



PF-PTD-250

Potassium	7880 Accredited to ISO 15189:2012
Synonyms	К+
Clinical Indication	Diagnosis of hypokalaemia and hyperkalaemia or monitoring patients at risk of developing hypokalaemia or hyperkalaemia.
Part of Profile / See Also	Urea and Electrolytes (UE)
Request Form	Combined Pathology manual Blood form or ICE request
Availability / Frequency of	On request
Analysis	
Turnaround Time	Same day
Patient Preparation	
Sample Requirements	
Specimen Type	Serum and plasma
Volume	2 ml
Acceptable Containers	Yellow top (SST) tube
	Green top (lithium-heparin) tube
	paediatric orange top (lithium-heparin)
	paediatric green top (lithium-heparin)

Plain serum samples may also be used.

Reference Range & Units	Age	Potassium (mmol/L)			
	Neonate (< 1 month)	3.4 - 6.0			
	1 - 12 months	3.5 - 5.7			
	1 - 16 years	3.5 - 5.0			
	Adult	3.5 - 5.3			
	Plasma potassium levels are slightly lower than serum potassium levels Reference: Pathology Harmony Group, Clinical Biochemistry Outcomes, January 2011 (www. pathologyharmony.co.uk)				
Interferences	cause an artefactual increas	delay in separation of serum from rec ie in potassium concentration. In pati im is released from platelets leading	ents with		



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Interpretation	&	Clinical	
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Decision Value (if applicable)

## References

Test code

Lab Handling

Critical Difference 14%

Beckman kit insert.

## UE

Analysed from primary tube and stored at 4°C. Sample must be separated within 6 hours. Serum and plasma stable for 7 days at 2-25°C.