

AmiShield® Kidney Plus Panel

For Veterinary Use Only

For Professional Use Only

Catalog Number : 001-37GI (For VCA-TE-100)

001-37GK (For VCA-TE-300)

-----Please follow the instructions before use-----

Intended use

The disposable AmiShield® Kidney Plus Panel in conjunction with the AmiShield® Veterinary Clinical Analyzer utilizes dry and liquid reagents to provide quantitative determinations of albumin (ALB), blood urea nitrogen (BUN), creatinine (CRE), glucose (GLU), phosphorus (PHOS), total calcium (CA). At the same time through the calculation, can get the other one parameter (BUN/CRE) information. After each test, biochemical test results can be obtained 7 to provide a basis for rapid diagnosis in lithium heparinized whole blood, lithium heparinized plasma or serum.

Clinical Significance

The disposable AmiShield® Kidney Plus Panel and the AmiShield® Veterinary Clinical Analyzer assist the veterinarian in diagnosing the following disorders:

Albumin (ALB): Hepatic and kidney diseases

Blood urea nitrogen (BUN): Hepatic and kidney diseases

Creatinine (CRE): Renal disease

Glucose (GLU): Diabetes, hyperglycemia, hypoglycemia, hepatic disease

Phosphorus (PHOS): Nephrotic disease, hypoparathyroidism and nutritional disorders

Total calcium (CA): Parathyroid, bone and chronic renal disease ; tetany

Urea Nitrogen/Creatinine (BUN / CRE) (calculated): Kidney disease

As with any diagnostic test procedure, the clinical samples or other test procedures should be considered prior to final diagnosis

Principles of Procedures

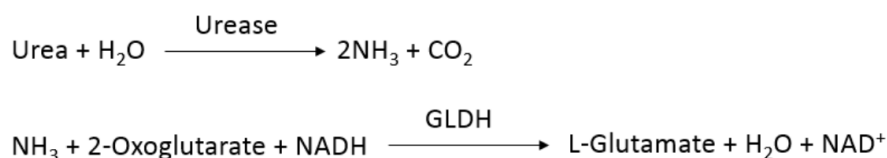
(1) Albumin (ALB)

Dye-binding method is used to detect the quantity of albumin. Albumin is bound by the Bromocresol green (BCG) dye to produce an increase in the blue-green color measured at 620 nm. The color increase is proportional to the concentration of albumin present.



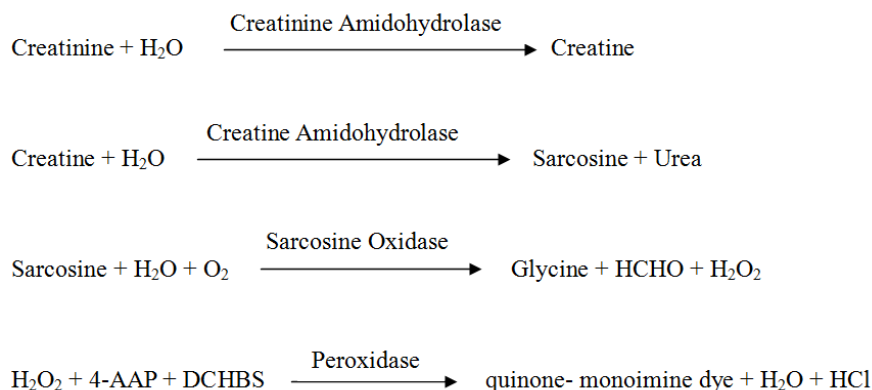
(2) Urea Nitrogen (BUN)

Urea is hydrolyzed by urease to produce ammonia and carbon dioxide. The liberated ammonia reacts with α -ketoglutarate and NADH in the presence of enzyme glutamate dehydrogenase to yield glutamate and NAD. NADH undergoes oxidation during the reaction resulting in a decrease in absorbance at 340 nm that is directly proportional to the urea nitrogen concentration in the sample.



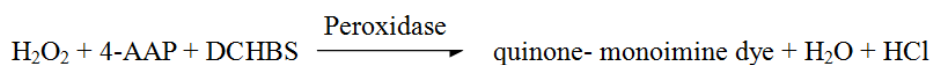
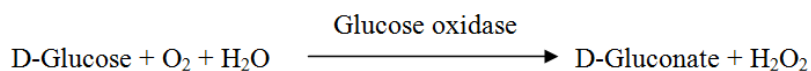
(3) Creatinine (CRE)

Creatinine is catalyzed to creatine by creatinine amidohydrolase. Under the catalyzation of creatine amidohydrolase, sarcosine and urea is produced from creatine. Sarcosine is oxidized to glycine and hydrogen peroxide (H_2O_2) in the presence of sarcosine oxidase. Hydrogen peroxide reacts with the substrates 3,5-Dichloro-2-hydroxybenzenesulfonic acid (DCHBS) and 4-aminoantipyrine (4-AAP) to form the color complex quinone- monoimine dye that absorbs at 510 nm. The rate of formation of color is proportional to the creatinine in the sample. Potassium ferrocyanide and ascorbate oxidase are added to the reaction to minimize the interference from bilirubin and ascorbic acid.



