and analytical performance
- Proprietary state of the art core biologicals Antigens, antibodies and enzyme conjugates
- Ready-to-use color-coded unified reagents
- Automation compatibility with all common ELISA processors
- Short and unified running protocols
- Convenience Based on break-apart 8 wells strip format, equally suited for both high or low volume use
- For quantitative or semi-quantitative assays Built-in controls for standard curve

Product	Description	Features Sample Type	Ordering Info & Approva		provals
			Tests/ kit	Catalog Number	
SeroCP	ELISA for qualitative detection of	Highly purified <i>C. pneumoniae</i> (TWAR-183) elementary bodies as antigens • Species-specific	IgG 96T	A191-01	CE ₀₄₈₃
ala	serum IgG, IgM and IgA antibodies to Chlamydia pneumoniae	• For IgG - 95%/97% sensitivity/specificity as	IgM 96T	A192-01	
	to Chiamyula pheumomae	compared with MIF (gold standard) • For IgM - 91%/95% sensitivity/specificity as compared with MIF • For IgA - 98%/97% sensitivity/specificity as compared with MIF Serum	IgA 96T	A193-01	
SeroELISA	SeroELISA ELISA for qualitative detection of serum IgG, IgM and IgA antibodies to Chlamydia genus	Based on purified L2 serovar MOMP antigen • Chlamydia genus-specific • Highly sensitive • Simple interpretation of results • Ready to use conjugate • For all antibody classes - 95% agreement with IPAzyme Serum	IgG 96T	A111-01	FDA CE ₀₄₈₃
m 24			IgM 96T	A112-01	CE ₀₄₈₃
			IgA 96T	A113-01	FDA CE ₀₄₈₃
SeroMP	ELISA for semi-quantitative	Based on highly purified M. pneumoniae native	IgG 96T	A261-01	FDA
ala	determination of serum IgG, IgM and IgA antibodies to <i>Mycoplasma</i>	antigen enriched with the pathogenic-state P1 adhesin • A set of 3 calibrators (P10, P50 and P75)	IgG 192T	B261-01	CE
	pneumoniae	allows for semi-quantitative determination of results (as expressed in BU/mL) • Ready to use conjugate	IgM 96T	A262-01	
		• For IgG - 95%/100% sensitivity/specificity as compared with consensus results (2 commercial	IgM 192T	B262-01	
		assays) • For IgM - 98%/96% sensitivity/specificity as compared with concensus • For IgA - 92%/80%	IgA 96T	A263-01	
		sensitivity/specificity as compared with concensus Serum	IgA 192T	B263-01	



Product	Description	Features Sample Type	Ordering Info & Approva		provals
			Tests/ kit	Catalog Number	
SeroPertussis	ELISA for semi-quantitative	Based on highly purified <i>B. pertussis</i> native antigen enriched with pertussis toxin and FHA • A set of	IgG 96T	A231-01	CE
64	determination of serum IgG antibodies, and qualitative determination of IgA/IgM antibodies to Bordetella pertussis	3 calibrators (P10, P50 and P75) allows for semi- quantitative determination of IgG (as expressed in BU/ mL) • Can detect both <i>B. pertussis</i> and <i>B. parapertussis</i> Serum	IgA/IgM 96T	A233-01	
Haptoglobin Typing ELISA	ELISA for qualitative detection of Haptoglobin phenotypes (Hp 1-1, Hp 2-1, or Hp 2-2) in human serum/ plasma of diabetic patients as an aid in predicting risk of coronary arterial and cardiovascular disease	The only available EIA for high-throughput Haptoglobin phenotyping • Based on patented monoclonal antibodies which differentialy bind to the different Hp isomers • A highly valuable tool for prognosis of CVD risk in diabetic patients and as a pharmacogenetic tool to assess the benefit of high dose vitamin E in such patients • Manual and automated protocols • Ready to use reagents • 98%/99% sensitivity/specificity for Hp 1-1, and Hp 2-1, 100%/99.5% sensitivity/specificity for Hp 2-2 as compared with standard electrophoretic techniques Serum/Plasma	96T	710-01	CE
CoproELISA H. pylori	ELISA for detection of Helicobacter pylori	Based on monoclonal antibodies to <i>H. pylori</i> Compatible with a variety of common preservatives (including formalin and SAF • No cross-reactivity with other enteric pathogens • Manual and automated protocols • Ready to use reagents • 96% agreement as compared with commercial FDA-approved EIA Fecal Sample	96T	774-01	CE





