



A GAME CHANGING IN VETERINARIAN MORPHOLOGY TECHNOLOGY

Jiaying Liang
DVM



TOOTOO MEDITECH CO., LTD



What is a Game Changing Technology?



**Two essential factors for innovation:
Needs and Challenges!**

- What about in veterinary Diagnostics?

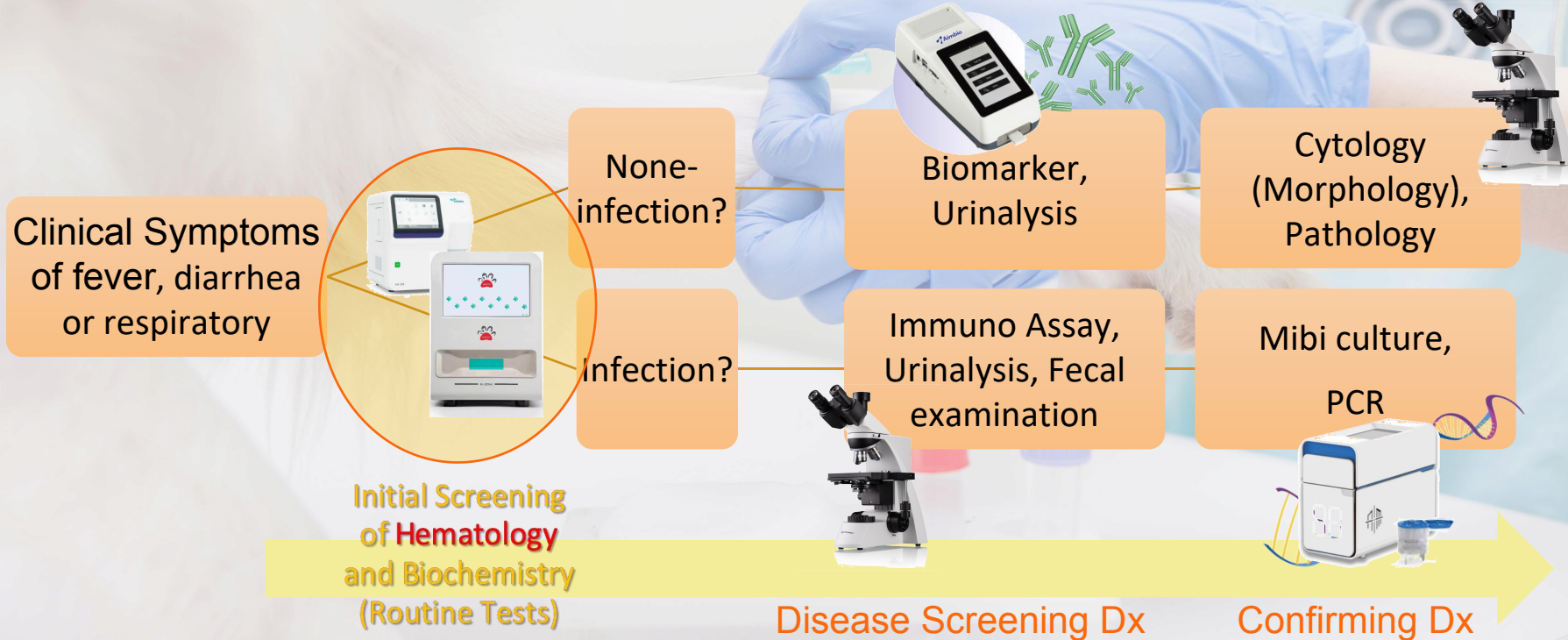


Most Common Laboratory Tests (In-Vitro-Diagnostics) in Small Animal Veterinary

- Blood Counting (5-6%)
- Urinalysis (1-2.5%)
- Faecal Examination (region depends 0.2-2%)
- Blood Biochemistry
- Rapid tests for infectious diseases
- Skin and other sample Cytology
- Pathology, microbiology, PCR, Immunoassay.....

Resource: Robinson NJ, etc. Diagnostic testing in first opinion small animal consultations. Vet Rec. 2015

Common Internal Medicine Diagnostics Work Flow in Small Animal Clinics



The Most Common Laboratory Device in Small Animal Clinics?

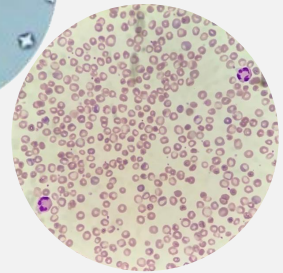
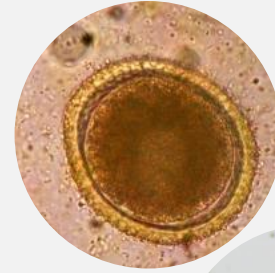


- **Cheap**
- **Multiple Function**
- **Morphology**





Why do we need morphology?



Feces Examination (Coproscopy)



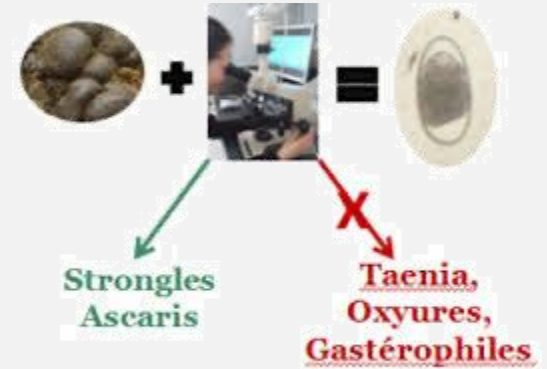
Purpose of Feces Examination and Coproscopy in Small Animal Veterinary

- Parasitology examination!!!
- Maldigestion/malabsorption
 - exocrine pancreatic insufficiency [EPI]
- dysbiosis
- other reasons



Common Challenges of Coproscopy in Small Animal Veterinary

- Time and labor consuming
- Results vary with personal skill and carefullness/patience
- Contaminative
- Flotation or not?



Challenges of Flotation Methode

Advantages

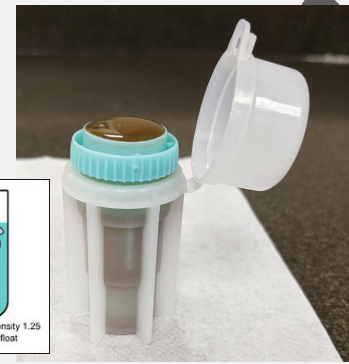
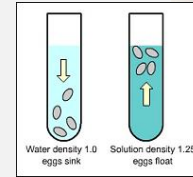
- Effectively concentrate the parasites eggs
- Provide possible clean views
- Reduce the misdiagnostics

Native(direct) smear is almost useless in parasitology!

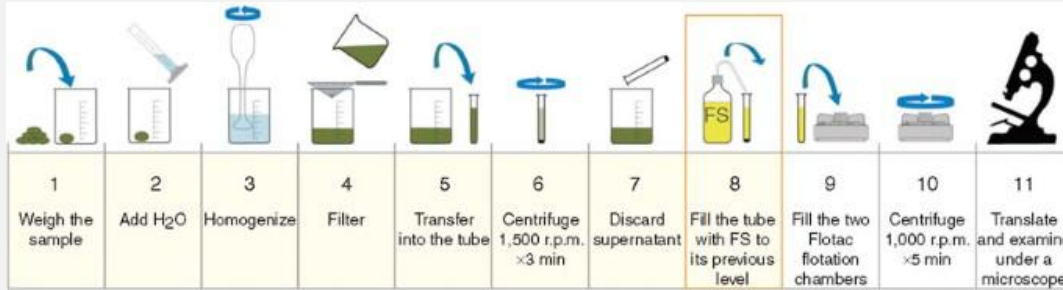


Disadvantages

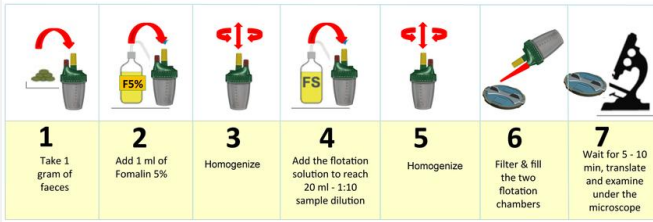
- 10-30min flotaion (+5min centrifuge)
- Operation complicated
- Test kits / flotation solution costs
- Oocysts deformation
- Trichomonas break
- Some eggs (trematodes) don't floate



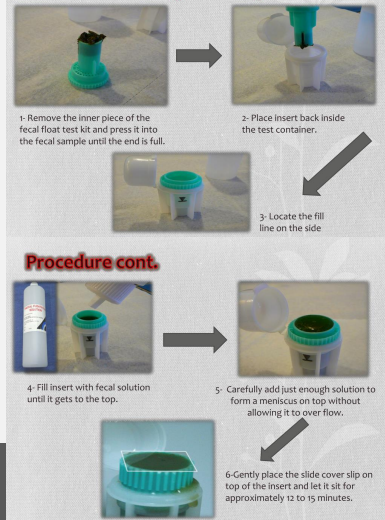
Challenges of Flotation Methode



Mini- FLOTAC basic technique + formalin 5%



How to use the FECALYZER? (procedure)

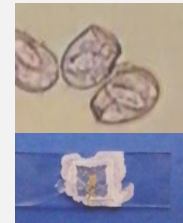


Procedure cont.

10-30 min floating time!

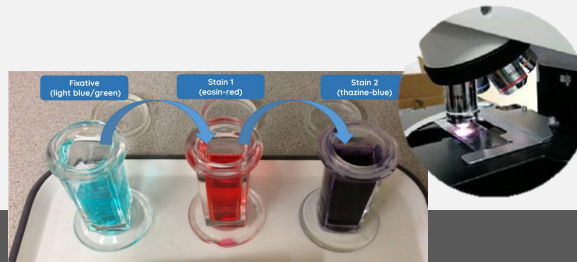
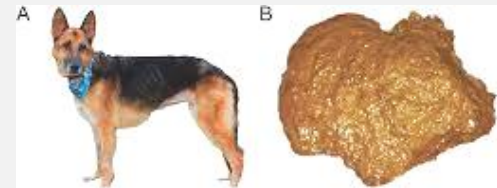
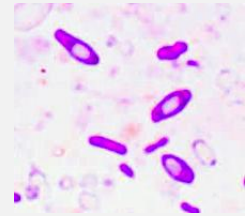
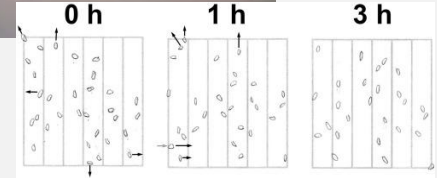
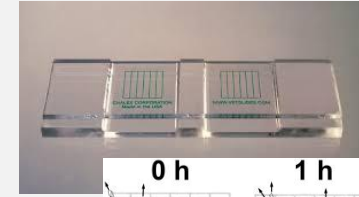
(Sedimentation: 30min)

Common problems:



And... more questions:

- How do you evaluate/monitor the therapy effect during a long-term antiparasitica therapy?
- Will you check the fecal bacteria types before making the decision of giving antibiotica or only probiotica?
- How important is the cytological and digestive information for you?



Challenges of Feces Examination

Any automation can
take over these “**dirty
jobs**” from the
vets/vet assistants?

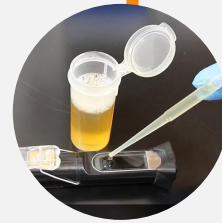




Urinalysis



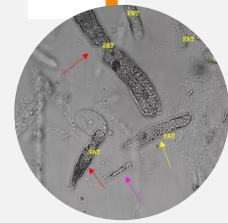
Three Essential Initial Tests of a Clinical Urinalysis for Small Animals



Specific Gravity

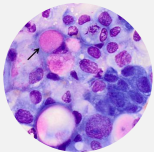


Dry Chemistry (Strip-test)



Sediment Analysis

➔ Further urine tests according to the initial examination results:



cytology (staining)



Microbiology (Reference lab)

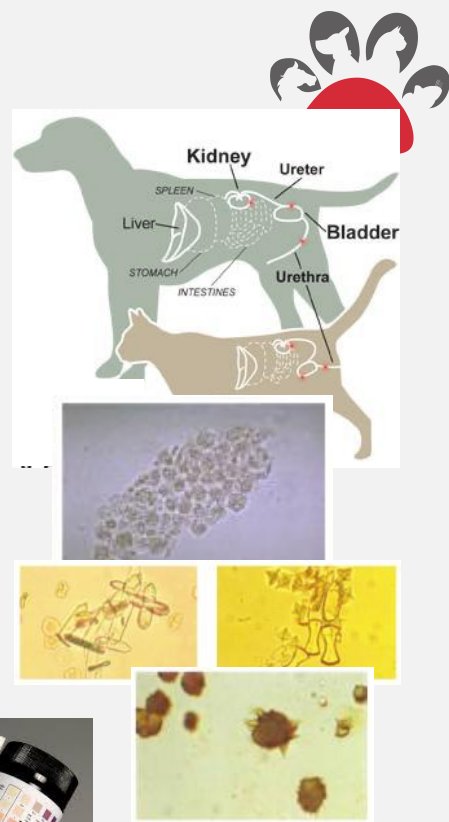


PCR, Immunoassay

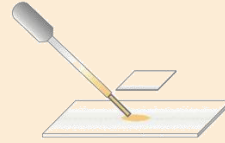


Purpose of the Urine Morphology in Small Animal Veterinary

- UTI is one of the most common disease in small animal - but with or without bacteria infection?
- Acute kidney damage can only be found through urine
- Urolith monitoring and preventure
- Poisoning diagnostics, liver failture, transitional cell carcinoma...
- Strips are not always relaiable!
 - Blood or hemolysis
 - False negative and false positive of WBC and protein
 - No info about bacteria/fungi



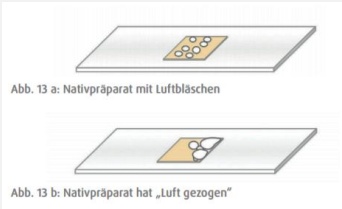
Manual Procedures (without staining)



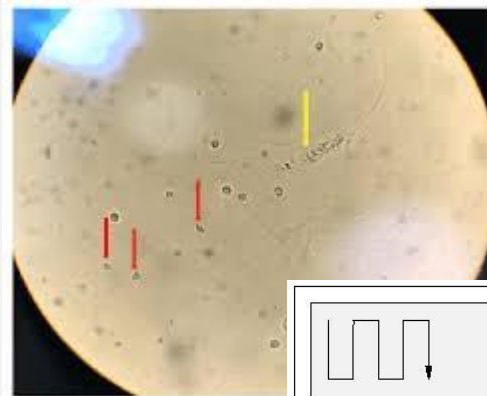
Comparison, reading and documentation

Centrifuge, preparing and manual microscopy of 20 fields (10 min)

