





InSight V-CHEM Quick Reference Guide

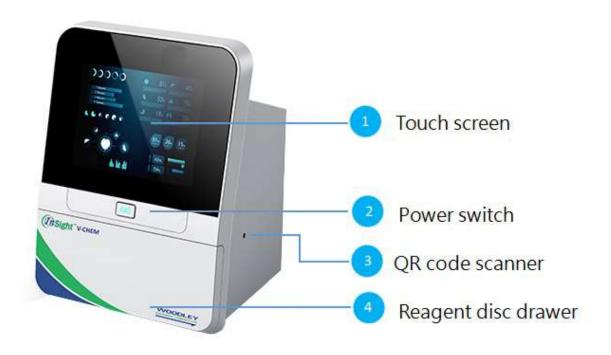


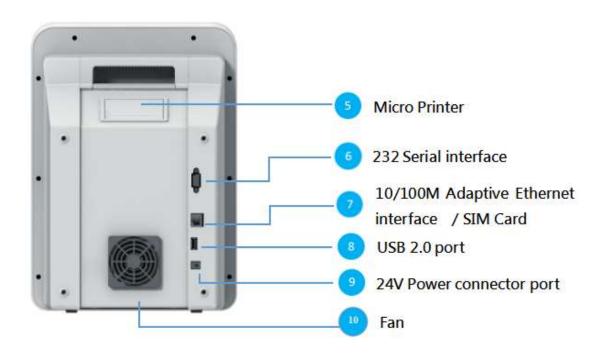
Quick Reference Guide Contents

External features of the analyser	3
InSight V-CHEM Function Buttons	4
Testing a Sample	5
Interpreting the results	6
Accessing previous results	7
Date and Time setting	8
Troubleshooting	9
InSight V-CHEM Rotors	11



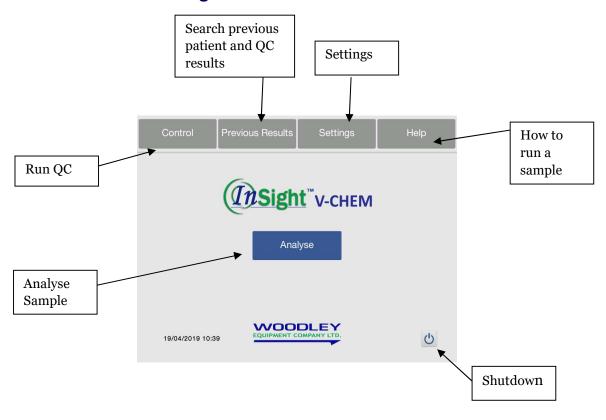
External features of the analyser







InSight V-CHEM Function Buttons



Warning – To avoid scratching the screen do not use sharp or hard objects to touch the screen



Testing a sample

Sample Preparation

- Sample type Serum, Lithium-Heparinised Whole blood or Plasma
- Sample Volume 100ul per test panel
- Testing must be conducted within 60 minutes (at room temperature)
- To prevent haemolysis of the sample, do not refrigerate, freeze, or shake whole-blood sample.
- Plasma or serum can be stored at Room Temp for 5 hours prior to testing or up to 48 hours if stored at 2-8C

Rotor Preparation

- Rotors should be stored at 2-8C and can be used straight from the refrigerator without the need for warming
- Avoid placing unopened reagent discs in temperatures greater than 25C (77F) for more than 48 hours.
- 1. Before using a reagent disc, carefully check the foil pouch for any damage. Tear open the package from the upper right side of the pouch.
- 2. Remove the reagent disc gently with finger and thumb
- 3. After removal use your other hand to hold the disc by the edge. Avoid contact with disc surface where optical measurement takes place.
- 4. Check the reagent disc for any damage do not use reagent discs that have been dropped or damaged.
- 5. After opening the pouch, the disc must be used within 20 minutes. Do not place the disc back in the refrigerator for later use.
- 6. Remove the foil tab from the rotor to release the diluent by pulling the tab at a 45° angle.

Dispensing the sample

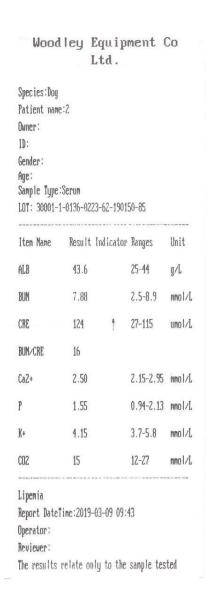
- 1. Attach a new pipette tip to the end of the 100ul pipette provided. Do not touch the end of the tip.
- 2. Using your thumb push down the plunger on the top of the pipette to the stop position.
- 3. Immerse the pipette tip below the surface of the sample.
- 4. Slowly release the plunger to draw up the sample, avoid drawing in any air, as this will cause the disc to fail.
- 5. Remove the pipette from the sample tube.
- 6. Insert the pipette tip into the sample port (indicated by a red triangle) on the rotor
- 7. Keep the reagent disc level and hold the pipette at a 45-degree angle to the surface of the disc.
- 8. Gently push down the plunger at the top of the pipette to the stop position. All of the sample should have been expelled into the sample chamber. Keeping the plunger held down gently remove the pipette tip from the sample port. The plunger may now be released, remove the disposable tip from the pipette and discard the tip.
- 9. Press the **Analyse** icon on the touch screen
- 10. Select the sample type (whole blood, serum or plasma)
- 11. When indicated on the screen, Scan the QR code on the foil pack that the rotor came in
- 12. Hold a reagent disc that contains a sample by its edge and the disc kept level. Avoid touching the surface of the reagent disc. Place the disc into the drawer until it clicks and gently remove the blue film, press **Close** to close the disc drawer.
- 13. Press the 'Yes' icon to confirm the blue film has been removed.
- 14. Use the touch screen to enter the species and Patient ID
- 15. The fully automated analysis will now commence once the disc drawer closes
- 16. When the analysis is completed, the system will store the analysis results and display them on the screen.