Savyon Diagnostics offers a wide portfolio of rapid tests for professional and self-use. We offer a wide range of tests for upper and lower respiratory tract infections, gastrointestinal pathogens, sexually-transmitted infections, urinary tract infections as well as tests for self- or professional assessment indications associated with women's health.

- Rapid Lateral Flow Immunochromatographic Assay (LFIA) & Enzymatic (Uriscreen) test
- · Reliable results
- Literature-supported superior performance
- · Simple and rapid operation
- Minimal (1-2 min) hands-on time
- Sample collection disposables included (e.g., swabs, vials, pipettes)
- Integrated and/or external controls
- Long shelf life
- Clear cut visual interpretation
- No need for instrumentation
- Selected combo tests for multiple targets
- CE-IVD
- Affordable
- Results obtained in 10 min

Product	Description	Features	Ordering Info & Approvals		
		Sample Type	Tests/ kit	Catalog Number	
Coprostrip C. difficile GDH+ToxA+ToxB	Rapid combo test for simultaneous detection of <i>C. difficile</i> GDH, Toxin A and Toxin B	LFIA • 3-in-1: Simultanous detection of GDH, Toxin A and Toxin B allowing for screening in confirmation of CDAD in one test • Differentiation of Toxin A and Toxin B - allowing for differentiation of hypervirulent strains • Compatible to the guidelines recommendation of GDH screening in combination with toxin testing to improve sensitivity • No cross reactivity with other enteric pathogens or other clostridial glycosyltransferases • Over 99% agreement with commercial rapid test for detection of C. difficile GDH and toxins Fecal Sample	20T	41220	CE
Coprostrip C. difficile Positive Control Set	External Quality Control Set for Coprostrip C. difficile GDH+ToxA+ToxB (Catalog # 41220)	Ready to Use Positive control set for GDH, Toxin A and Toxin B • Easy to use - supplied in dropper bottles • Sufficient for 20 individual tests for each parameter • Compatible with EU guidelines for laboratory external quality control Ready to Use Control Set	20T	41220-10	CE
Coprostrip H. pylori	Rapid test for detection of <i>H. pylori</i> antigen in stool samples	LFIA • No cross reactivity with other enteric pathogens • 94% sensitivity and 99% specificity as compared with commercial ELISA test for detection of H. pylori stool antigen (HpSA) Fecal Sample	20T	41221	CE
Coprostrip H. pylori Positive control	External Quality Control for Coprostrip <i>H. pylori</i> (Catalog # 41221)	Ready to Use Positive control set for Coprostrip H. pylori • Easy to use - supplied in a dropper bottle • Sufficient for 20 individual tests • Compatible with EU guidelines for laboratory external quality control Ready to Use Control Set	20T	41221-10	CE
Coprostrip Giardia	Rapid test for detection of Giardia lamblia	LFIA • No cross reactivity with other enteric pathogens • >99% agreement with microscopic examination of wet mounts Fecal Sample	20T	41217	CE