Errors:



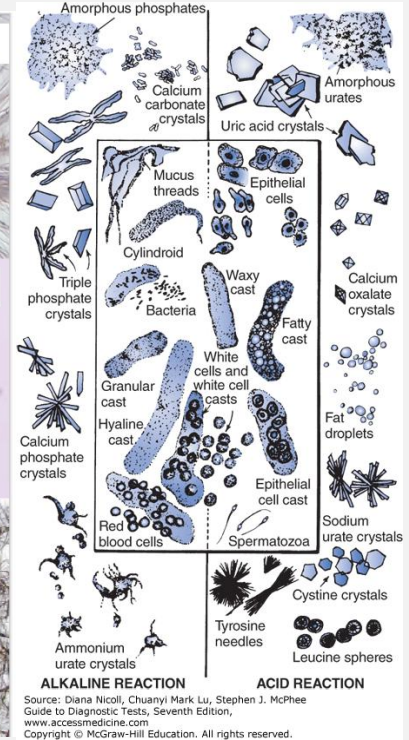
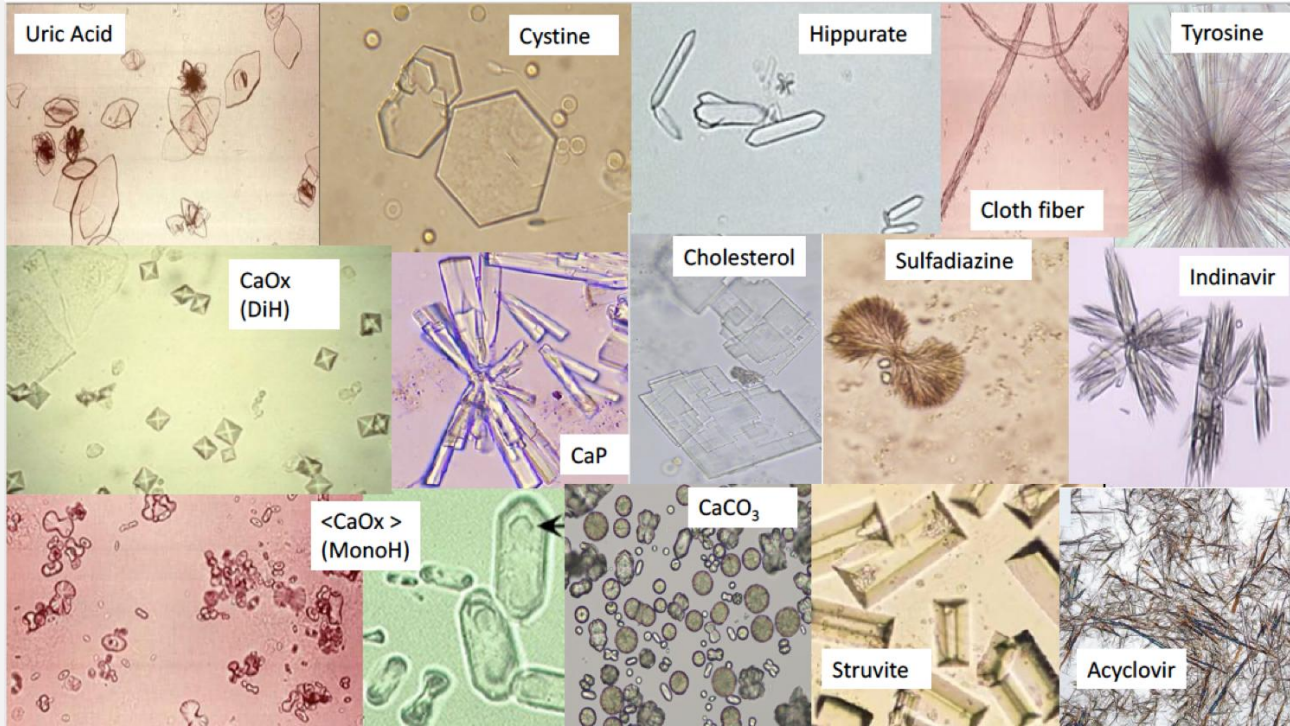
Count **RANDOMLY**
in 10 LPF and 10-20 HPF
ALL the sediments



EumEc:	0-1	/HPF	SqEc:	0-1	/HPF
DysEc:	-	/HPF	Bact:	(+)	/HPF
Lc:	0-1	/HPF			

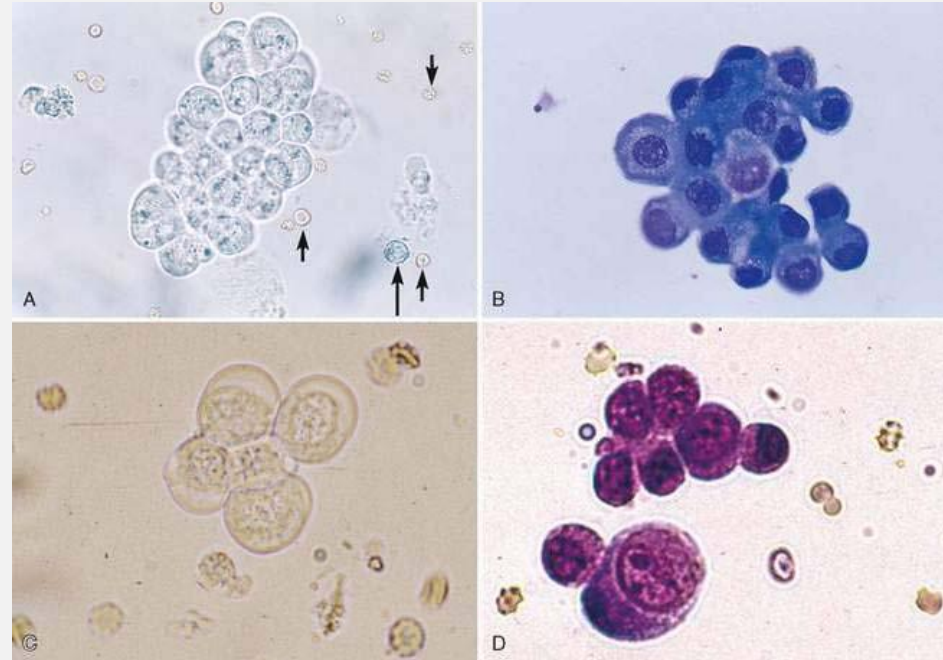
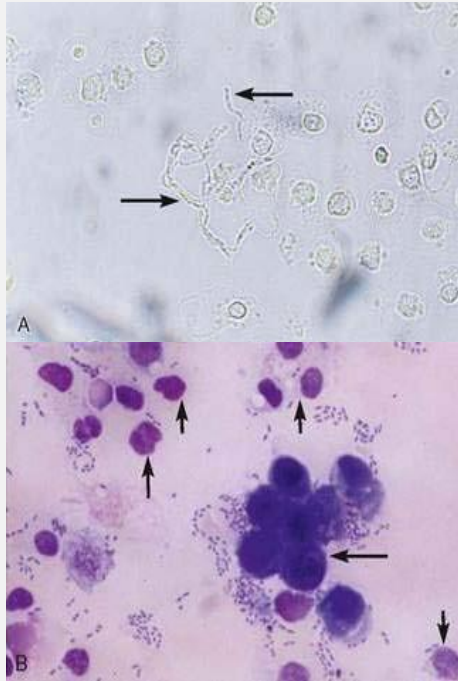
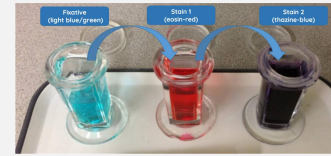
semi-quantitative!

Native (unstained) urine sediment



Source: Diana Nicoli, Chuanyi Mark Lu, Stephen J. McPhee
 Guide to Diagnostic Tests, Seventh Edition,
 www.accessmedicine.com
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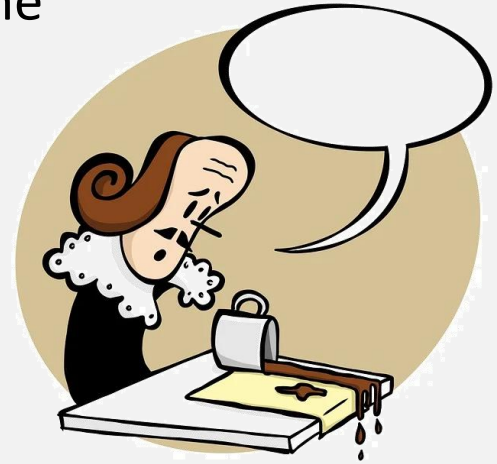
Urine cytology (stained)



transitional cell hyperplasia vs carcinoma

Challenges of Urine Morphology

- 5-10ml samples required for microscopy
(but animals don't cooperate like humans do!)
- 5-10 min centrifuge + 10-20 fields counting time
- Interpretation skills required
- Semi-quantitative results
- Stain or not stain? (crystals lost during cytological staining)
- Transport or not?
(Urine sediment can change in 30 min-4h)





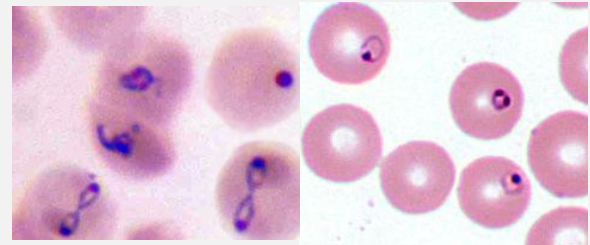
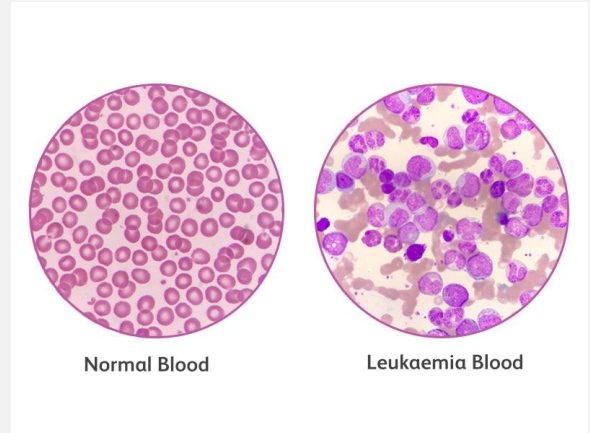
Blood Morphology



Purpose of the Blood Morphology in Small Animal Veterinary

- Anemia diagnostics and classification!
 - 20% of hospital small animal patients in USA
- Thrombocytopenia diagnostics
- WBC count, inflammation classification
- Leukemia/lymphoma
- Blood “parasites”
- Poisoning, hereditary blood disease

Luckily, CBC is invented! :D





Blood - Clinical Question 1



- If you get a patient with anaemia (low HCT), which further diagnostics will you run, or will you start a general treatment directly?





Anemia, but why?



**Non-regenerative =
not produce enough**



**Regenerative = waste
too much**



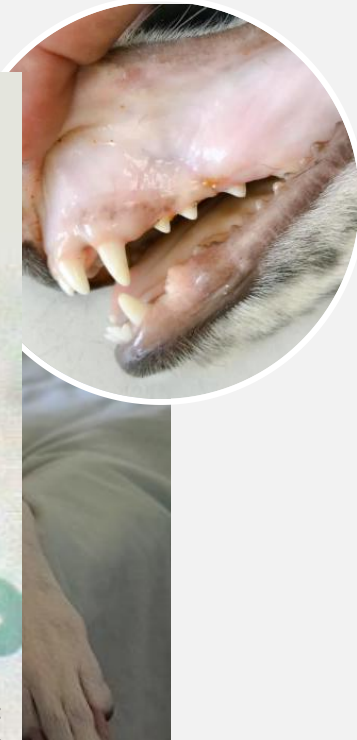
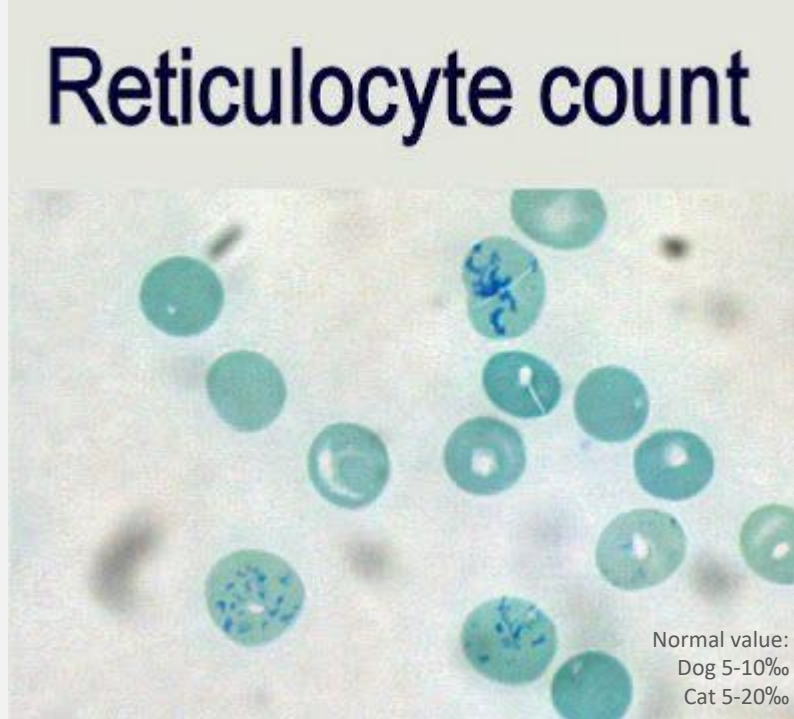
Anemia: Regenerative or Nonregenerative?

(aplastic/hypo-plastic)



IF regenerative → Haemolysis or coagulation disorder / blood losing?

IF nonregenerative → Iron deficiency or bone marrow disorder?



Causes of Regenerative Anemia

- **Blood loss**

→ moderate regenerative,
eventually total protein (TP) ↓

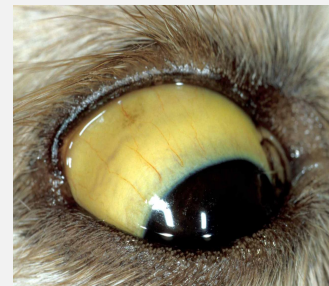
- Acute (haemorrhage)
- Chronic
 - tumor, infection, inflammation
 - Coagulation disorder



- **Hemolysis**

→ often great regenerative
with icterus & bilirubin↑

- Autoimmune
 - Non-autoimmune
- More common and more dangerous,
can die in 1-3 months without targeted therapy



Non-Regenerative Anemia



Non-regenerative =
not produce enough

- The production function is damaged and stopped
- The production function is defected, products has poor quality and can't be used
- The production is intact, but no materials for work
- The production is intact, materials are available, but no orders

intra bone marrow issues (aplastic anemia)

extra bone marrow issues

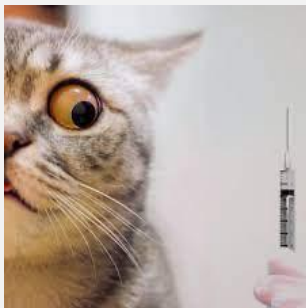




Blood - Clinical Question 2



- If you have a patient with raised WBC after a surgery, how do you distinguish a stress reaction with a real infection?



