# Limitations of Point-of-Care Testing



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# **Objectives:** Monitoring

- Understand the purpose and limitations of point-of-care testing
   Understand which monitors are useful for quick information gathering and what the information means
- Understand the limitations of the monitoring equipment

# **Point-of-Care Testing**

- Purpose: To obtain immediate (within 5-10 minutes), cage-side patient information that will compliment findings from the initial triage examination
- When: Initial presentation and dynamic patients
- Focus: Life-threatening problems
- Limitations: Know your machines!

#### **Recognizing Life-Threatening Problems: Triage and Point-of-Care Testing**

- Shock
  - Hypovolemic
  - Distributive
  - Cardiogenic
- Sepsis/SIRS/MODS
- Respiratory distress
- Cardiac disease
- Diabetic Ketoacidosis
- Addison's Disease
- Hypoglycemia
- Hypo/hypertension

- Acute abdomen
- Seizuring
- Head trauma
- Blood disorders
- Electrolyte abnormalities
- Renal failure
- Post-renal obstruction
- Toxins
- Acid-base disorders
- Saddle Thrombus

#### The Extended Data Base

#### PCV/TP

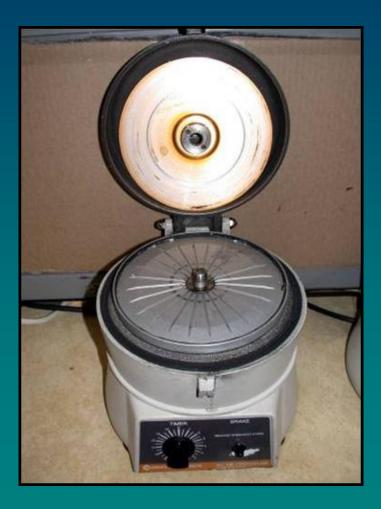
- Glucose
- BUN or Creatinine
- Electrolytes
- Lactate
- VBG or ABG

- Urine s.g. & dipstick
- Blood smear
- Coagulation status
- ECG
- Blood pressure
- Pulse oximeter

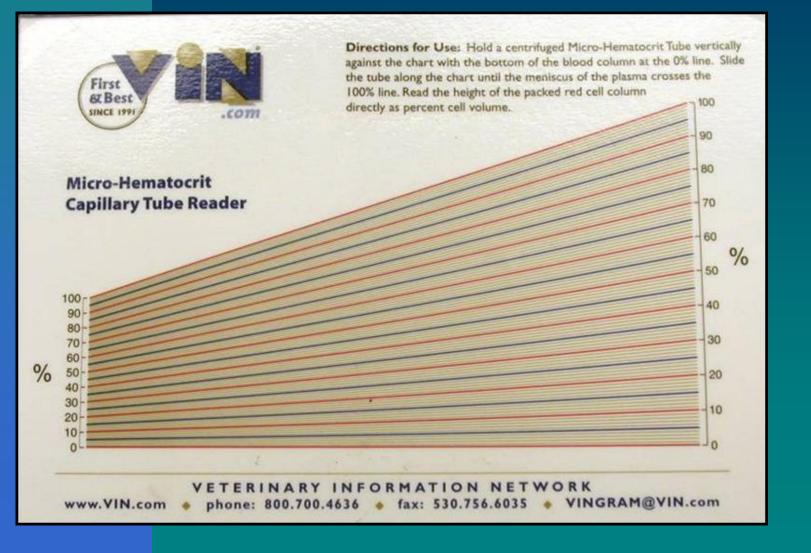
# Packed Cell Volume

 Percentage of whole blood that contains RBC's compared to plasma
 Centrifuged in

- hematocrit tubes
- Measured on chart

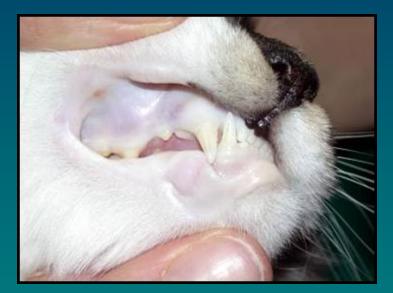


# Packed Cell Volume



# Packed Cell Volume - Anemia

- Loss, destruction, decreased production, sequestration
- During hemorrhage, TP will drop before PCV due to splenic contraction
- Plasma color and TP can help differentiate causes of anemia
- Look at how the patient is handling the anemia



