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Eppendorf Purity Grades Selection Guide

				eppendorf PCR clean	
				certified purity grade	
	amondar	eppendorf	conceptor	eppendorf	annoniar
	guaranteed	sterilie purity grade	PCR clean	sterile certified purity grade	biopur
		punty grace	purity grade	punty grace	pornty grade
					×
	Eppendorf Quality	Sterile*	PCR clean	PCR clean and sterile*	Biopur*
Continuous quality control for the followin	g relevant criteria				
Function, tightness,precision					
Low wetting					
High chemical resistance					
High thermal resistance					
High resistance to centrifugation forces**					
High transparency			=		
Precisely shaped			=		=
Lot testing (certified) for the following purity cri	iteria				
Human DNA-free					-
DNA-free (Human- + bacterial DNA)					
DNase-free					
RNase-free					
PCR inhibitor-free			-		
ATP-free					
Pyrogen-free (endotoxin-free)					
Sterile (Ph.Eur./USP)				=	
Methods (Examples)					
Methods where high quality consumables are needed without special purities					
Bacteria and yeast culture					
Cell and tissue culture					
Isolation and storage of DNA					
Isolation and storage of RNA					
DNA analysis (PCR, qPCR, restriction analysis, hybridization, microarrays, sequencing)					
Mitochondrial DNA analysis					
Bacterial DNA analysis					
RNA analysis					
Application Areas (Examples)					
Routine Application					
Molecular biology					
Microbiology					
Cell technology					
> Stem cell research					
> Transgenic animals / plants					
Research > Medical Research > Agriculture & Aquaculture Research					
Quality control					
> Food and beverage > Water supply					
> Environmental monitoring					
Forensic					

Recommanded
Highly recommanded
Increased safety due to availability of individually packaged / single-blistered products

** For accurate details regarding resistance to centrifugation, please refer to the product individual instruction for use.

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Importance of Purity Criteria

Sterility

Per definition, a sterile product does not harbor any living organisms on its surface. The degree of sterilization is described by a residual probability of contamination. This probability is expressed as SAL (Sterility Assurance Level). Thus, a SAL value of 10-6 indicates the presence of one non-sterile item among 106 (1,000,000) sterilized items.

Pyrogen-free (endotoxin-free)

Thermostable substances (glycoproteins) from the outer membrane of bacteria and other microorganisms can cause fever in humans and impair the growth of cell cultures.

Bacterial DNA-free (E. coli)

DNA is found in all cells of living entities, and it is the carrier of all genetic information. The highly sensitive PCR technique enables the detection of individual molecules.

Human DNA-free

To eliminate this potential source of contamination, the consumables are tested for the presence of human DNA. Even a single cell (e.g. skin particles) would be detected in the test. Manufacturing is virtually fully automated and monitored by staff wearing protective clothing.

DNase-free

DNases are enzymes which degrade DNA.

RNase-free

RNases are enzymes that degrade RNA. These enzymes are extremely resistant, even to autoclaving and irradiation.

ATP-free

ATP is a part of all living cells; therefore, its presence can indicate biological contamination.

PCR inhibitor-free

PCR—the replication of DNA—has developed into one of the most important and commonplace molecular biology methods used in medical diagnostics, genetic counseling and all basic biological research. However, there are also substances that impair this reaction, so lab products must be free of these inhibitors.

Importance

Sterile products are required whenever the presence of germs may have a negative effect; for example, in the prevention of infection of samples or incorrect test results for microbiological experiments that would be caused by unsterile lab equipment.

Importance

Absence of pyrogen prevents endotoxin-based contamination in drug manufacture, cell culture and medical laboratories.

Importance

The presence of DNA could lead to falsepositive results for different applications involving DNA. Note: Autoclaving does not remove traces of DNA.

Importance

Tests for human DNA prevent consumables from containing DNA that could lead to falsepositive results (e.g. genetic tests in forensics)

Importance

DNase contaminations can influence DNA analysis.

Importance

RNase-free products are an absolute must in the field of molecular biology because RNA is highly sensitive and can be destroyed very quickly by RNases.

Importance

The test procedure for the quantitative and qualitative detection of ATP is already an integral part of hygiene monitoring, e.g. in the pharmaceutical industry.

Importance

It is essential that consumables used contain no impurities that could adversely affect PCR. This is particularly crucial for the amplification of minute quantities of genetic substances and for quantitative PCRs.

Your local distributor: www.eppendorf.com/contact Eppendorf AG · 22331 Hamburg · Germany eppendorf@eppendorf.com · www.eppendorf.com

www.eppendorf.com/consumables

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