VS



WBC ↑ : infection or stress? acute or chronic?



- If you have a patient with raised WBC after a surgery, how do you distinguish a real infection?

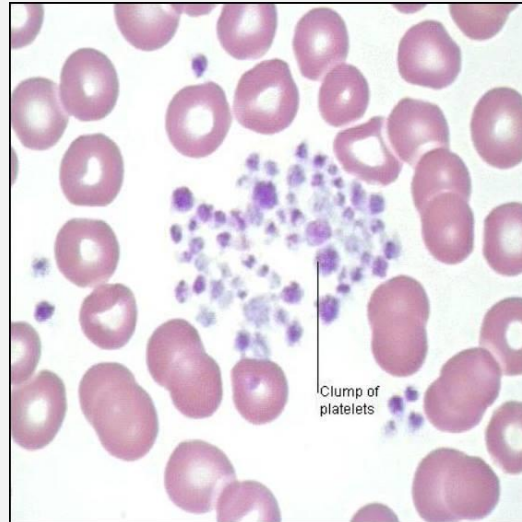




Blood - Clinical Question 3



- If a CBC shows low platelets, will you re-view the sample under microscope to rule out the in-vitro agglutination?



Is CBC enough for vet clinics?

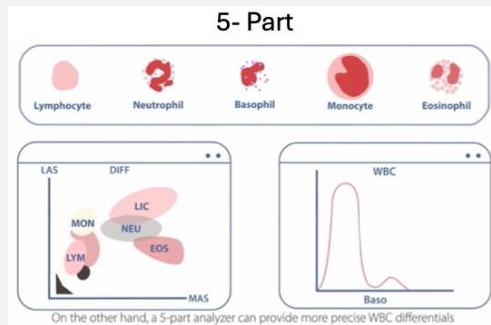
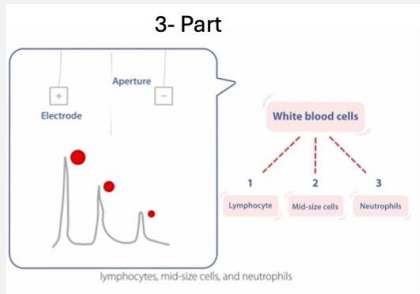


Advantages of Flow-cytometry Hematology (CBC machine)

- Easy to use
- Quantitative results
- Small amounts of sample required
- Very quickly

Challenge of in-house CBC in small animal clinics

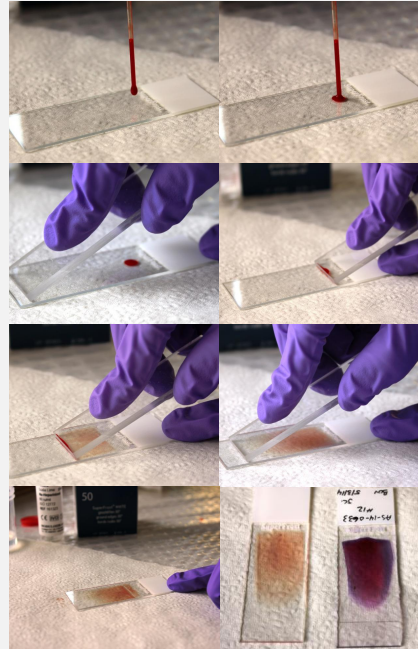
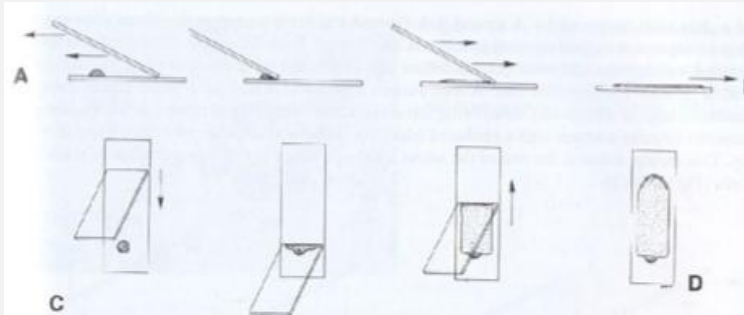
- Most don't have reticulocytes
- WBC Only 3-part or 5-part
- Not visual methods, unable to examine the morphology
- Machine limit/error/interference
- Re-inspection with manual microscope
- Only for mammals blood sample, unsuitable for reptils and birds



Manual Procedure of Blood Morphology

Blood smear procedure:

1. Blood collecting (EDTA)
2. Gently mixing
3. Smearing and drying



Common mistakes:

TOO SHORT



TOO THICK OR LONG



HESITATION



TOO THIN



RIDGES, WAVES OR UNEVEN ENDS



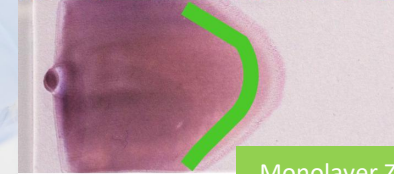
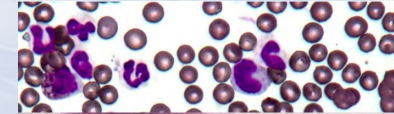
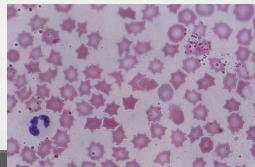
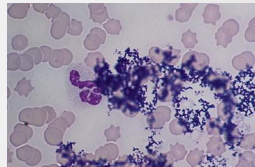
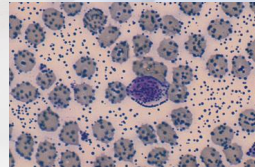
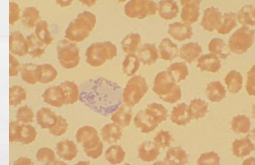


Manual Procedure of Blood Morphology

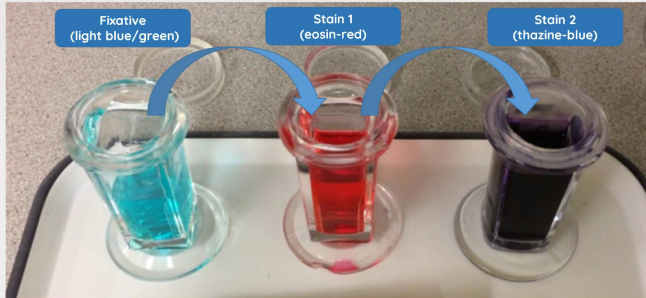
Blood smear procedure:

1. Blood collecting (EDTA)
2. Gently mixing
3. Smearing and drying
4. Staining and drying again
5. Manual examing and counting

Common mistakes:



Monolayer Zone



Count total WBCs/platelets/special RBCs with 1000 RBCs;
Count 100-200 WBCs for differential

Challenges of Blood Morphology

30 min +
at least!



**AND YOU NEED TO
KNOW WHAT YOU ARE
LOOKING FOR**

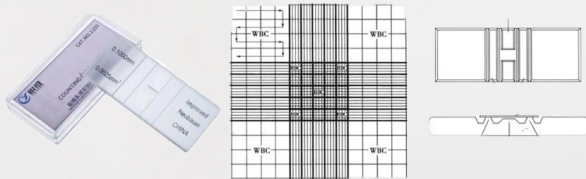


Challenges of morphology diagnostics in small animal clinics



Advantages of Microscopy

- Cheap
- visual
- multiple function



Challenges of Microscopy

- Time consuming!!!
- Personal skill training required!
(Sample preparing & interpretation)
- Intra- and Inter-observer variability
- Results not recordable (difficult)
- Quantitative complicated
- contamination



Challenges of morphology diagnostics in small animal clinics



Advantage of Reference Laboratory

- Accurate
- Morphologist available
- Professional interpretation
- Order of further diagnostics

Challenge of Reference Laboratory

- Turn-around-time too long!
- sample changes during transport (urine, oocytes parasites)
- Price

What if there's a equipment which is...

- Easy and Quickly
- Save the labor
- Accurate
- More clinical information
- Visual and recordable
- Multiple function
-
- Like a morphology expert in-house?!

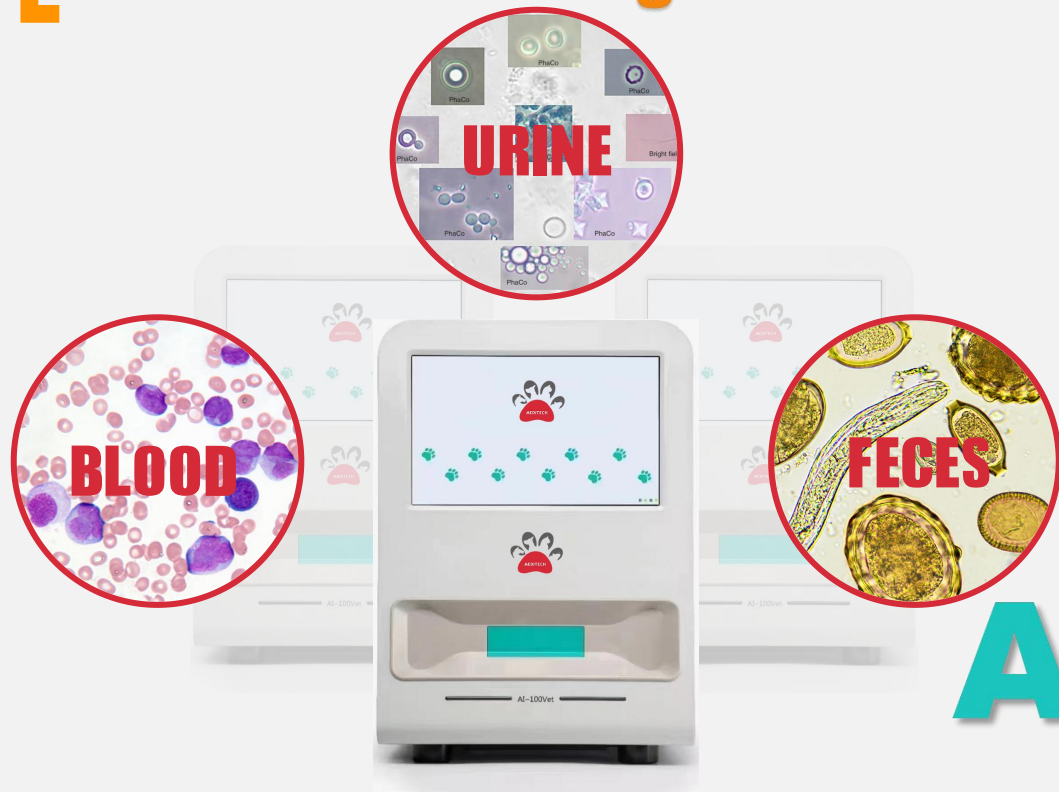


The Game Changer:

Small Animal Veterinary

3-in-1

**Automated AI
Morphology Analyzer**



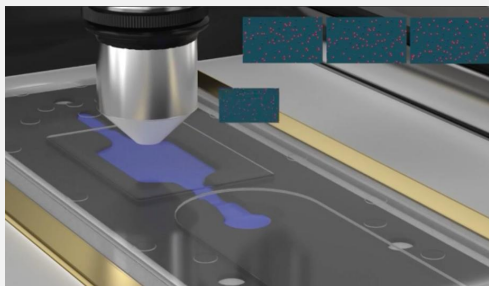
AI-100Vet

Your In-house AI morphology expert





What is AI-100VET?



An automated microscope



A multiple-function sample preparing and examining platform



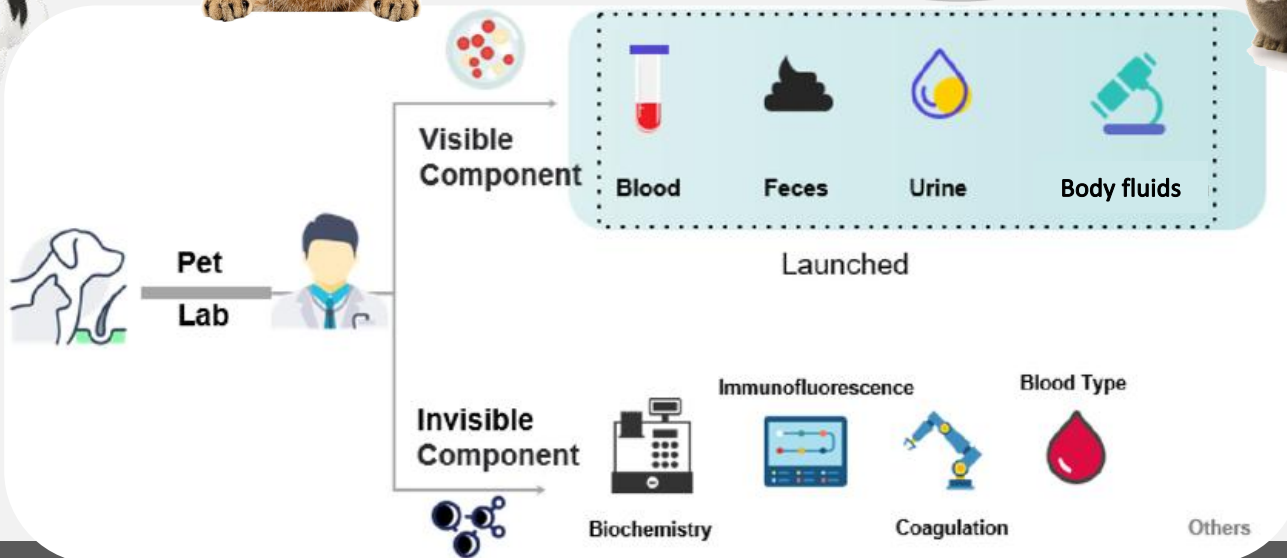
A deep-learning AI morphologist and parasitologist