#### Packed Cell Volume - Polycythemia

Hemoconcentration
 Chronic hypoxia
 Primary polycythemic disorder

# **Packed Cell Volume Limitations**

- Diluted samples are common, especially if sample was from a catheter
- Centrifuge speed and length of time can alter results
- Minor subjective errors in reading
- May be excessively hemodilute if recently received lots of fluids
- Blood clots can alter results

# **Total Protein**

Measured on refractometer
 Can be artificially elevated with lipemia





## **Total Protein**

Hypoproteinemia

Loss (GI, hemorrhage, or renal)
Decreased production (liver)

Hyperproteinemia

Dehydration
Hyperglobulinemic

#### **Total Protein Limitations**

- Can't differentiate between panhypoproteinemia vs. hypoalbuminemia
- Can't differentiate panhyperproteinemia and hyperglobulinemia
- Lipemia, icterus, and hemolysis can artificially raise TP

# **PCV/TP** Case Examples

PCV = 45%; TP = 7.5 g/dL - Normal PCV = 55%; TP = 8.5 g/dL - Dehydration PCV = 40%; TP = 4.0 g/dL - Early hemorrhage - Hypoproteinemia PCV = 25%; TP = 4.0 g/dL Late hemorrhage

#### **PCV/TP** Case Examples

PCV = 68%; TP = 5.2 g/dL - Hemorrhagic gastroenteritis PCV = 20%; TP = 7.5 g/dL - RBC destruction or lack of production – Look at plasma color! PCV = 70%; TP = 7.0 g/dL - Polycythemia PCV = 45%; TP = 11 g/dL- Hyperglobulinemia Lipemia

#### Glucose

- Measured on glucometer or blood gas machine
- Glucometers are designed for human diabetics
  - Read approximately 20 mg/dL low during hypoglycemia



#### Glucose

Look at your patient! - Clinical signs? - Pediatric vs. adult? - Patient size/species? Seizuring doesn't typically occur unless below 40 mg/dL Detrimental effects of glucose supplementation?

#### Glucose

#### Hypoglycemia

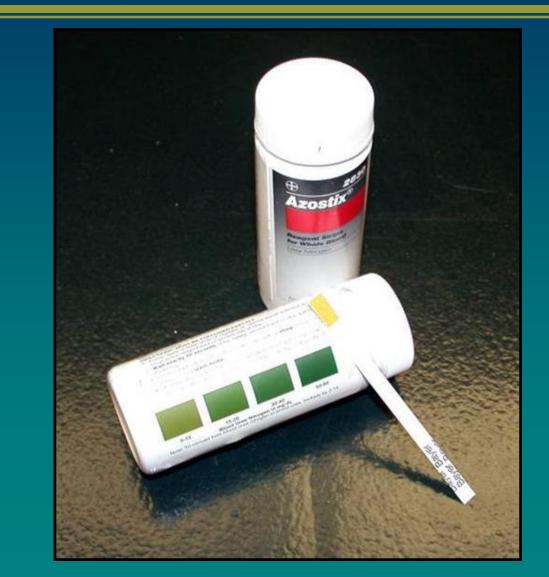
- Lab error
- Sepsis/SIRS
- Insulinoma
- Starvation
- Pediatric
- Hyperglycemia
  - Stress (cats)
  - Diabetes mellitus
  - Massive epinephrine release
  - Head trauma



#### **Renal Values**

BUN
May be elevated with Gl hemorrhage
Creatinine
More renal specific than BUN
Pre-renal vs. renal vs. post-renal

# Measuring BUN - Azostix



#### **Urine Specific Gravity and Dipstick**

- Pre-fluids s.g. is most useful
- Be comfortable with no coag abnormalities prior to cystocentesis
- Don't withhold fluids just to get s.g.
- Plasma can be used in lieu of urine to look for ketones
- Leukocyte/nitrite not accurate on dipstick in animals