(4) Glucose (GLU)

Glucose is oxidized by glucose oxidase to gluconate and hydrogen peroxide. DCHBS, 4-AAP and hydrogen peroxide, in the presence of peroxidase, produces a quinoneimine dye that is measured at 510 nm. The absorbance at 510 nm is proportional to the concentration of glucose in the sample. Potassium ferrocyanide and ascorbate oxidase are added to the reaction to minimize the interference from bilirubin and ascorbic acid.



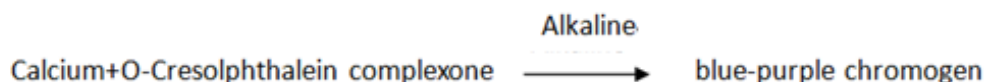
(5) Inorganic Phosphorus (PHOS)

Inorganic phosphorus reacts with ammonium molybdate in a strongly acid solution to form a phosphomolybdate complex that absorbs light at 340 nm. The absorbance at this wavelength is directly proportional to the amount of inorganic phosphorus present in the sample.



(6) Total calcium (CA)

Total calcium is coupled with o-Cresolphthalein complexone (Phthalein purple) in an alkaline solution to form a blue-purple color complex. The amount of Calcium in the sample is proportional to the absorbance. For calcium determination, 8-hydroxyquinoline was added to the reagent to eliminate the interference of magnesium ions in the sample.



Storage

1. Store the rotor that sealed in their foil pouches at 2 – 8 °C (36 – 46 °F). When stored as described above, all reagents in the rotor are stable until the expiration date which printed on the rotor foil pouch.
2. Do not expose opened or unopened rotor to direct sunlight or temperatures above 30 °C (86 °F).
3. Do not use a rotor after the expiration date.
4. Do not use a rotor from a damaged foil pouch. Because, a torn or otherwise damaged foil pouch may lead moisture to reach the unused rotor and adversely affect reagent performance.

Materials Required but not Provided

1. AmiShield® Veterinary Clinical Analyzer
2. Sample collector

3. Pipette and tip
4. Controls

Instructions for Reagent Handling

1. The rotor should be used for assay immediately following take out from refrigerator.
2. Open the sealed foil pouch and remove carefully the rotor. Don't touch the barcode located on the top of the rotor. The contaminated or scratched barcode will not be scanned by analyzer.
3. The rotor should be used within 20 minutes after opening the pouch. The rotor in opened pouches can't be placed back into the refrigerator for reuse.
4. The rotor would be firmly pressed onto the spindle of AmiShield® Veterinary Clinical Analyzer.
5. Transfer 0.14 mL (140 µL) sample to rotor inlet through the sample port by pipette.
6. **Use only lithium heparinized whole blood, plasma or serum.**
7. The analyzer maintains the rotor at a temperature of 37 °C over the measurement interval. The analysis time is about 13-15 minutes. In addition, the AmiShield® System operates at ambient temperatures between 15°C and 30°C.

Sample Collection and Preparation

1. The minimum required sample size is 0.14 mL (140 µL) of lithium heparinized whole blood/plasma/serum or control.
2. Use no-additive (red stopper) evacuated specimen collection tubes or serum separator tubes (yellow or red/yellow stopper) for serum samples. Use only lithium heparin (green stopper) evacuated specimen collection tubes for whole blood/plasma samples.
3. Whole blood samples obtained by venipuncture must be homogenous. Gently invert the collection tubes several times just prior to sample transfer. Do not shake the collection tube. Shaking may cause hemolysis.
4. Release both the needle of syringe and the stopper of collection tube before transferring whole blood sample to collection tube.
5. The test must be started once sample is transferred into the rotor. A long delay time may affect the analytical performance.
6. Lithium heparinized whole blood samples should be run within 60 minutes of collection; if this is not possible, separate the sample and transfer it into a clean test tube. Run the separated plasma or serum sample within 5 hours of centrifugation. If this is not possible, refrigerate the sample in a stoppered test tube at 2 – 8 °C (36 – 46 °F) for no longer than 48 hours.

Precautions

- Wear a laboratory coat and gloves to avoid the biohazard and puncture injury.
- The medical waste should be disposed following the local regulations.
- See the AmiShield® Veterinary Clinical Analyzer Operator's Manual for complete information on using the analyzer.

Warnings

1. When rotor embeds onto the spindle the diluent container is open. A rotor with an opened diluent container can't be reused. Ensure that the sample or control has been placed into the rotor before running the test.
2. The AmiShield® products used only with the AmiShield® Veterinary Clinical Analyzer, vice versa. Before START the test, please confirm the rotor is properly and evenly embedded into the spindle, in addition, the assembled holder should be well placed on the spindle in the Analyzer.
3. Please avoid colliding or falling damages. In this case, the rotor can't be used.
4. Reagents in the rotor may contain acids or caustic substances. The operator does not come into contact with the reagents when following the recommended procedures. In the event that the reagents are handled (e.g., cleaning up after dropping and cracking a rotor), avoid ingestion, skin contact, or inhalation of the reagents.
5. Some reagents contain sodium azide, which may react with lead and copper plumbing to form highly explosive metal azides. Reagents will not come into contact with lead and copper plumbing when following recommended procedures. However, if the reagents do come into contact with such plumbing, flush with a large volume of water to prevent azide buildup.