Product	Description	Features	Ordering Info & Approvals		
		Sample Type	Tests/ kit	Catalog Number	
Quickstripe Chlamydia Ag	Rapid test for detection of Chlamydia trachomatis specific antigen	LFIA • No cross reactivity with other STIs or with other chlamydial species • LoD = 1x10E5 Chlamydia EB/test • As compared vs. a commercial rapid immunoassay (INSTALERT) the test exhibits 97%/98% sensitivity/specificity for female cervical swabs, 98%/97% sensitivity/specificity for male urethral swabs, and 99%/94% sensitivity/specificity for urine samples Urine, Cervical or Urethral Swab	20T	41101	CE ₀₄₈₃
Quickstripe Chlamydia Ag with Positive Control	Rapid test for detection of Chlamydia trachomatis specific antigen (External Positive Control Incl.)	All features as in 41101 • External Positive control Reagent for Laboratory quality control Included in the kit Urine, Cervical or Urethral Swab	20T	41115	CE ₀₄₈₃
StrepA Stick	Rapid test for detection of Group A Streptococcus pyogenes antigen	LFIA • No cross reactivity with a broad list of other respiratory pathogens • Includes external positive and negative controls • LoD = 1X10E4 CFU • As compared with culture, the test exhibits >98% sensitivity and specificity Pharyngeal Swab	25T	R-6010	CE
No Step Strep A Test	Rapid patented test device for detection of Group A Streptococcus pyogenes antigen	LFIA • Patented easy to use no step device • Test exhibits >95.1% sensitivity and 97.8 specificity as compared with culture Pharyngeal Swab	1T	R-6009	CE
StrepB Stick	Rapid test for detection of Group B Streptococcus (GBS) antigens	LFIA • As compared with culture, the test exhibits 91% sensitivity and 98% specificity Vaginal / Rectal / General Swabs	25T	R-6012	CE
hCG Pregnancy Test	Rapid test for detection of human chorionic gonadotropin (hCG) at the sensitivity of 15 mIU/mL	LFIA • LoD = 15 mIU/mL • As compared with a reference commercial rapid EIA, the test exhibits 100% sensitivity and 100% specificity on both urine and serum (N=72) samples • No cross reactivity or interference by LH (300 mIU/mL), FSH (1,000 mIU/mL), or TSH (1,000 mIU/mL) Urine / Serum	25T	R-6012-U	CE
Savvycheck Vaginal Yeast Infection Test (POC)	Rapid test for the detection of vaginal yeast infection (candida species) for professional use	LFIA • All features and characeristics as the OTC version (42013) • A box containing 20 tests for professional (OB/GYN) use Vaginal Swab	20T	41013	CE
Savvycheck Vaginal Yeast Infection Test (Self-Testing)	A rapid, easy-to-use, at-home test for the detection of vaginal yeast (<i>Candida spp.</i>) in vaginal discharge	OTC - for self testing of Vaginal Candidiasis: Positive result - self antimycotic treatment (OTC), Negative result - medical consultation • One-step assay • Results obtained in 10 min • Vaginal swab included • >94% sensitivity vs. culture • >98% specificity Vaginal swab	1T	42013	CE ₀₄₈₃
Uriscreen™	Rapid Urinary Tract Infection (UTI) Screen test for the detection of bacteriuria and presence of somatic cells	Easy to Use • Results in 10 sec • CE and FDA (OTC/POC) cleared • CLIA-weaved • LoD = 5X10E4 CFU/mL • Fits POC, OTC, clinical laboratory and veterinary use (large and small animals) • Negative Predictive Value (NPV) >95% Urine	20T	101-01	FDA/CE
SensAheart	Rapid test for detection of hFABP and Troponin	LFIA • Testing for the two cardiac markers (hFABP and Troponin) simultaneously allows for earlier detection of Acute Myocardial Infarction (AMI) • h-FABP LOD 5ng/mL, cTnT LOD 0.5 ng/mL Blood sample	2T	R-6503	CE





Savyon Diagnostics SeroFIA™ Microimmuoassays offers a gold standard detection method for determination of IgG, IgM and IgA antibodies to Chlamydia species.

- Superior clinical and analytical performance
- Proprietary state of the art biological antigens, antibodies and enzyme or fluorophore conjugates
- Convenient slide design

Product	Description	Features Sample Type	Ordering Info & Approvals		
			Tests/ kit	Catalog Number	
SeroFIA	"Semi-quantitative Microimmunofluorescence	The "gold-standard" in Chlamydia serodiagnostics • Species or antibody isotype-specific assays • Results	IgG 105T	511-01 512-01	CE ₀₄₈₃
, (C)	Assay (MIF) for the differential determination of <i>C. pneumoniae</i> , <i>C. trachomatis</i> and <i>C. psittaci</i> specific IgG, IgM or IgA antibodies"	expressed in endpoint titer • Convenient slide design • Excellent correlation with semi-quantitative ELISA tests • 93%/93%/100% agreement with reference commercial MIF for <i>C. trachomatis, C. pneumoniae</i> and <i>C. psittaci,</i> respectively • 99%/97%/100% agreement with reference commercial MIF for IgG, IgM and IgA, respectively Serum	IgA 105T	513-01	





The Savvygen extraction kits are to be used with the Savvygen Extractor instrument to provide high yield and quality nucleic acid.