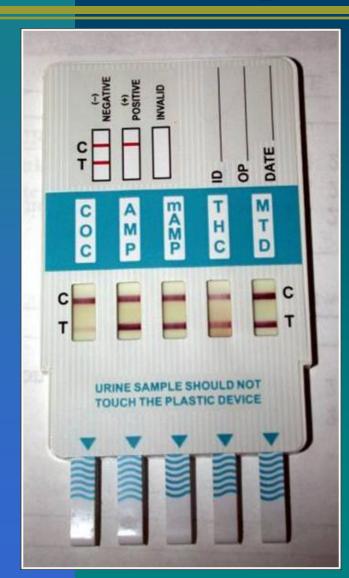


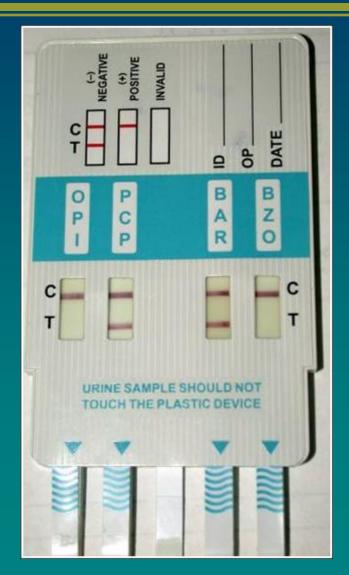
# **BUN/Specific Gravity Examples**

**BUN** = 100 mg/dL, Creatinine = 4.0 mg/dL, Urine sg = 1.040Prerenal azotemia **BUN** = 100 mg/dL, Creatinine = 4.0 mg/dL, **Urine sg = 1.010** – Renal azotemia – Prerenal azotemia? BUN = 100 mg/dL, Creatinine = 1.5 mg/dL, Urine sg = 1.025- GI hemorrhage



# Urine Drug Testing





#### Electrolytes

Sodium Potassium Chloride Ionized calcium - Reads artificially low if sample over-heparinized - Acidosis increases ionized calcium

Glu	60	mg/dL
Na	135	mmol/
К	>9.0	mmol/
TC02	8	mmol/
iCa		mmol/
Hct	37	%PCV
нь*	13	9/dL
*via	Hct	

#### Electrolytes

Make sure samples from catheters are not contaminated with fluids or additives Hct/Hb on blood gas machines usually not accurate

		20.0
Glu	60	mg/dL
Na	135	mmol/
К	>9.0	mmol/
TC02	8	mmol/
iCa		mmol/
Hct	37	%PCV
нь*	13	9/dL
*via	Hct	

#### Lactate

- A measure of perfusion
- Predictor of gastric necrosis in GDV if > 6 mmol/L
- How quickly lactate drops may be more important than absolute numbers
- Great tool to assess response to resuscitation



#### Venous or Arterial Blood Gas



At 37C	
PH6.927	
PC0235.4	mm
P0268	i mmi
HC037	mm
BEecf25	i mm
s02*78	%
*calculated	1

## Venous vs. Arterial Blood Gases

- Venous blood gases are reliable for acidbase status and ventilatory status
- VBG gives a more accurate picture of global acid-base status during CPR
- pH < 7.2 leads to cellular dysfunction</p>
- Only arterial blood gases can tell you how well a patient is oxygenating
- Venous oxygen tensions are only useful for calculating oxygen extraction
- Venous oxygen < 20 mmHg has been associated with poor prognosis in people

#### **Blood Smear**

WBC estimate -Normal estimate 5-15 WBC/10x RBC morphology Platelet estimate -Normal estimate 8-12 plts/100x Poor feathered edge and not looking for platelet clumps can alter results

# **Coagulation: Primary hemostasis**

Platelet count Platelet function -Von Willebrands disease Buccal mucosal bleeding time, as long as platelet count is normal Typically causes mucosal bleeding, epistaxis, and petechiae, not cavity bleeding

#### **Coagulation: Secondary Hemostasis**

Activated clotting time – intrinsic pathway (n < 120s)</li>
 Can affected by low platelets and can

be user dependent



#### **Coagulation: Secondary Hemostasis**

# Prothrombin time (PT) – extrinsic Activated partial thromboplastin time (APTT) - intrinsic

	S	CA	200		1	2	3)	1
	VET	RINARY CO	AGULATION	ANALYZER	4	5	6	
STAI	17 E	NAL ST		ALC: NO.	7	8	9	
	ID	QC	Data Base	Print	CANCEL	0	ENTER	
<u> </u>								

# Bilateral epistaxis, melena, hyphema, and petechia on mucous membranes



PT/aPTT WNL Thrombocytopenia on blood smear (only 3-5 platelets/hpf) Destruction (ITP) Use (hemorrhage, DIC) Non-production (bone marrow) disease)

Dyspnea, dull lung sound ventrally, pale mucous membranes, hemothorax on thoracocentesis, PCV = 20%, TP = 3.9 g/dL
 Elevated PT only
 Rodenticide toxicity

#### Vomiting, icterus, bruising ventrally



Pictures courtesy of Dr. Robin VanMetre

# **Coagulation Case Examples**

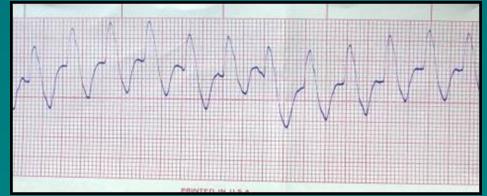
Elevated PT/aPTT
 DIC
 Increased PT/aPTT
 Low platelets
 Increased FDPs/D-dimers
 Liver failure