An effective screening tool for clinics/hospitals



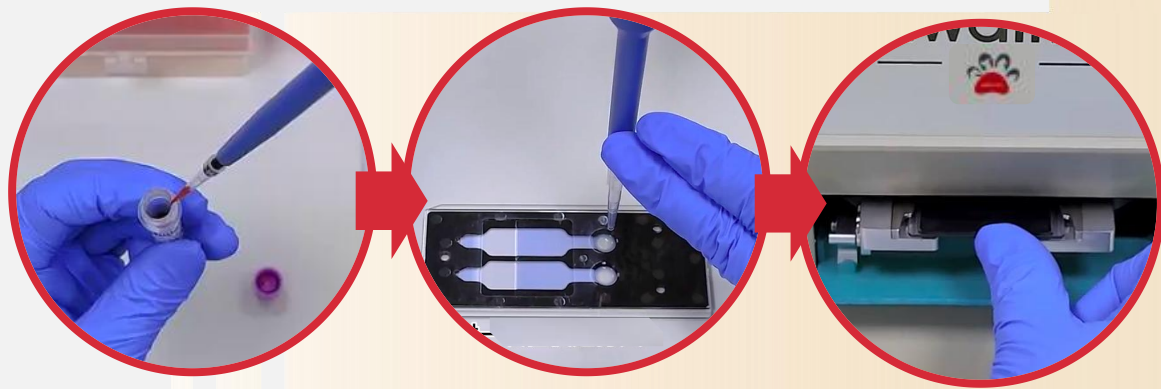
Multiple species and Multiple functions



Fully Automated with 3 Simple Steps



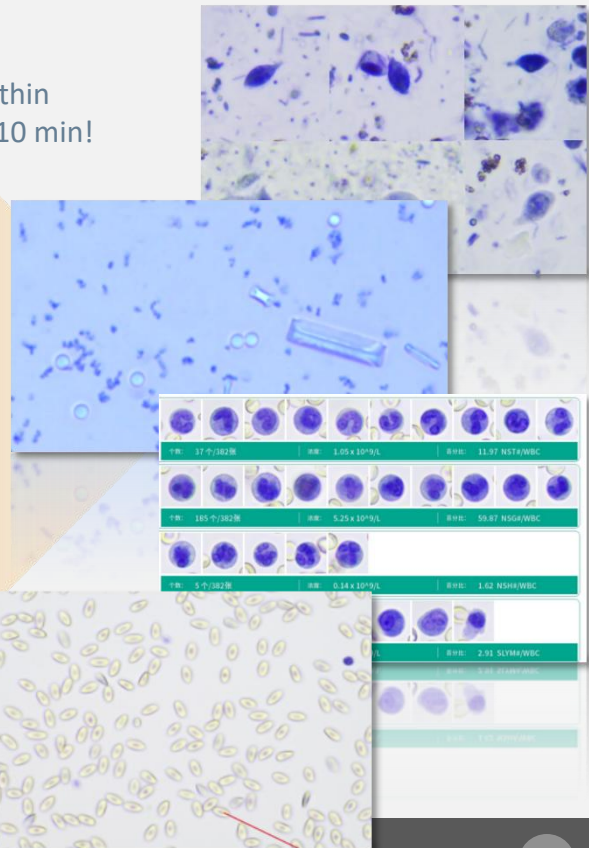
within
7-10 min!

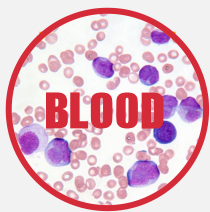


1. liquid staining
(10-500ul sample)

2. chamber injection

3. plug in
and start





Main Features



10uL of Blood

Test requires just **10 μ L** blood sample.



7.5 Min of Testing

The entire test, including PBS, is completed within **7.5 minutes**



11+ Specieses

Capability to detect various animal with **individual algorithms for each species**, including:

Mammals: dog, cat, rabbit, chinchilla, rat, mouse, hamster, ferret (and others).

Reptiles: bird, turtle, (other reptils).



46 blood parameters





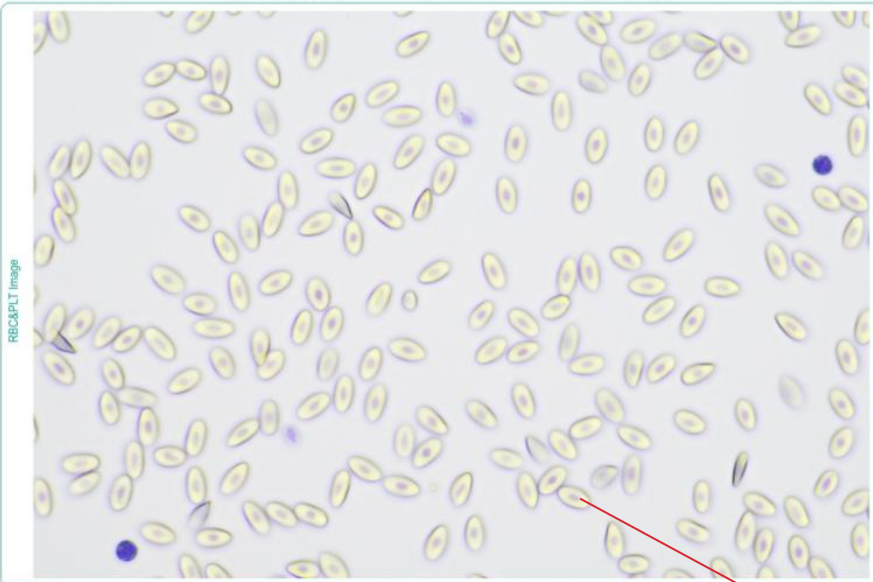
Kakaparrot Blood Morphology



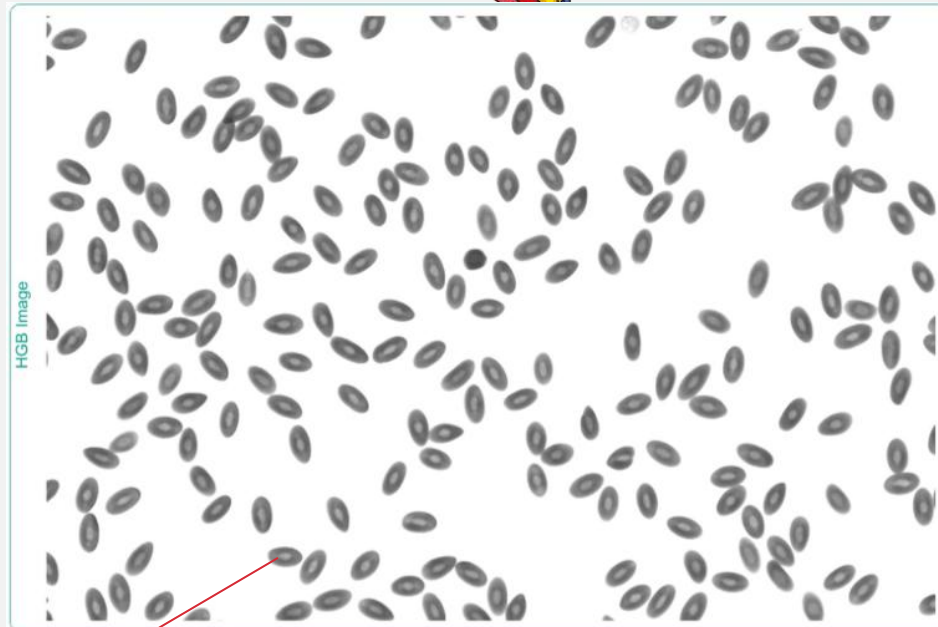
Blood Morphology Report



No.: 240726001	LIS:	Doctor: Admin	Sample: Blood(1x)	Owner: ZooMelaka
Pet name: Kakaparrot	Species: others.	Gender: M	Pet age: 0Yrs0M	Weight: 0.4kg



(whole-field-of-view with special blood staining)

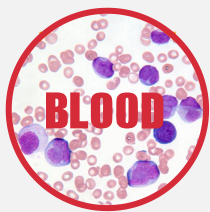


This report is solely responsible for this sample and is provided for reference purposes only. No.: 240726001 Pet name: Kakaparrot Time: 2024/07/26(10:51:10) 2/3Page

(whole-field-of-view of same position in hemoglobin channel)

Nucleated red blood cells





Main Features



10uL of Blood

Test requires just **10µL** blood sample.



7.5 Min of Testing

The entire test, including PBS, is completed within **7.5 minutes**



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Capability to detect various animal with **individual algorithms for each species**, including:

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46 blood parameters

Surpassing **9 part** diff. diagnostics including:

Left shift and right shift

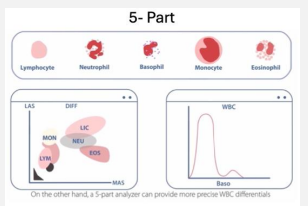
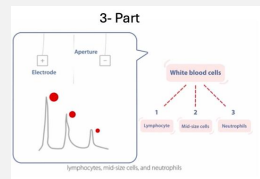
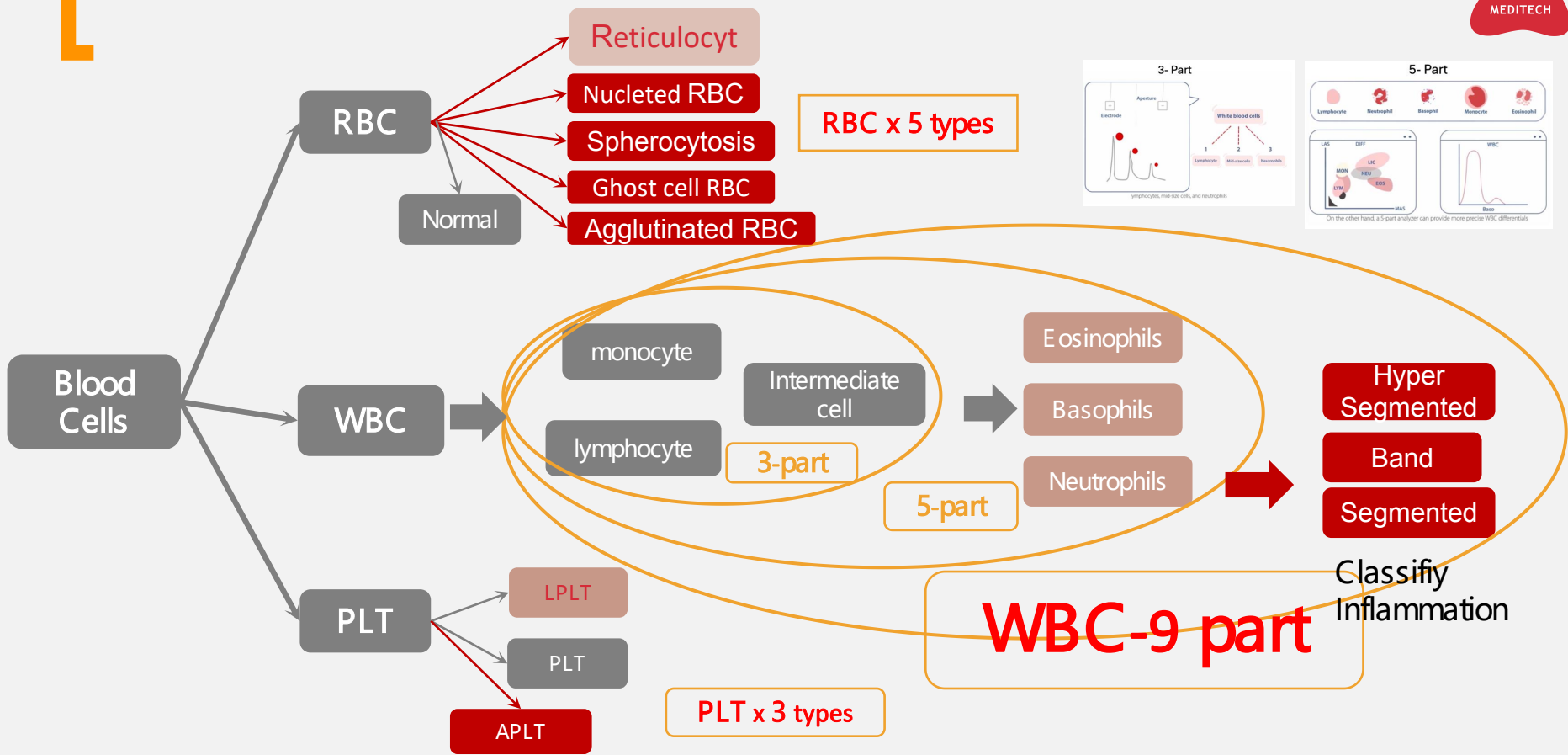
Nucleated red blood cells and Reticulocytes

Ghost cells and spherocytes

Large platelet and agglutinated platelet



Introduction and Classification of Blood Cells



WBC-9 part

Classify Inflammation

Why vets need these parameters?



01 Anemia classification

02 Monitoring the disease progress

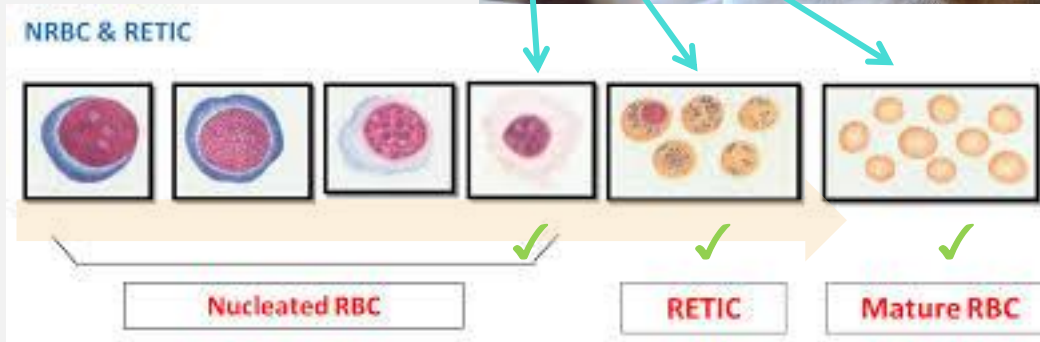
03 Avoid machine errors (misdiagnostics) and manual re-inspection



Anemia: Regenerative or Nonregenerative?

IF regenerative → Hemolysis or coagulation disorder / blood losing?

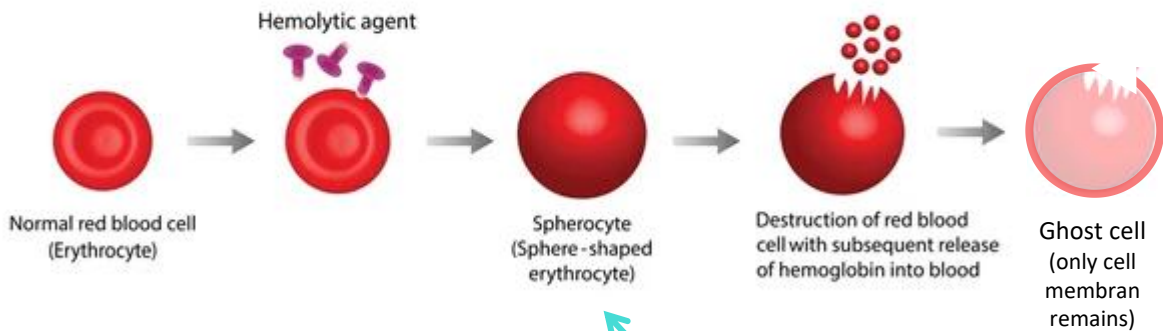
IF nonregenerative → Iron deficiency or bone marrow disorder?