Features:

- Validated with Savvygen RT-PCR kits
- 22 min fully automated protocol
- Simultaneous extraction of 1-48 samples

Product	Description	Features	Ordering Info & Approvals		
		Sample Type/ Dimensions & Weight	Tests/ kit	Catalog Number	
Savvygen S			96T (4X24)	671-01	CE
Urogenital Extraction Kit	Urine / Urogenital samples	RT-PCR kits (615-01, 617-01, 618-01) • 22 min fully automated protocol for 1 to 48 samples • High DNA yield for downstream PCR-basd applications SAMPLE: Urine / Genital, Vaginal or Cervical Swab / Semen / LBC	48T (6X8)	672-01	
Savvygen S	DNA/RNA Extraction kit for	To be used on the Savvygen Extractor instrument (670-01)	96T (4X24)	677-01	RUO
Respiratory Extraction Kit	Respiratory samples	Compatible for extraction of nucleic acids for Savvygen RT-PCR kits (612-01, 613-01, 623-01) • 22 min fully automated protocol for 1 to 48 samples • High DNA/RNA yield for downstream PCR-basd applications SAMPLE: Plasma, Serum, Nasal swab, Throat swab, Nasal swab, CSF and UTM. For COVID-19 samples	48T (6X8)	678-01	
Savvygen	DNA Extraction kit for	To be used on the Savvygen Extractor instrument (670-01) • 22	96T (4X24)	683-01	RUO
S Genomic DNA	Genomic DNA samples	min fully automated protocol for 1 to 48 samples • High DNA/RNA yield for downstream PCR-basd applications SAMPLE: Whole Blood, Buffy coat swab, Buccal swab and Saliva	48T (6X8)	684-01	
Savvygen™ Stool NA Extraction kit	DNA/RNA extraction kit for isolation of bacteria from stool samples	The kit is compatible with manual workflow or with Hamilton / Tecan automated extraction systems) • High nucleic acid yield & purity) • Load & Run) • Ready to Use) • Fast Extraction-90 min (incl. PCR setup) for 96 preps Compatible with HAMILTON STARIet / Star robots SAMPLE: Stool	96 T	690-01	CE
Savyon NA Lysis Buffer	A Lysis buffer based on Gu-Thiocyanate and Triton X-100 For cell lysis and inactivation of viruses (e.g., SARS-CoV-2) and bacteria	For cell lysis, binding of total nucleic acids and inactivation of nucleases SAMPLE: Nasal swabs, oropharyngeal swabs	100mL	FC800176	
Savvygen Transport &	A Lysis buffer based on Gu-Hcl for use in SARS-	The Transport & Lysis Buffer is intended to be used for inactivation of SARS CoV-2, as well as other respiratory viruses and the	1 tube- 3mL buffer	A800177	
Lysis Buffer	CoV-2 PCR sampling	release of viral nucleic acids. Can be used for direct sampling SAMPLE: Nasal swabs, oropharyngeal swabs	1 tube 10mL buffer	B800177	

Instruments

Product	Description	Features	Dimensions	Oracining into a Appr	
		Sample Type	& Weight	Catalog Number	
Savvygen Extractor (NX-48S)	Automated Ultra-fast (22 min for 1 - 48 samples) DNA/RNA Extractor	Ultra-fast - Up to 48 samples in 22 min • Extremely simple to operate • Very small footprint • Setup and hands-on time <5 min for 48 samples • Magnetic silica microparticles chmistry ensure high yeald and high quality of PCR-ready DNA/RNA • Pre-filled Cartridge type of Reagent– 8 or 24 Extraction Cartridge • UV Lamp/Auto door Lock • 7' Touch Screen based User interface - Easy to operate without training • Pre-programmed and User define protocols	Height: 0.39 m Depth: 0.42 m Width: 0.36 m Weight: 25 kg	670-01	CE