# Electrocardiogram



### Electrocardiogram

- Tachycardia
  - Shock
  - Pain
  - Primary cardiac arrhythmia
- Bradycardia
  - Electrolyte abnormalities (K<sup>+)</sup>
  - Primary cardiac arrhythmia (AV block)



### Electrocardiogram

Arrhythmias (VPC's, A-fib)
Primary vs. secondary
Electrical alternans
Pericardial effusion

### **Electrocardiogram Limitations**

- Poor contact, shivering, and patient movement can lead to background noise, making ECG difficult to interpret
- Electrical activity can continue long after the heart has stopped (PEA/EMD)!
- ECG is NO substitute for an audible pulse or heartbeat
- Alcohol as contact agent can cause fire if patient needs to be defibrillated

### Indirect Blood Pressure

#### BP = CO x SVR

- One of the best objective measurement of perfusion we have
- May be difficult to measure during hypotension/poor perfusion
- Target MAP > 80 mmHg in awake patients
- Indirect methods
  - **Doppler**
  - Oscillometric

# **Blood Pressure: Doppler**

- Only gives systolic blood pressure
- May give mean pressure in cats under anesthesia
- Allows for audible pulse throughout procedure
- May not work well during severe vasoconstriction



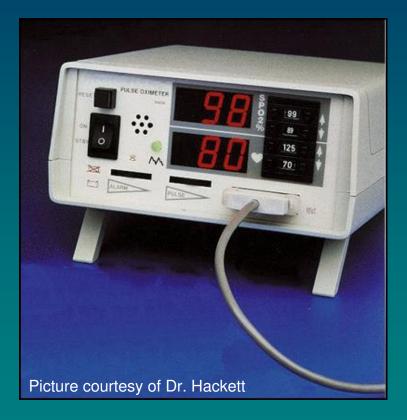
### **Blood Pressure: Oscillometric**

- Gives systolic, diastolic, and mean
- Must match heart rate to be considered accurate
- Cuff size important
- Cardell® only oscillo. BP accurate in cats
- Won't read well with poor perfusion and hypotension



# **Pulse Oximetry**

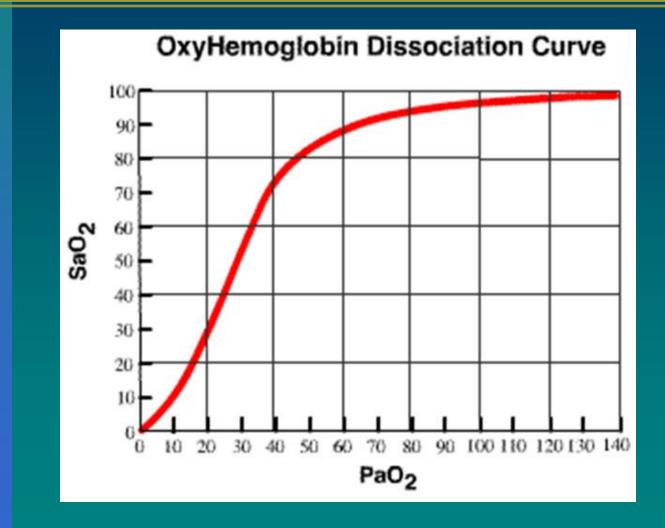
- Indirect method of measuring hemoglobin saturation
- Two wavelengths of light emitted to determine oxyhemoglobin and deoxyhemoglobin levels
- Accounts for tissue absorption by assuming only arterial blood pulsates



### **Pulse Oximetry - Artifacts**

- Ambient light
- Low perfusion
- Motion
- Look for matching heart rate and good plethysmograph waveform
- Reading frequently defaults to 85% when artifacts are present
- Carboxyhemoglobin/methemoglobin

#### **Oxyhemoglobin Dissociation Curve**



Graph

#### Beyond the EDB: Additional Monitoring

Respiratory -Capnography -Respirometry – Apnea monitors -Tidal flow-volume loops -Transcutaneous oxygen monitoring

# Capnography

In the normal healthy dog or cat, the ETCO<sub>2</sub> is a good representation of the PaCO<sub>2</sub>
 The ETCO<sub>2</sub> is usually 4-6 mmHg lower than the PaCO<sub>2</sub>



The ETCO<sub>2</sub>-PaCO<sub>2</sub> gradient is due to mixing of gas from alveoli that are being perfused with alveoli that are not being perfused <u>– Dead space ventilation</u>