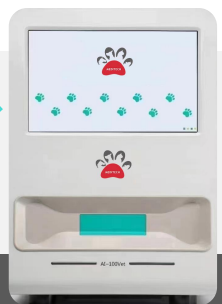
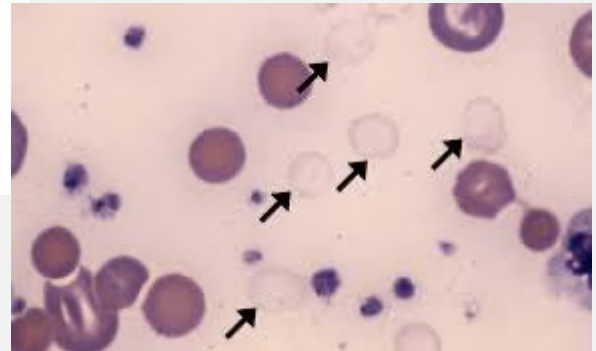


Regenerative Anemia: Hemolysis or not?

Hemolysis

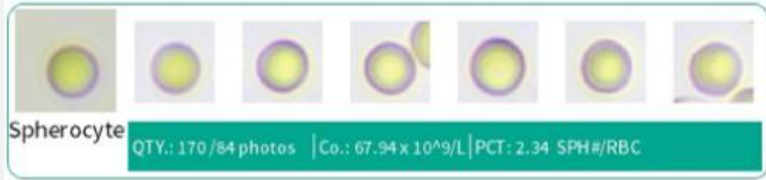


shutterstock.com · 2391470211



Hemolysis Anemia: Cortisol or not?

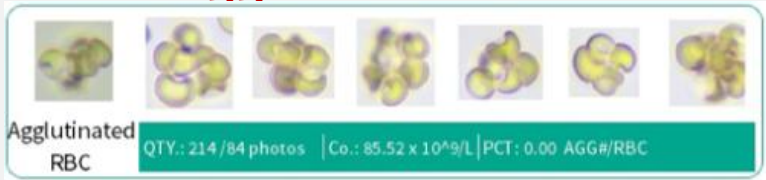
Spherocytes



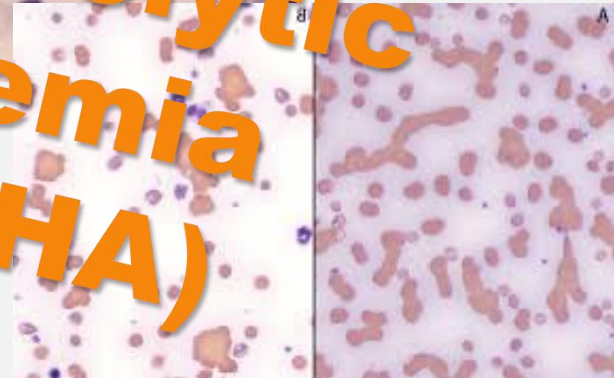
Erythrocyte Ghost Cell



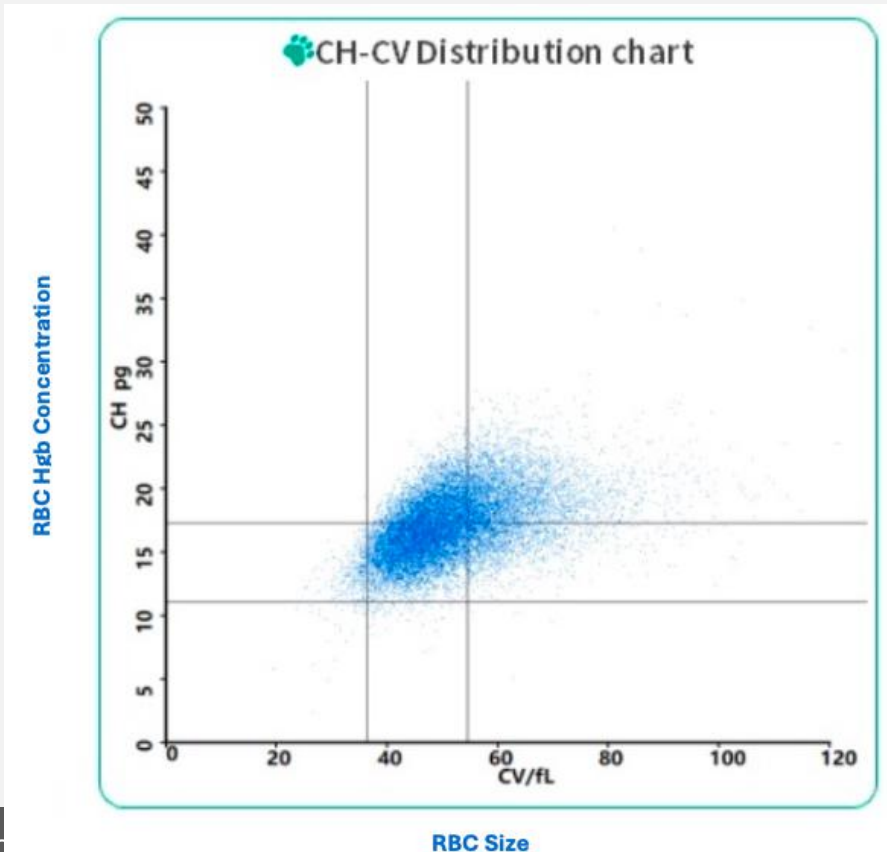
RBC Agglutination



**Autoimmune
Haemolytic
Anemia
(IMHA)**



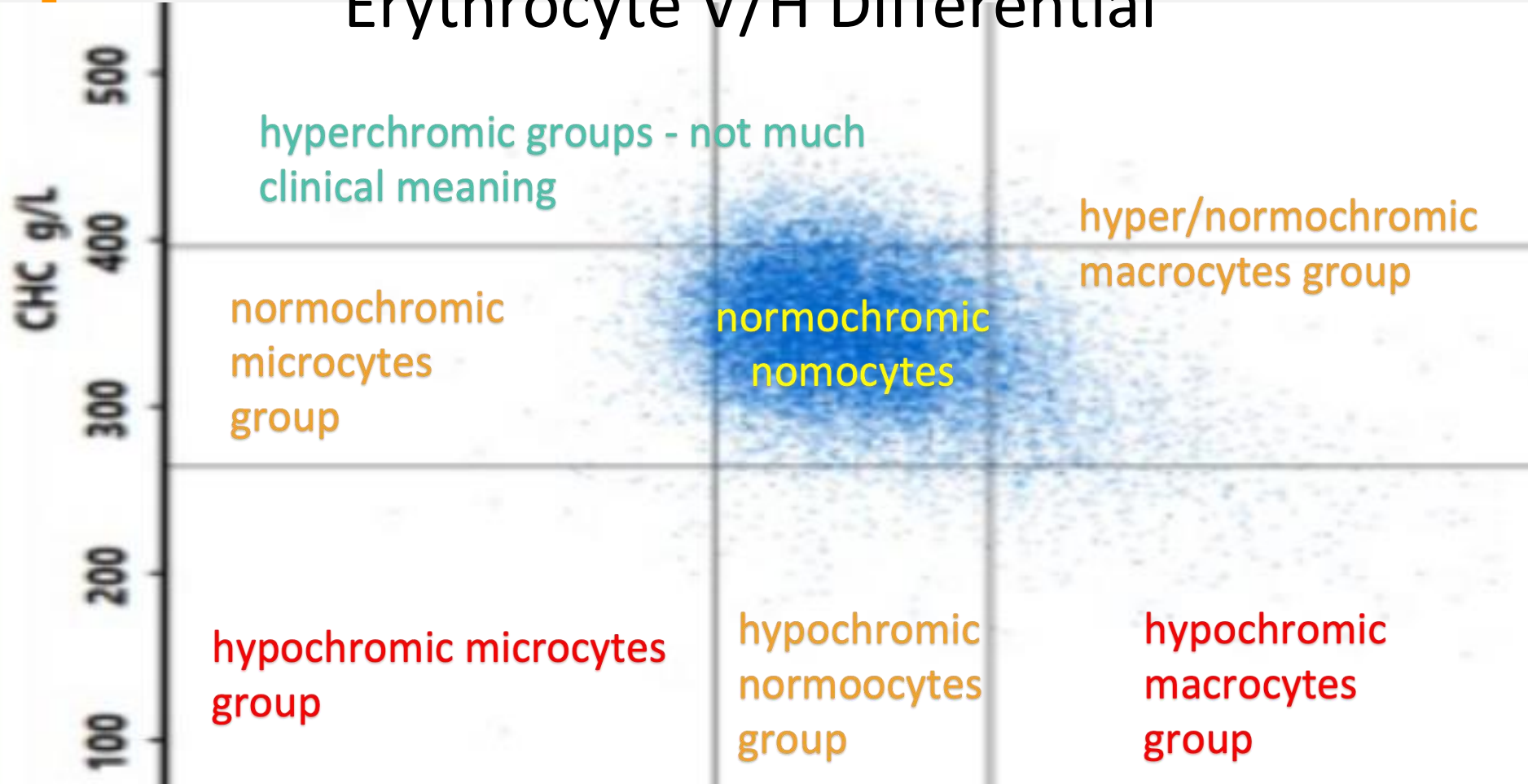
Non-regenerative Anemia: Causal?



0.01	24.32	21.62
2.01	42.58	8.87
0.20	0.35	0.03

- **Erythrocyte V/H Differential:** the first real RBC-differential analyzer for small animal vets
- Direct information of **micro-/normo-/macrocytes** and **hypo-/normo-/hyperchromic** indicate different **causals of anemia**
- A further step of erythrocyte indices (MCV/MCH/MCHC): more sensitive and easy to understand

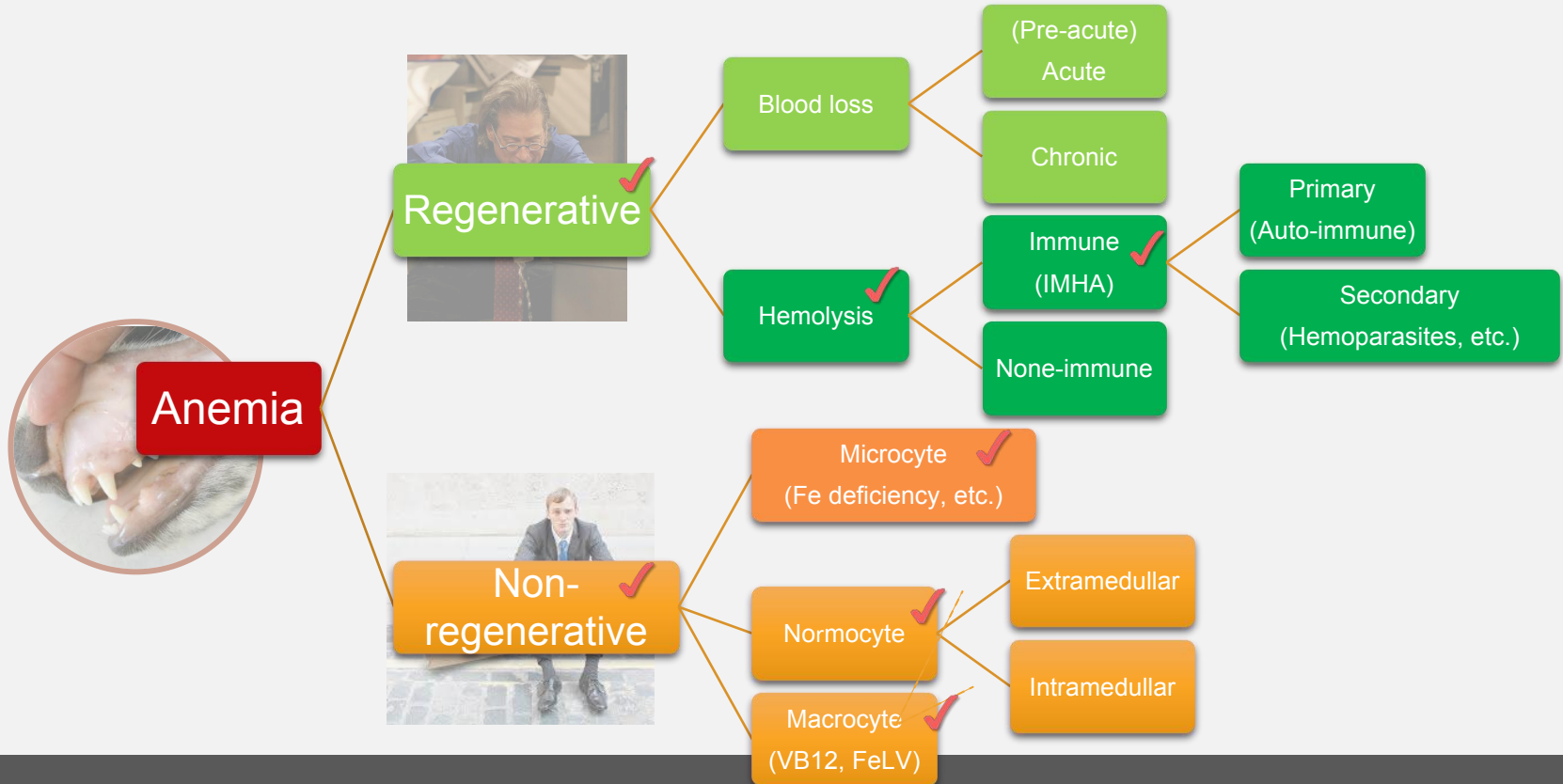
Erythrocyte V/H Differential



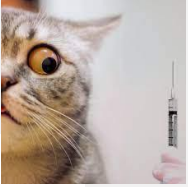
Erythrocyte V/H Differential

	Microcytic	Normocytic	Macrocytic
Hyperchromic		Dehydration (MCH \uparrow , MCHC normal)	Megaloblastic: VB12 or folate deficiency (e.g. dysbiosis); Myelodysplasia (leukemia, FeLV); Sulfonamid, methotrexat; Regenerative response (Reticulocyte)
Normochromic	Portosystemic Shunt (PSS) (Akita + Shiba NORMAL)	Hemolysis Acute blood loss Acute pancreatitis Chronic renal disease, heart-kideny-failure (EPO \downarrow) Endocrine disease (hypothyriasis, Addison syndrom, estrogen) Protein deficiency/cachexia (tumor, chronic disease/inflammation) Bone marrow failture (immunogic aplastic, FIV, FIP, Leishmaniosis, Ehrlichiosis, tumor metastasia into BM, chemo therapy, radiation damage, intoxication e.g.lead, chloramphenicol)	
Hypochromic	Iron deficiency Cupfer deficiency Lead/cupfer poisoning Sideroblastic anemia Chronic blood loss Chronic inflammation	Mild iron deficiency (Acute IMHA)	Megaloblast or reticulocyte with 2nd. iron deficiency (Chronic IMHA)

