

# UriSed Cuvettes

## REF

URS-9961HU  
URS-9961-1  
URS-9971  
URS-9972  
URS-9974  
URS-9961CH-1  
URS-9971CH

## Instructions for use

### Intended use:

UriSed Cuvettes are disposable, single use polycarbonate specimen receptacles used to analyze uncentrifuged, human urine samples with UriSed sediment analyzers. It is intended for professional, laboratory use. It is intended for in vitro diagnostic use.

### Test principle:

UriSed Cuvettes are specimen receptacles allowing for microscopic analysis of urine or body fluid samples.

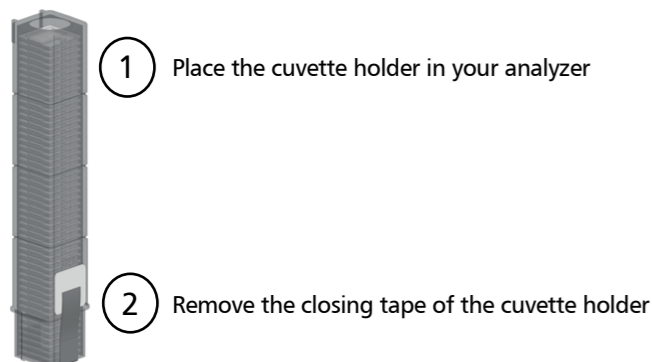
### Materials provided:

600 cuvettes in 12 cuvette holders of 50 cuvettes each

### Materials not provided:

- Compatible urine sediment analyzer (UriSed, UriSed2, UriSed 3, UriSed 3 PRO, UriSed mini)
- General laboratory equipment

### Using cuvettes:




## Environmental Conditions

Storage temperature	0 – 45°C
Transport temperature	-25°C – 60°C
Transport humidity	20 – 80 %
Operation conditions	According to your analyzer's conditions

## Warnings and cautions










- Do not store cuvettes in direct sunlight
- Do not remove closing tape from the cuvette holder before installing in your analyzer
- Do not remove partially full cuvette holders from your analyzer
- Each cuvette is single use, never perform a test with previously used cuvette
- Since urine is a fluid of human origin, it may be infectious and may constitute a potential biological risk
- Handle used UriSed cuvettes and urine contaminants with care
- Dispose of waste according to accepted laboratory instructions and procedures.
- Contact your distributor to make sure to order cuvettes compatible with your specific analyzer
- Use cuvettes before expiration date

 Check your analyzer's instructions for use for details on specimen collection, potential preparatory steps, result calculation, analytical and performance characteristics, interferences, limitations, quality control procedures, specific warnings and cautions.

## Incident reporting

Report any serious incidents which may occur when using this product to your 77 Elektronika service representative and your local competent authority.

## Symbols:

	Unique Device Identifier
	In vitro diagnostic medical device
	Catalogue Number
	Lot Number
	The CE mark identifies that the product complies with the applicable directives of the European Union
	Use by
	Temperature Limitation
	Manufacturer
	Keep away from sunlight
	Consult instructions for use
	Humidity Limitation
	Caution
	Biological Risks
	600 Contents sufficient for 600 tests
	Do NOT Reuse
	Do not use if package is damaged
	English Language
	Batch number

## Version history

Version	Date	Changes
3	2022.03.25	IVDR compliance update
2	2020.11.25	General update of content
1	2009.02.17	First release

## Manufacturer:

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