Savyon Diagnostics offers a wide portfolio of Real Time PCR tests. Savyon's RT-PCR SAVVYGEN product line includes a wide range of tests for the detection of Gastrointestinal Pathogens, Sexually Transmitted Infections, Respiratory Tract Infections and Tropical Diseases

- User friendly Setup time < 5min
- Results obtained in 60min
- Full compatibility with most common RT-Thermocyclers
- Compatibility with most manual and automated DNA extraction techniques
- Integrated internal control guarantees appropriate validated results

Product	Description	Features	Ordering Info & Approvals		
		Sample Type	Tests/ kit	Catalog Number	
Savvygen STI CT/NG/TV	Multiplexed RT-PCR for simultanious detection and differentiation of <i>Chlamydia trachomatis (CT)</i> , <i>Neisseria gonorrhoeae (NG)</i> and/or <i>Trichomonas vaginalis (TV)</i> infections	Ready to use master mixes • Common thermocycling profile -can be tested together with other STI panels (615-01, 616-01, 617-01) • As compared with two commercial molecular techniques (RT-PCR and microarray), the tests exhibits sensitivity/specificity levels of 99%/100%, 100%/100% and 100%/100% for CT, NG and TV, respectively (N = 376) Urine / Urogenital Swab	48T	618-01	CE ₀₄₈₃
Savvygen STI MG/MH	Multiplexed RT-PCR for simultanious detection and differentiation of <i>Mycoplasma genitalium (MG)</i> and/or <i>Mycoplasma hominis (MH)</i> infections	Ready to use master mixes • Common thermocycling profile -can be tested together with other STI panels (616-01, 617-01, 618-01) • As compared with two commercial molecular techniques (RT-PCR and microarray), the test exhibits sensitivity/specificity levels of 100%/100% for both MG and MH (N = 241) Urine / Urogenital Swab	48T	615-01	CE
Savvygen STI UU/UP	Multiplexed RT-PCR for simultanious detection and differentiation of <i>Ureaplasma urealyticum (UU)</i> and/or <i>Ureaplasma parvum (UP)</i> infections	Ready to use master mixes • Common thermocycling profile -can be tested together with other STI panels (615-01, 616-01, 618-01) • As compared with two commercial molecular techniques (RT-PCR and microarray), the tests exhibits sensitivity/specificity levels of 100%/100% for both UU and UP (N = 230) Urine / Urogenital Swab	48T	617-01	CE
Savvygen HSV 1+2/VZV	Multiplexed RT-PCR for simultanious detection and differentiation of <i>Herpes</i> Simplex Virus (Types 1 and 2) and Varicella Zoster virus (VZV)	Ready to use liophyllized in-well master mixes • Transport and storage at ambient temperature • LoD = 10 copies • As tested on 5 QCMD panels and 2 INSTAND panels (HSV-1 N=13, HSV-2 N=12, VZV N=14), 100% accordance was found as compared with the validated EQA reports UTM / Genital Swabs / Urine / Plasma	48T	622-01	RUO
Savvygen H. pylori & Antibiotic Resistance	Multiplexed RT-PCR for the identification and differentiation of <i>H. pylori</i> bacteria and a panel of 11 mutations related to its antibiotic resistance to Clarithromycin from stool or biopsy (colony) samples of symptomatic patients.	Ready to use master mixes • Results obtained in 60 min • A first line screening assay for H. pylori & Two antibiotic resistances • Allows personalized antibiotic treatment by the GP • Reduce healthcare expenses due to accurate antibiotic treatment • Reduce doctor appointments and prevention of unnecessary endoscopy procedure • Integrated internal controls guarantee appropriate validated results • Compatible with CFX-96 (Bio-Rad), LC96 (Roche) Stool or Biopcy specimens	96T	621-01	CE