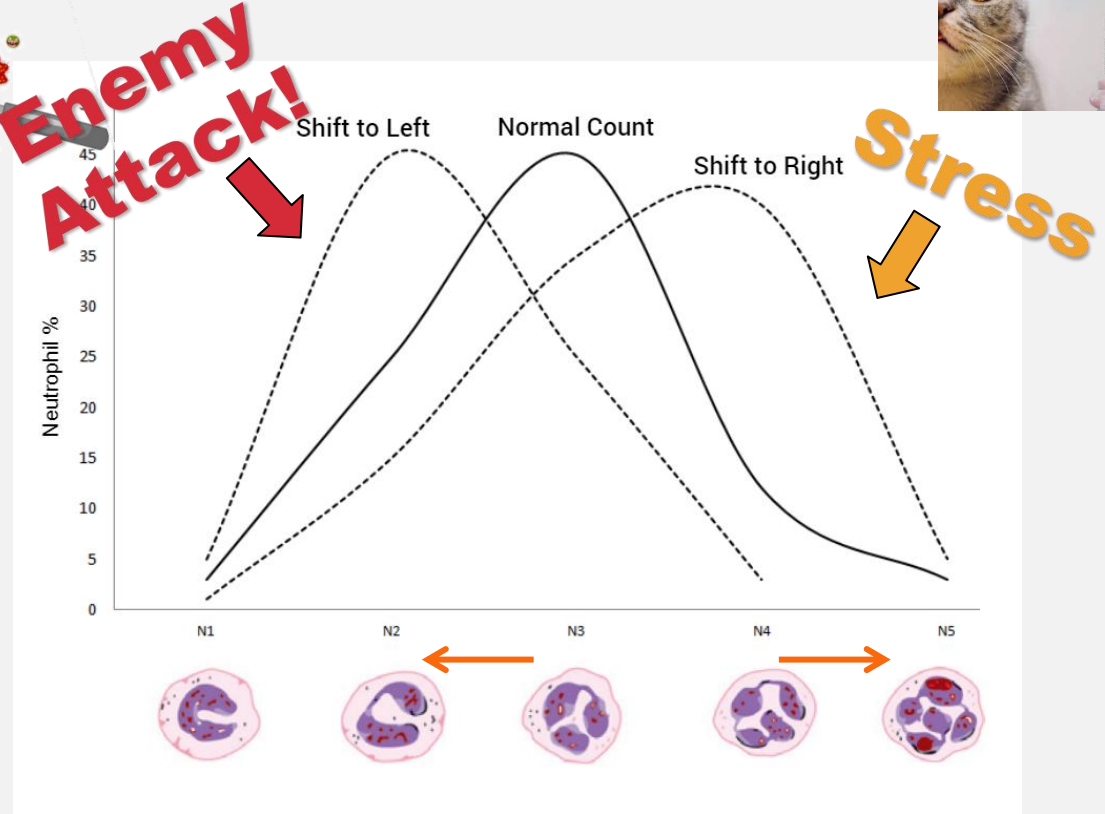
Powerful Screening Tool for Anemia Diagnostics



Infection or Stress? Acute or chronic?

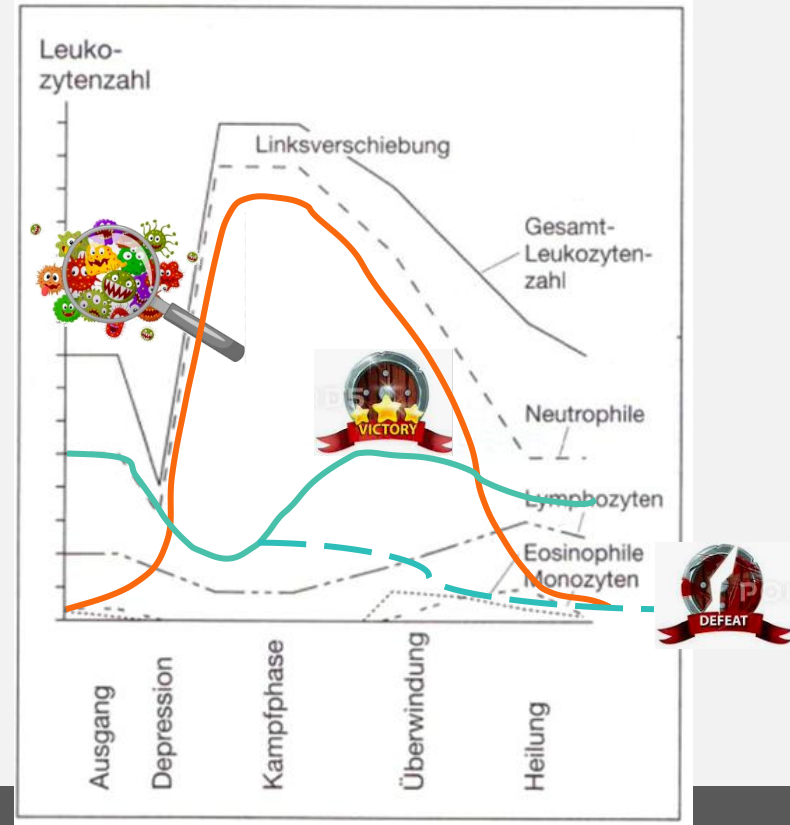


- **Left Shift:**
increasing of **immature band neutrophils** suggests more severe inflammation, often associated with a bacterial infection (Productive leucocytosis)
- **Right Shift:**
stress/excitement leukogram (Pseudoleucocytosis)



Infection: Acute or Chronic? Prognosis?

- Infection and recover process:
 1. WBC depression (initial phase)
 2. WBC (band neutrophil) increasing - fighting phase
 3. WBC decreasing (neutrophil↓, lymphocyt↑) (recover phase)



AI-100Vet: First WBC 9-part Differential



Detection items	Result	Unit
1.WBC	2.RBC	3.PLT
1-1.NEU#	2-1.HGB	3-1.PCT
1-2.NST#	2-2.HCT	3-2.MPV
1-3.NSG#	2-3.MCV	3-3.LPLT#
1-4.NSH#	2-4.MCH	3-4.P-LCR
1-5.LYM#	2-5.MCHC	3-5.APLT#
1-6.MON#	2-6.RDW-SD	3-6.PDW-SD
1-7.EOS#	2-7.RDW-CV	3-7.PDW-CV
1-8.BAS#	2-8.HDW-SD	
1-9.NEU%	2-9.HDW-CV	
1-10.NST/WBC%	2-10.RET#	
1-11.NSG%	2-11.RET%	
1-12.NSH/WBC%	2-12.NRBC#	
1-13.LYM%	2-13.NRBC/WBC%	
1-14.MON%	2-14.ETG#	
1-15.EOS%	2-15.ETG%	
1-16.BAS%	2-16.SPH#	
1-17.NST/NEU%	2-17.SPH%	
1-18.NSH/NEU%	2-18.AGG#	


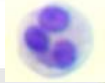

band neutrophile

segmented neutrophile

hypersegmented neutrophile

nucleated RBC

WBC Differentiation: 9-part VS 3-/5-part

3 part	5 part		AI 100VET—7 Types, 9 Part
Granulocyte	Neutrophil		Band neutrophil
			Segmented neutrophil
			Hypersegmented neutrophil
	Eosinophil		Eosinophil
	basophil		basophil
Lymphocyte	Lymphocyte		Lymphocyte
monocyte	monocyte		monocyte

**For Better Monitoring
and Accurate Prognosis**

DDx According to the WBC 9-PART Differential

Disease	WBC	<u>Segmented Neutrophils</u>	<u>Band Neutrophils</u>	Lym	Mon	Eos	(CRP/SAA)
Acute inflammation	Stongly Increased	<i>Increased</i>	<i>Increased</i>	Decreased/ Normal	Varies	Decreased	increased
		<i>Strongly Increased</i>					
Severe inflammation	Decreased /NoChange	<i>Decreased/ Normal</i>	<i>Increased</i>	Decreased/ Normal	Varies	Varies	(strongly) increased
		<i>Normal/Slight Increased</i>					
Mild/chronic inflammation	Increased	<i>Increased</i>	<i>Slightly Increased/ Normal</i>	Varies	Increased	Increased/ Varies	slightly increased
Excitement Leukogram	Increased	<i>Increased/ Normal</i>	<i>Normal</i>	Normal/ Increased in cats	Varies	Varies	Normal
		<i>Normal/Slight Increased</i>					
Stress leukogram	Increased	<i>Increased</i>	<i>Normal</i>	Decreased	Normal/ Increased in dogs	Decreased/ Varies	Normal/ increased in aseptic inflammation
		<i>Increased</i>					



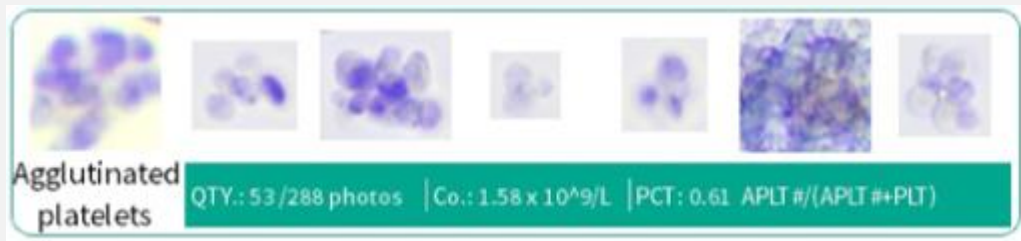
Avoid machine errors (misdiagnostics) and manual re-inspection

Thrombocytopenia: real or misdiagnosed?

Large Platelet (LPLT) differential with RBC

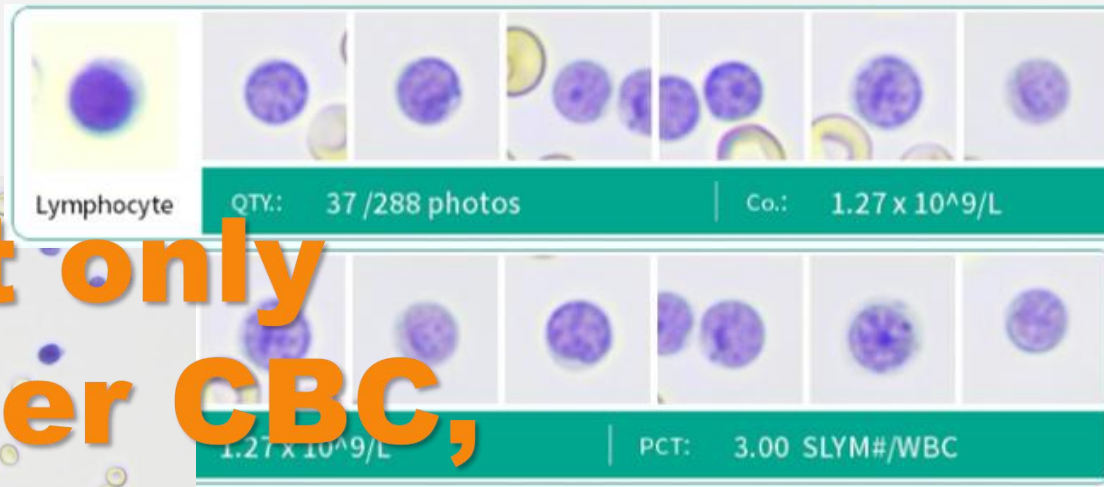


Agglutinated Platelet differential with WBC





Moreover...



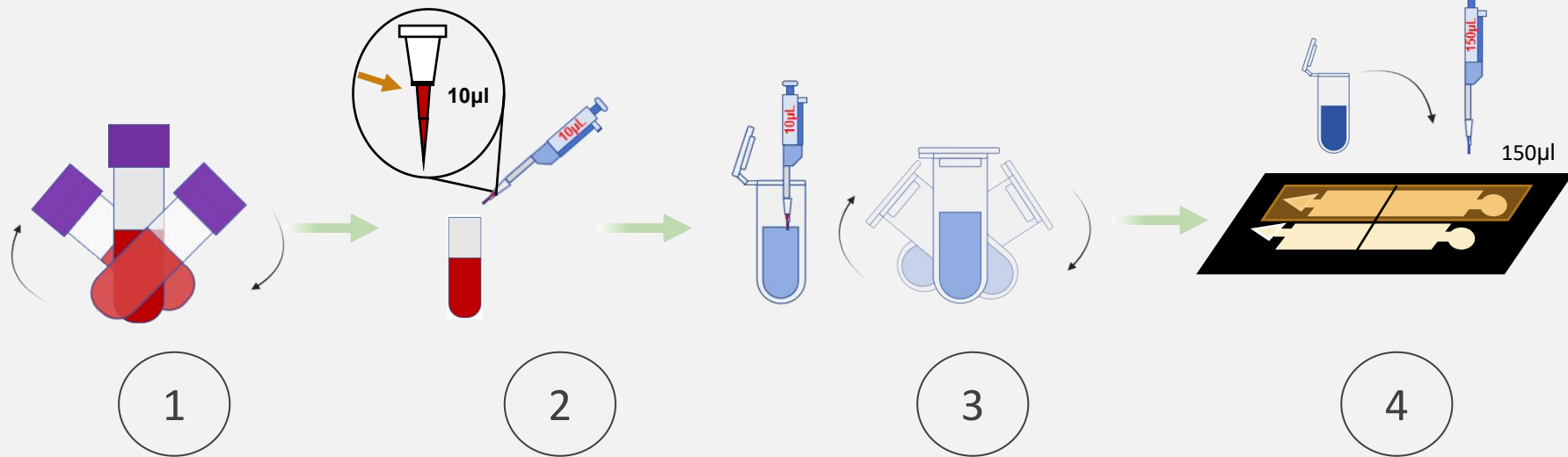
**Not only
another CBC,
but
Morphology!**

Discovery of Leukemia!
(highly possible small-cell lymphoma)





4-Steps Procedure Blood Testing for Dogs and Cats



1

Before starting the test, gently and thoroughly mix the EDTA-blood sample (reptiles:Heparin-blood) and ensure there are no obvious blood clots.