#### **Cardiac Arrest**

- During cardiac arrest the ETCO<sub>2</sub> frequently is less than 10 mmHg due to little to no pulmonary capillary perfusion
- The ETCO<sub>2</sub> is therefore a great tool for determining:
  - Cardiac arrest
  - Adequacy of chest compressions during CPR
  - Return to spontaneous circulation

### Beyond the EDB: Additional Monitoring

Cardiovascular
 Direct arterial blood pressure
 Central venous pressure

### **Direct Arterial Blood Pressure**

Arterial blood gas sampling **Continuous real-time monitoring** Intentional pharmacological or mechanical cardiovascular manipulation Failure of indirect BP monitoring Supplementary diagnostic clues

#### **Central Venous Pressure Monitoring**

- Need central line in right atrium
- Normal  $< 5 \text{ cmH}_2\text{O}$ , but look for trends
- Determines preload to the heart
  - High CVP = Volume overload or low cardiac output
    - Low CVP = Hypovolemia
- Uses
- Evaluate response to and capacity for fluid loading
- Aids in determining success of pericardiocentesis for cardiac tamponade

# **CVP** Monitoring

- Can be artificially high if patient in dorsal recumbency or receiving PPV
- Water manometers read in cmH<sub>2</sub>O and pressure transducers read in mmHg (1 mmHg = 1.36 cmH<sub>2</sub>O)
- Animal should be in lateral recumbency and transducer at level of the heart
- Kinked catheters can lead to artificially high CVP's

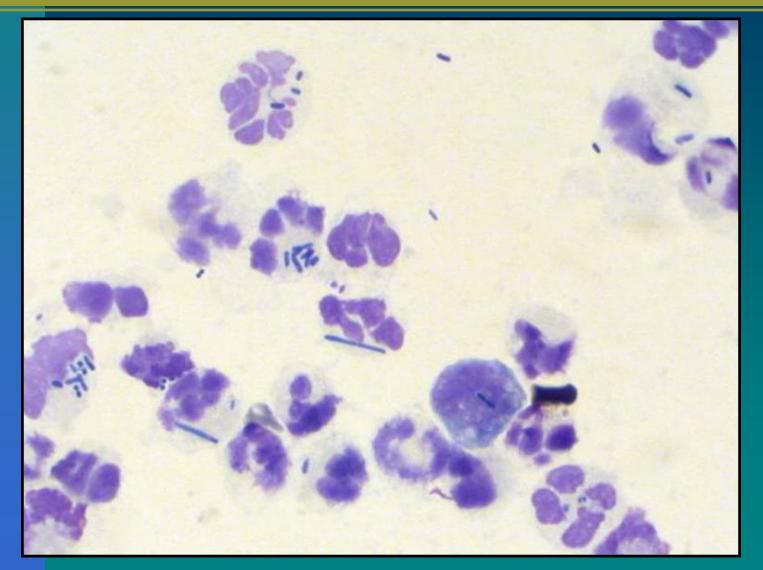
### Beyond the EDB: Additional Monitoring

Abdominal monitoring
Cytology
Colloid osmotic pressure

# **Abdominal Monitoring**

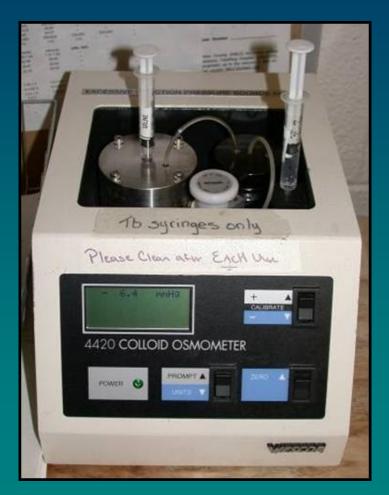
Intra-abdominal pressure Abdominal girth measurements Abdominocentesis -Generally need 20-30 ml/kg of fluid in the abdomen to have a positive blind abdominal tap Paired lactate and glucose levels Diagnostic peritoneal lavage





# **Colloid osmotic pressure**

- Measures the colloid osmotic pressure in whole blood
- Normal COP = 18-22 mmHg
- Useful to monitor response to artificial colloid use



#### Conclusions

- Point-of-care testing is essential to aid in quick diagnosis of lifethreatening diseases
- The limitations of point of care testing should be recognized and questionable results should be verified

#### Conclusions

Continued monitoring and assessment of dynamic patients (repeating blood tests, urine output, continuous ECG, BP, CVP) is necessary in order to remain proactive in treatment of critical patients