To cancel a test in progress, press the X icon on the screen, the system will prompt for confirmation. A reagent disc cannot be re used if a test has been cancelled.





Interpreting the results



A typical report printout is shown above. The heading of a report printout includes information such as Sample Patient Name, Owner Name, ID Number, Gender, Age, Sample Type and rotor Lot Number. The test results section is printed in five columns: Analyte Name, Analyte Result, Indicator, Reference Range and Specified Units. The end of the report printout includes information such as the sample indices, test date and time and blank areas for the operator and reviewer to sign.





Accessing Previous Results

The InSight V-CHEM automatically stores tests results for display, printing or Export to a computer.

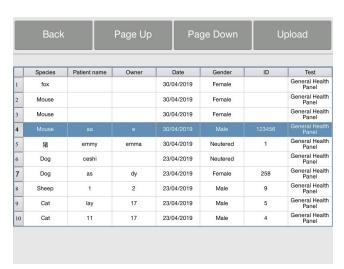
To access previous results: From the Home Screen > press 'Previous Results' then select patients.



Enter the patient ID or time range to search for the result.



The analyser will display all results as filtered, use the up and down arrows to scroll through the list.



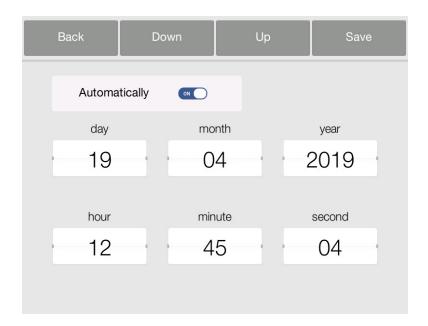


Date and Time setting

The system date and time will usually be set before installation and the system remembers the date and time between uses.

To change the date and time, from the home screen select 'Settings' then press 'Date/Time'.

Use the up and down arrows to change the day, month, year, hour, minute and seconds. Press 'Save' once correct date and time has been set.





Troubleshooting

When there is a problem with the analyser, a warning and error message will be shown.

Error Code	Problem Description	Solution
0101	Multi-switch fault	
0102	+12V power fault	
0103	-12V power fault	
0105	AD (AD fault)	
0107	Optical module fault	
0108	LED fault	
0109	Optical output is unstable	
0202	Drawer open fault	Plana contact
0203	Drawer close fault	Please contactWoodley Equipment
0206	Motor speed fault	or your authorised
0213	Temperature fault	dealer.
0214	Insufficient lamp exposure	
0215	Dark current	
0216	Light intensity saturation	
0220	Temperature data transmission fault	Refer to the Maintenance Manual
0221	PT100 fault	— (Authorised dealers — only)
0222	Upper NTC fault	
0223	Lower NTC fault	
0224	Upper heating film fault	
0225	Lower heating film fault	
0301	FPGA fault	
0302	Scan module fault	
0303	606 board fault	



Error Codes for Reagent Disc and Troubleshooting

Error Code	Problem Description	Solution
0208	Insufficient sample	According to Section 3.3 ,
0209	Insufficient diluent	repeat the analysis with a new
0210	Sample and diluent not mixed correctly	reagent disc
0211	Sample interference, severe haemolysis, severe lipaemia or severe jaundice	Confirm sample quality and repeat the analysis with a new reagent disc. If symptom persists, use an alternative testing method or send the sample to a reference laboratory.
0212	Disc expired	Please use the reagent disc within the validity period.



InSight V-CHEM Rotors



Analytes Profile	Items	ALP	ALT	AST BUN	_	CREA	9	ALB	£	TBIL	AMY	3	PHOS	·e	<u>.</u>	ь	BA	GGT	ž	CHOL	tco;	Others
General Health Panel	13	ALP	ALT		BUN	CREA	GLU	ALB.	TP	TBIL	AMY	S	PHOS		П				X	CHOL	П	
Preanaesthetic Panel	9	ALP	ALT	1	BUN	CREA	OID	П	Д	П			П	П	Н	Н		Н			П	
Preanaesthetic Plus Panel	6	ALP	ALT	AST	BUN	CREA	CLU		ТР	П	Н	П	П	П	Н	Н			×	П	П	HO1
Liver & Renal Panel	6		ALT	AST	BUN	CREA	OIO	ALB	ТР	TBIL	Н	П	П	П	Н	Н		GGT	Н	П	П	
Liver Function Panel	00	ALP	ALT	AST				ALB.	ТР	TBIL		П	П	П	Н	Н		GGT		П	П	DBIL
Renal Function Panel	7				BUN	CREA		ALB				3	PHOS		<u>*</u>						tCO2	
Electrolytes Panel	7											S	PHOS	· eN	<u>*</u>	.o				-	tCO2	Mg
Critical Care Panel	00		ALT		BUN	CREA	OID							Na.	¥	Ö.					1002	
Ammonia Test	,-						П	П	П							П	П		П	П		NH ₃
Avian & Reptile Panel	13		ALT	AST			Offi	ALB	4			3	PHOS	*e	÷	ō.	A8		×			UA
Large Animal Panel	10	ALP		AST	BUN			ALB	4			S	PHOS					GGT	X		П	Mg
GLU & Lipid Panel	S						CIO								Н					CHOC		TG, HDU-C, HCY
Triple Tests Panel	12	ALP	ALT		BUN	CREA	П	ALB	Д	TBIL	AMY	S	PHOS	Н	Н	Н	Н		×	CHOL	П	