Microbiology tests play a pivotal role in diagnosing infectious diseases and guiding treatment decisions. Accurate interpretation of results informs physicians on appropriate antimicrobial therapy, aiding in patient management and infection control. Savyon offers innovative tools to support and improve the diagnostic processes of microbiology laboratories

- No false positive results
- Simple clean and rapid streaking
- Minimal urine sample required
- · Rapid detection of mixed culture
- Visual diagnosis of UTI pathogens by colony color
- Excellent isolation of colonies needed for subsequent AST analysis

Product	Description	Features Sample Type	Ordering Info & Approvals		
			Tests/ kit	Catalog Number	
ChromoStreak	For transport, screening, isolation and identification of bacterial contamination in urine samples	Culturing Device • Two-sided agar including MacConkey Agar & Chromogenic agar • Streaking device for minimal use of urine • Tube shape device Urine sample	10T	BD-915	CE/ FDA
Dialide	For transport, screening, isolation and identification of bacterial contamination in urine samples	Culturing Device • Two-sided agar including MacConkey Agar & Chromogenic agar • Streaking device for minimal use of urine • Flat shaped device Urine sample	10T	DS-105	CE/ FDA





The quality of the results obtained during the analysis at a clinical laboratory depends to a large extent on the quality of the laboratory reagents used. This is especially important when it comes to ensuring the quality of results and the safety of the technical staff. Savyon offers common high quality laboratory reagents to meet the specific laboratory needs.

- High quality
- Convenient containers
- No spilling
- Barcoded

Product	Description	Features	Ordering Info & Approvals		
		Sample Type	Tests/ kit	Catalog Number	
10% Formalin Cup	Prefilled specimen containers with Neutral Buffered Formalin 10%	High quality formalin solution • Useful for surgical tissues and necropsy specimens • Leakage proof • Prefilled with 20mL or 30ml	120T	BM-603/BM - 601	
50% Medical Ethanol	Prefilled containers with 50% medical Ethanol	Prefilled cups with 40mL 50% Ethanol • Leakage proof • For cytology & general medical purposes • 40mL • Barcoded	ЗТ	800173	
UniSep	Filtered centrifuge tube for density gradient isolation of lymphocytes	Ready to use • Sterile • Maximize yield of mononuclear cells • Prefilled with aseptic solution of 5.7% polysucrose and 9.1% sodium metrizoate	10T	U-05/ U-10	RUO



Savyon Diagnostics has been offering contract manufacturing and development services to biotech companies, varying from startup companies to medium and large enterprises worldwide for the last three decades. Savyon Diagnostics provides high quality products, efficient manufacturing facilities, proficient regulatory affairs services and specialized personnel at all levels. Thus providing the highest level of competence in a cost effective manner.

Savyon Diagnostics offers professional expertise in a variety of fields:

Microbiology • Immunology • Molecular biology • Protein and nucleic acid chemistry • Biochemistry • Glyco-biology

Our assay development services, include, but are not limited to assay development, prototype design, scale-up to production, compilation of SOPs and preparation of technical files for regulatory purposes.

Parameter	Features		
Facilities & Instruments	Environment controlled spaces (Class 10,000 (ISO07)), Dry Room (<20% humidity)		
	Fully automated ELISA microplate coating, blocking, drying and labeling line		
	Fully automated production line for lateral flow tests		
	Automated microplate processor		
	Biosafety level 3 laboratory		
	Automatic dispensing and labeling		
	Validated sterilization process		
Procedures	Sterile filtration (uL to Liters)		
	Bacterial, viral and animal cell culture culture		
	Native and recombinant antigen production		
	Coating of different surfaces		
	Biochemistry and protein purification		
	Downstream molecular biology compartmentalization		
	Top-notch QA & RA services		
Product Design	Packaging		
	Labeling Manuals (print & digital)		
Logistics	Procurement		
g	• Supplies		
	Temperature and humidity-controlled Storage		
	ERP-controlled inventory		
	Worldwide transportation and shipping		
	- Trondmas adireportation and empping		



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The Haptoglobin Typing ELISA (Cat# 710-01) is manufactured under a license from Bio-Rap Technologies Ltd.