Quality Control and Calibration

1. The AmiShield® Veterinary Clinical Analyzer is calibrated by the manufacturer before shipment.
2. The barcode printed on the upper cover provides the analyzer with rotor-specific calibration data.
3. Controls may be run periodically on the AmiShield® Veterinary Clinical Analyzer to verify the accuracy of the analyzer by user.
4. A control is only available from producer. Run controls on the rotor in the same manner as for patient samples. See the AmiShield® Veterinary Clinical Analyzer Operator's Manual to run controls.
5. The QA/QC should be conducted following the local regulations or the laboratory guideline.

Known Interference Substances

1. The only anticoagulant recommended for the AmiShield® Veterinary Clinical Analyzer is lithium heparin. Sodium heparin must not be used when collecting blood sample for use with this rotor. The EDTA, fluoride, oxalate, and any anticoagulant containing ammonium ions will interfere with at least one chemistry in the AmiShield® Kidney Plus Panel.
2. Physical interferents (hemolysis, icterus, and lipemia) may cause changes in the reported concentrations of some analytes. The sample indices are printed on the bottom of each result card to inform the operator about the levels of interferents present in each sample.
3. Bilirubin may interfere with the peroxidase used in the creatinine reaction. Creatinine results are lowered when bilirubin levels are > 10 mg/dL.
4. Glucose concentrations are affected by the length of time since the patient has eaten and by the type of sample collected from the patient. To accurately interpret glucose results, samples should be obtained from a patient that has been fasted for at least 12 hours.
5. Interference may be seen in the total protein test when analyzing samples with a lipemic index. Samples with a triglyceride concentration >400 mg/dL may show an increased total protein level.
6. Hematocrit (Hct) > 60% of whole blood samples may affect the test results.
7. Hemolysis samples will release some substrate to the serum or plasma, may cause AST, LDH, K and other biochemical markers increased significantly, ALT, UA, CHOL, ALP, CK light to moderate the degree of increase. Severe hemolysis samples recommended not to use.

Reference Intervals

These normal intervals are provided only as a guideline. The most definitive reference intervals are established for your patient population. Test results should be interpreted in conjunction with the patient's clinical signs.

Analyte		Common Units		SI Units	
ALB	Canine	2.4–3.9	g/dL	24–39	g/L
	Feline	2.2–4.0	g/dL	22–40	g/L
BUN	Canine	6–27	mg/dL	2.14–9.64	mmol/L
	Feline	6–36	mg/dL	2.14–12.85	mmol/L
CRE	Canine	0.3–1.5	mg/dL	27–133	µmol/L
	Feline	0.3–2.4	mg/dL	27–212	µmol/L
GLU	Canine	70–140	mg/dL	3.89–7.77	mmol/L
	Feline	75–166	mg/dL	4.2–9.2	mmol/L

PHOS	Canine	2.1–6.5	mg/dL	0.68–2.1	mmol/L
	Feline	3.1–7.5	mg/dL	1–2.42	mmol/L
CA	Canine	7.9-12.0	mg/dL	1.97-2.99	mmol/L
	Feline	7.8-11.3	mg/dL	1.95-2.82	mmol/L

Dynamic range

The chemistry for each analyte is linear over the dynamic range listed below. The intervals below do not represent normal ranges.

Analyte	Common Units		SI Units	
ALB	1.0–5.0	g/dL	10.0–50.0	g/L
BUN	2–200	mg/dL	0.71–71.4	mmol/L
CRE	0.1–20.0	mg/dL	9–1768	μmol/L
GLU	10–500	mg/dL	0.56–27.7	mmol/L
PHOS	2.0–15.0	mg/dL	0.68–4.85	mmol/L
CA	4.0 – 40.0	mg/dL	1.00 – 9.98	mmol/L

Method Comparison

Field studies were conducted at a veterinary teaching hospital. The same serum samples were analyzed by the AmiShield® Veterinary Clinical Analyzer and a comparative method. Representative correlation statistics are shown in below.

Analyte	Correlation Coefficient	Slope	Intercept	Sample No.	Sample Range	
ALB	0.84	0.53	1.36	83	1.5-4	g/dL
BUN	0.97	0.89	3.34	83	8-124	mg/dL
CRE	0.97	0.92	0.09	83	0.4-9.1	mg/dL
GLU	0.98	1.01	-3.23	83	77-434	mg/dL
PHOS	0.93	0.87	0.52	83	2.2-10.5	mg/dL
CA	0.95	0.69	3.43	86	5.3-11.6	mg/dL

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Symbols



Consult Instructions for use



Caution



Temperature Limitation



Reference Number



Batch code



Manufacturer



Use by



Do Not Reuse

Manufacturer : ProtectLife international Biomedical Inc.
Address : 4F., No.8, Xinghua Rd., Taoyuan Dist., Taoyuan City 33068, Taiwan
Customer and Technical Service : 886 3 3775599
Official Website : www.protectlife-intl.com