2

Draw **10µl** of blood, ensuring the volume of blood is at the 10µl mark. Scrape off any excess blood from the outer wall at the mouth of the collection tube.

3

Add the blood sample to the blood staining solution (Reagent, 2.5ml) and gently invert to thoroughly mix the blood sample with the staining solution.

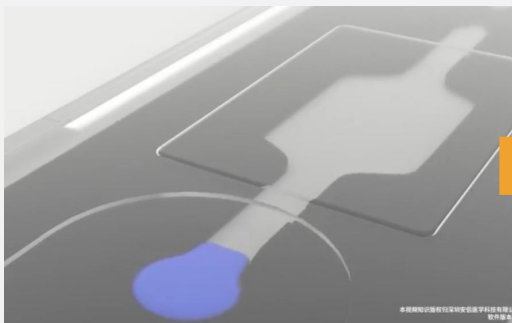
4

After mixing thoroughly, immediately draw **150µl** of the stained sample and quickly inject it into the chip channel in a single motion, then proceed with machine testing.

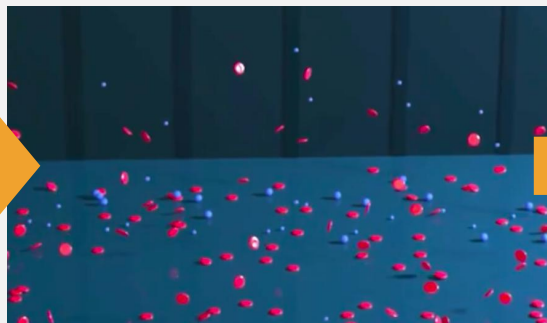




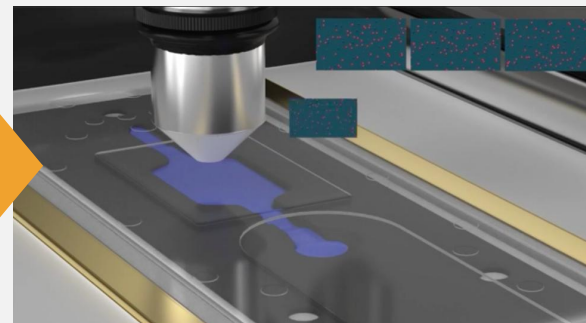
Automated Processing in 7.5 Minutes



Micro-Flow Chamber



Sedimentation to monolayer



Scan under 40x10 magnification
200-500K cells*, shooting 500+ full-
fields of view images

***Cell counting comparison:**

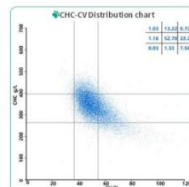
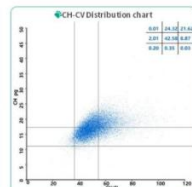
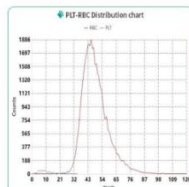
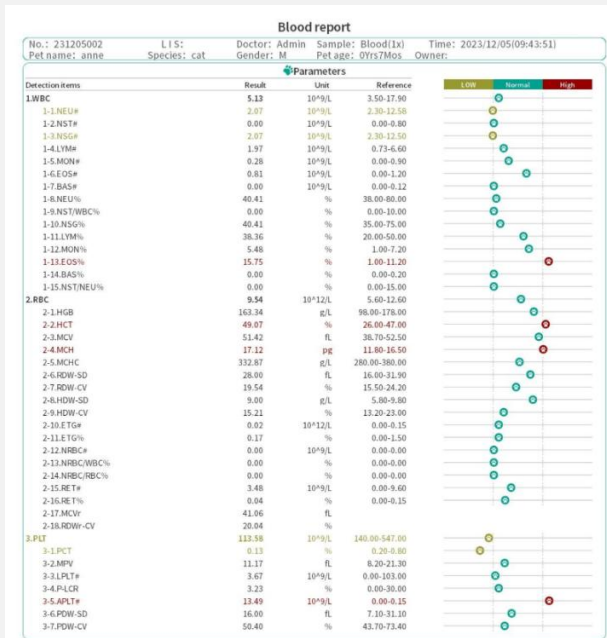
Microscope: 1000 cells

Flow-cytometry: 20K cells

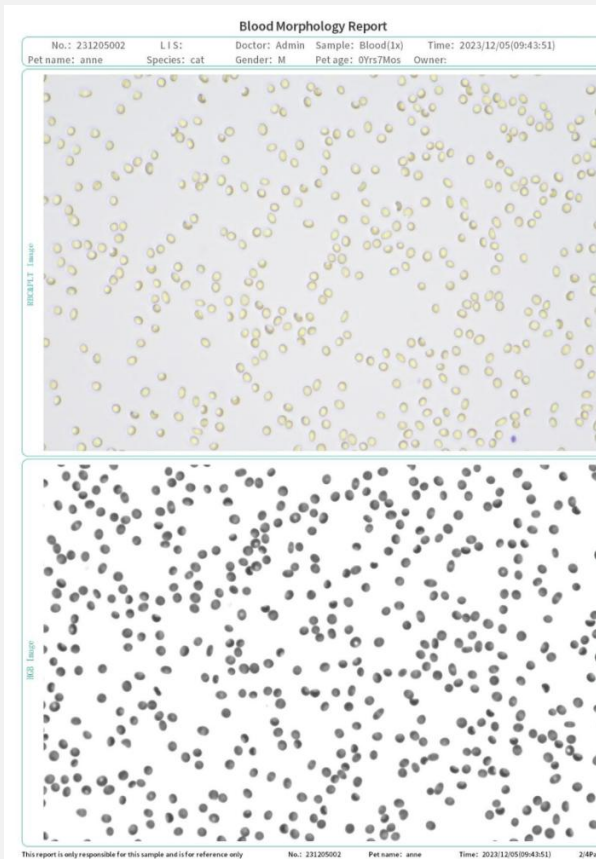




Blood Report with 45 Parameters

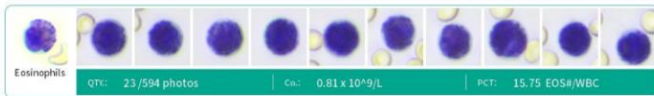
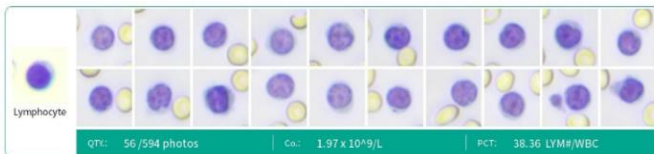
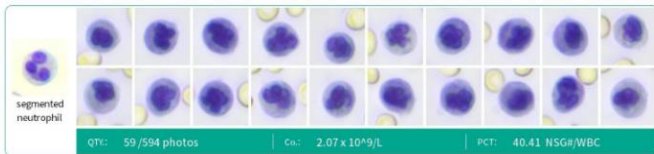


This report is only responsible for this sample and is for reference only. No.: 231205002 Pet name: anne Time: 2023/12/05/09:43:51 1/4Page





Blood Report



This report is only responsible for this sample and is for reference only

No.: 231205002

Pet name: anne

Time: 2023/12/05/09:43:51

3/4Page



Diagnostic recommendation

Single diagnosis:

一、[WBC]

1. [NEU<2.30] It is common in serious disease consumption, poisoning, physical and chemical damage and so on.

二、[RBC]

1. [HCT<47.00] It is common in dehydration, use of catecholamines, compensatory or pathological increase of erythropoietin, adrenocortical hyperfunction, hyperthyroidism, limb hypertrophy and so on.

三、[PLT<140.00] It is common in sample agglutination, hemorrhage, platelet destruction, organ detention, insufficient bone marrow formation, drug induction and so on.

1. [PCT<0.2] It suggests thrombocytopenia.

2. [APLT<0.15] It is suggested that coagulation and azotemia are common in immune-mediated thrombocytopenia, infectious diseases, malignant tumors, heart disease, drug-induced disorders and so on.

Smart- Interpretation of Diagnostics

This report is only responsible for this sample and is for reference only

No.: 231205002

Pet name: anne

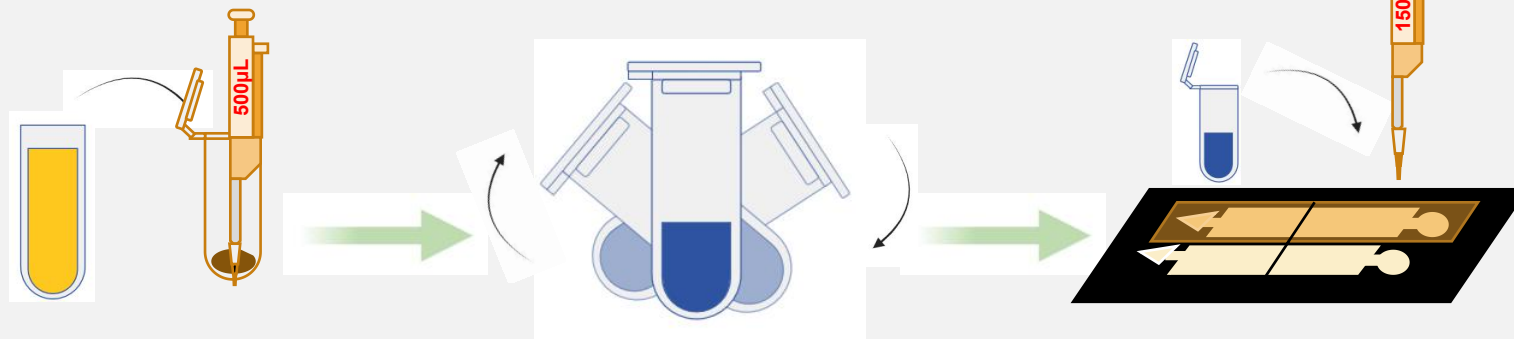
Time: 2023/12/05/09:43:51

4/4Page





3-Steps Procedure for Urine Test



1

Draw **500µl** of urine from the bottom of the test tube into the dye solution.

2

Mix the urine and dye solution thoroughly until there are no obvious dye sediments visible in the test tube.

3

Draw **150µl** of the sediment sample from the bottom of the test tube and add it to the chip for machine testing.



Getting report in **9 min**



21 Parameters of Urine Sediments:

Cast types x4

(chronic kidney disease inflammation infection)

hyaline, **cellular**, **granular**, **waxy**

For acute kidney injury!

Crystals x6

(Gout, kidney failure, urinary tract infection, hypothyroidism)

Magnesium ammonium phosphate crystal (Struvite), calcium oxalate dihydrate crystal, calcium oxalate monohydrate crystal, calcium phosphate crystal, uric acid crystal, cystine crystal

Cells x6

(Urinary tract infection, urinary tract stones, nephritis, etc.)

RBC, WBC, renal tubular epithelial cells, squamous epithelia cells, transitional epithelia cells, sperm cells

Pathogenic microorganisms x3

(Urinary tract infection)

cocci, bacilli, yeasts

Others x2

(Differential from Artefacts)

lipid drop, mucus, (unintentional objects)



Urine Morphology Report with 21 Parameters

Urine report

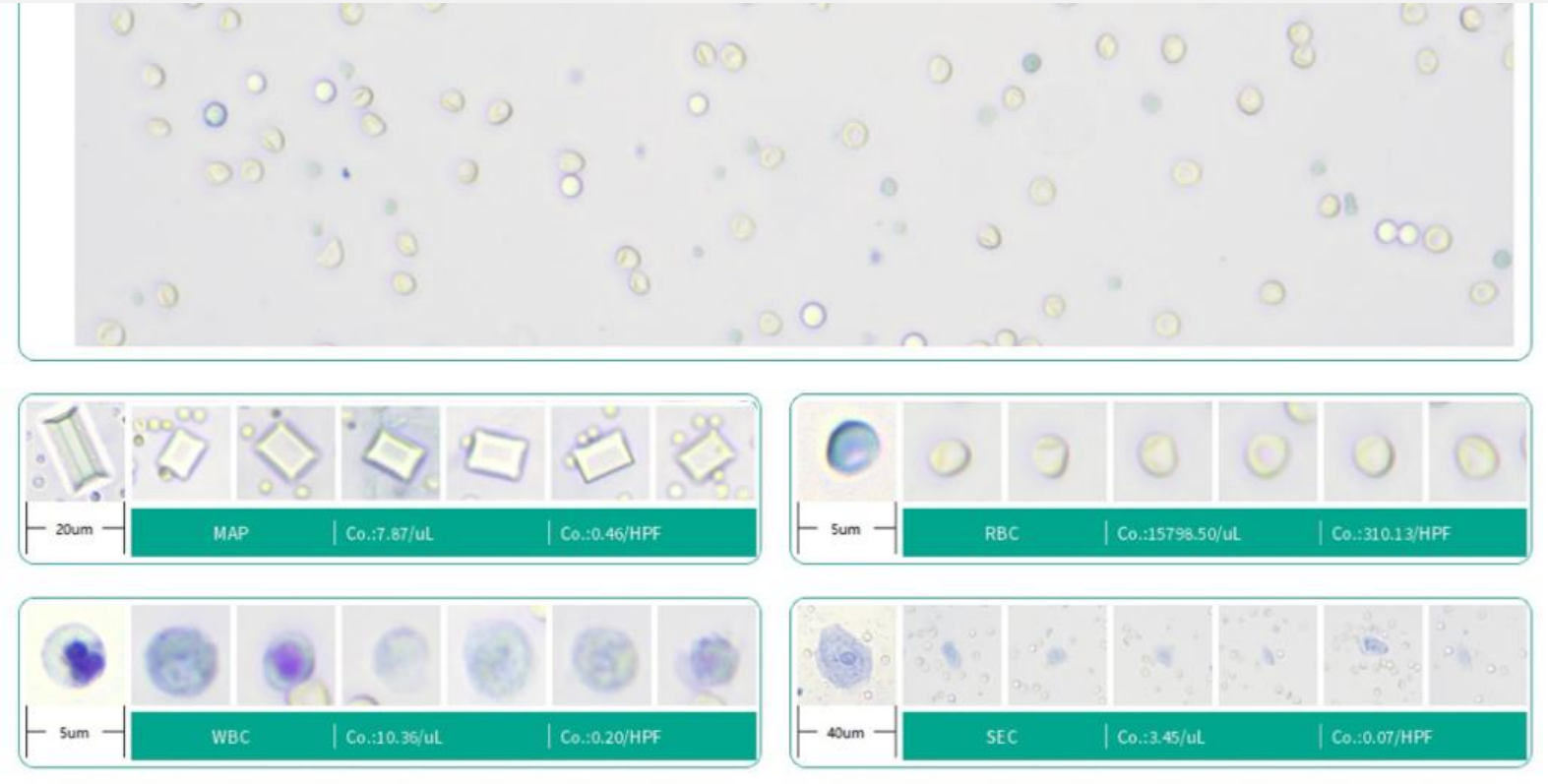
No.: 231205008 LIS: Doctor: Admin Sample: Urine Time: 2023/12/05(14:24:35)
 Pet name: Emmm Species: dog Gender: M Pet age: 0Yrs0Mos Owner:

Color: Yellow **Pellucidity: Clear and** **Co.: 1x**

Parameters

Detection items	Result/Unit	Result/Unit	Reference	Negative	Positive
1.Cast					
1-1.Hyaline cast	0.00/uL	0.00/LPF	0-0.8/uL	—	
1-2.Cellular cast	0.00/uL	0.00/LPF	0-0/uL	—	
1-3.Granular cast	0.00/uL	0.00/LPF	0-0/uL	—	
1-4.Waxy cast	0.00/uL	0.00/LPF	0-0/uL	—	
2.Crystal					
2-1.Struvite#	7.87/uL	0.46/HPF	0-5/uL		+
2-2.Calcium oxalate monohydrate#	0.00/uL	0.00/HPF	0-0/uL	—	
2-3.Calcium oxalate dihydrate#	0.00/uL	0.00/HPF	0-3/uL	—	
2-4.Calcium carbonate#	0.00/uL	0.00/HPF	0-0/uL	—	
2-5.Uric acid	0.00/uL	0.00/HPF	0-0/uL	—	
2-6.Cystine	0.00/uL	0.00/HPF	0-0/uL	—	
3.Cells					
3-1.RBC	15798.50/uL	310.13/HPF	0-25/uL		++++
3-2.WBC	10.36/uL	0.20/HPF	0-25/uL	—	
3-3.Renal tubular epithelial cell	0.00/uL	0.00/HPF	0-0/uL	—	
3-4.Squamous epithelial cell	3.45/uL	0.07/HPF	0-7/uL	—	
3-5.Transitional epithelial cell	0.00/uL	0.00/HPF	0-3/uL	—	
3-6.Sperm	0.00/uL	0.00/HPF	0-0/uL	—	
4.Germ					
4-1.Cocci	22.69/uL	1.34/HPF	0-0/uL		+
4-2.Bacillus	0.00/uL	0.00/HPF	0-0/uL	—	
4-3.Yeast	0.00/uL	0.00/HPF	0-0/uL	—	

Urine Morphology Report with 21 Parameters



Urine Morphology Report with 21 Parameters

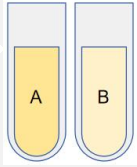
Diagnostic recommendation

- 1、【Yellow】 It is common in normal urine.
- 2、【MAP#>5】 It is common in urinary tract bacterial infection, low urine volume, alkaline urine or elevated magnesium level in diet.
- 3、【COS#>0】 There are many interference factors (such as amorphous crystallization, etc.) in bacterial detection of suspected cocci overgrowth or urinary tract cocci infection. This result is for reference only. It is recommended to confirm bacterial culture.
- 4、【RBC#>25】 Common in urinary tract bleeding (cystitis, urinary calculi, urinary crystallization).



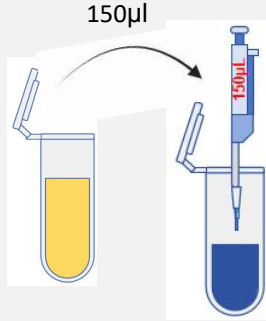


4-Steps Procedure for Feces Test



1

Use a turbidity card to determine if the sample concentration is appropriate.
(Solid feces should be dipped from variant sides by fecal wand and diluted in saline)



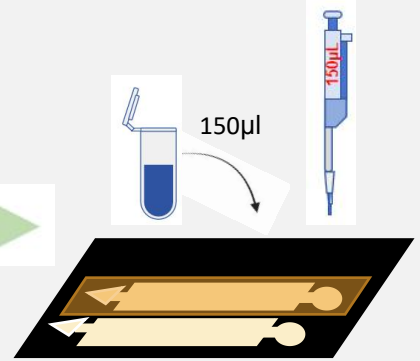
2

Draw **150µl** of the appropriately concentrated sample solution and add it to the dye solution, then invert to mix well.



3

Leave the test tube uncapped and let it stand for 1 minute.



4

Draw 150µl of the sample from the bottom of the test tube and add it to the chip for machine testing.



Getting report
in **9 min**



Feces Report with 31 Parameters



Faeces report				
No.: 231205006	LIS:	Doctor: Admin	Sample: Feces(150ul)	Time: 2023/12/05(12:07:49)
Pet name: mix	Species: cat	Gender: M	Pet age: 0Yrs0Mos	Owner:
Texture: Undefined	Smell: Undefined	Parameters		Color: Undefined
Detection items		Result/Unit	Result/Unit	Reference
1.Parasite egg				
1-1.Ascaris ascaris	0.00	0.00/LPF	0-0	
1-2.Hookworm	0.00	0.00/LPF	0-0	
1-3.Tapeworm	0.00	0.00/LPF	0-0	
1-4.Dipylidium caninum	0.00	0.00/LPF	0-0	
1-5.Sporozoa	0.00	0.00/LPF	0-0	
1-6.Alaria alata	0.00	0.00/LPF	0-0	
2.Intestinal protozoa.				
2-1.Trichomonas	2.00	0.40/LPF	0-0	
2-2.Giardia lamblia	15.00	3.01/LPF	0-0	
2-3.Isosporium coccidia	0.00	0.00/LPF	0-0	
2-4.Isosporium coccidia 0	0.00	0.00/LPF	0-0	
2-5.Isosporium coccidia 1	0.00	0.00/LPF	0-0	
2-6.Isosporium coccidia 2	0.00	0.00/LPF	0-0	
3.Germ				
3-1.Cocci	317.29/ug	28.81/HPF	145-729/ug	
3-2.Rods	7374.82/ug	650.97/HPF	516-13904/ug	
3-3.Br.evibacterium	5589.35/ug	493.36/HPF	321-12462/ug	
3-4.Crude bacilli	1490.49/ug	131.56/HPF	15-1650/ug	
3-5.Anaerofilum	294.99/ug	26.04/HPF	0-185/ug	
3-6.Cocci/Rods	0.043	0.043	0.017-0.156	
3-7.Campylobacter	2.83/ug	0.25/HPF	0-30/ug	
3-8.Bacillus	0.00/ug	0.00/HPF	0-40/ug	
3-9.Serpentine spiracheta	0.00/ug	0.00/HPF	0-0/ug	
3-10.Helicobacter	0.00/ug	0.00/HPF	0-0/ug	
3-11.Yeast	3.13/ug	0.28/HPF	0-150/ug	
4.Cells				
4-1.RBC	739.11/ug	65.24/HPF	0-5/ug	
4-2.WBC	4.88/ug	0.43/HPF	0-0/ug	
4-3.Epithelial cells	0.00/ug	0.00/HPF	0-12/ug	
5.Digestive function				
5-1.Starch granule	0.00/ug	0.00/HPF	0-9/ug	
5-2.Lipid drop	0.00/ug	0.00/HPF	0-1/ug	
5-3.Plant fiber	0.00/ug	0.00/HPF	0-0/ug	
5-4.Muscle fiber	0.00/ug	0.00/HPF	0-0/ug	
Diagnostic recommendation				
<ol style="list-style-type: none"> [FAT#1] Suspected Giardia lamblia infection. [RBC#1] Common in the colon, rectum, anus and other bleeding. [WBC#0] Common in all kinds of enteritis (bacterial, allergic, viral, parasitic, etc.), tumor and so on. [LFAT#1] Common in all kinds of enteritis, dyspepsia, pancreatic insufficiency, acute / chronic pancreatitis, pancreatic cancer and so on. 				
<p>Combined diagnosis:</p> <ol style="list-style-type: none"> [TR#6, TR#0] Suspected Trichomonas infection. The quantity is small, it is recommended to confirm by manual microscopic examination. <p>Due to various factors such as different stages of parasite infection, different parasite sites and different methods, operations and sites of specimen collection, eggs and bodies may be missed. It is recommended to reexamine fecal samples at different sites and at different times to improve the detection rate.</p> <p>—— (Methods for laboratory examination of parasitic diseases)</p>				

Faeces Morphology Report				
No.: 231205006	LIS:	Doctor: Admin	Sample: Feces(150ul)	Time: 2023/12/05(12:07:49)
Pet name: mix	Species: cat	Gender: M	Pet age: 0Yrs0Mos	Owner:

Parasite

Germs

Cells

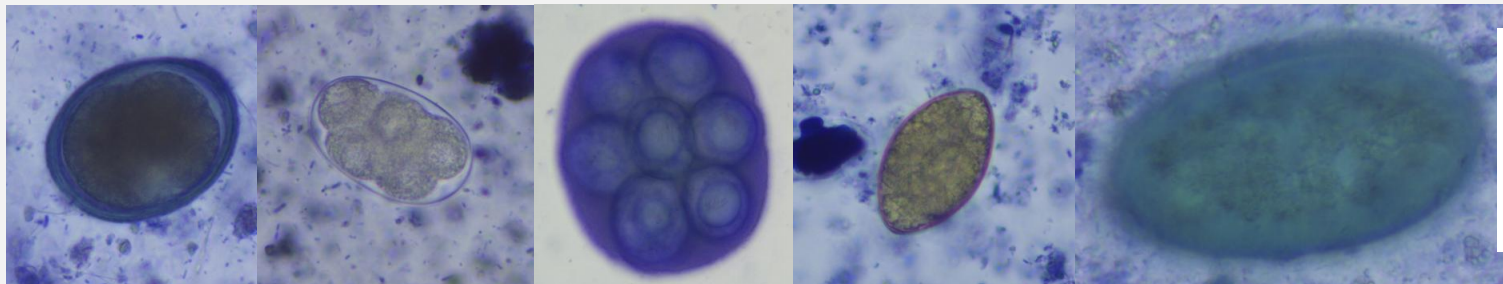
Plant fibers, muscle fibers, lipid droplets, starch granules



Feces Report with 31 Parameters

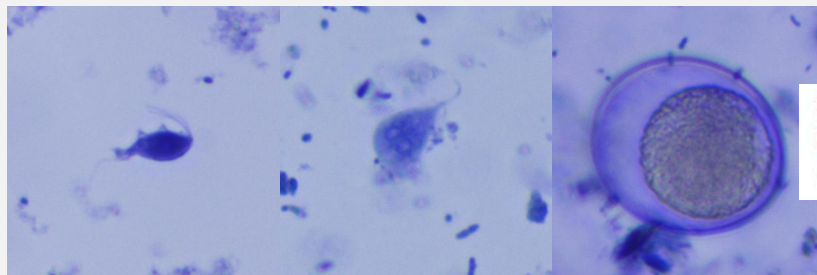


Helminth eggs: (x5)



- 1-1. *Ascaris ascaris*
- 1-2. Hookworm
- 1-3. Tapeworm
- 1-4. *Dipylidium caninum*
- 1-5. *Spirometra*
- 1-6. *Alaria alata*

OOcysts: (x3)



- 2-1. *Trichomonas*
- 2-2. *Giardia lamblia*
- 2-3. *Isosporium coccidia*



Dude, we're uncovered



Feces Report with 31 Parameters



3. Germ

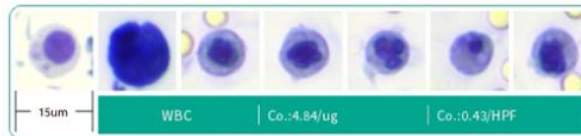
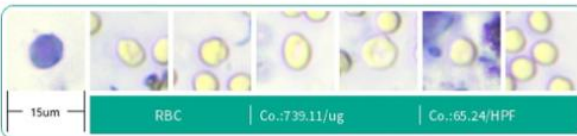
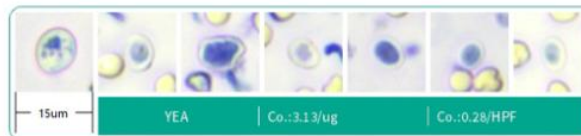
Parameter	Value	Reference Range
3-1. Cocci	317.29/ug	28.01/HPF
3-2. Rods	7374.83/ug	650.97/HPF
3-3. Brevibacterium	5589.35/ug	493.36/HPF
3-4. Crude bacilli	1490.49/ug	131.56/HPF
3-5. Anaerofilum	294.99/ug	26.04/HPF
3-6. Cocci/Rods	0.043	0.0
3-7. Campylobacter	2.83/ug	0.25/HPF
3-8. Bacillus	0.00/ug	0.00/HPF
3-9. Serpentine spirochete	0.00/ug	0.00/HPF
3-10. Helicobacter	0.00/ug	0.00/HPF
3-11. Yeast	3.13/ug	0.28/HPF



Antibiotics or Probiotics necessary?

4. Cells

4-1. RBC	739.11/ug	65.24/HPF
4-2. WBC	4.84/ug	0.43/HPF
4-3. Epithelial cells	0.00/ug	0.00/HPF



5. Digestive function

5-1. Starch granule	0.00/ug	0.00/HPF
5-2. Lipid drop	140.68/ug	12.42/HPF
5-3. Plant fiber	0.00/ug	0.00/HPF
5-4. Muscle fiber	0.00/ug	0.00/HPF



exocrine pancreatic insufficiency [EPI] Diagnostics



Diagnostic recommendation

1. 【FLA#>5】 Suspected Giardia lamblia infection.
2. 【RBC#>5】 Common in the colon, rectum, anus and other bleeding.
3. 【WBC#>0】 Common in all kinds of enteritis (bacterial, allergic, viral, parasitic, etc.), tumor and so on.
4. 【LFAT#>1】 Common in all kinds of enteritis, dyspepsia, pancreatic insufficiency, acute / chronic pancreatitis, pancreatic cancer and so on.

Combined diagnosis:

1. 【TRI#<6, TRI#>0】 Suspected Trichomonas infection. The quantity is small, it is recommended to confirm by manual microscopic examination.

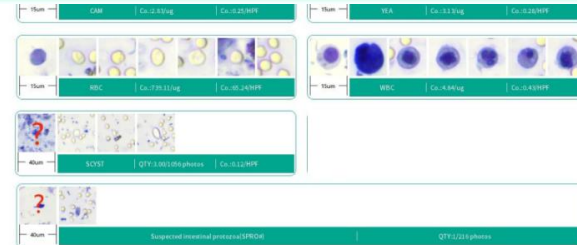
Due to various factors such as different stages of parasite infection, different parasite sites and different methods, operations and sites of specimen collection, eggs and bodies may be missed. It is recommended to reexamine fecal samples at different sites and at different times to improve the detection rate.

— 《Methods for laboratory examination of parasitic diseases》

Feces and Urine Report - Review and Edit



Automatical classify the "unknown" component for manual review and simply editing the report!

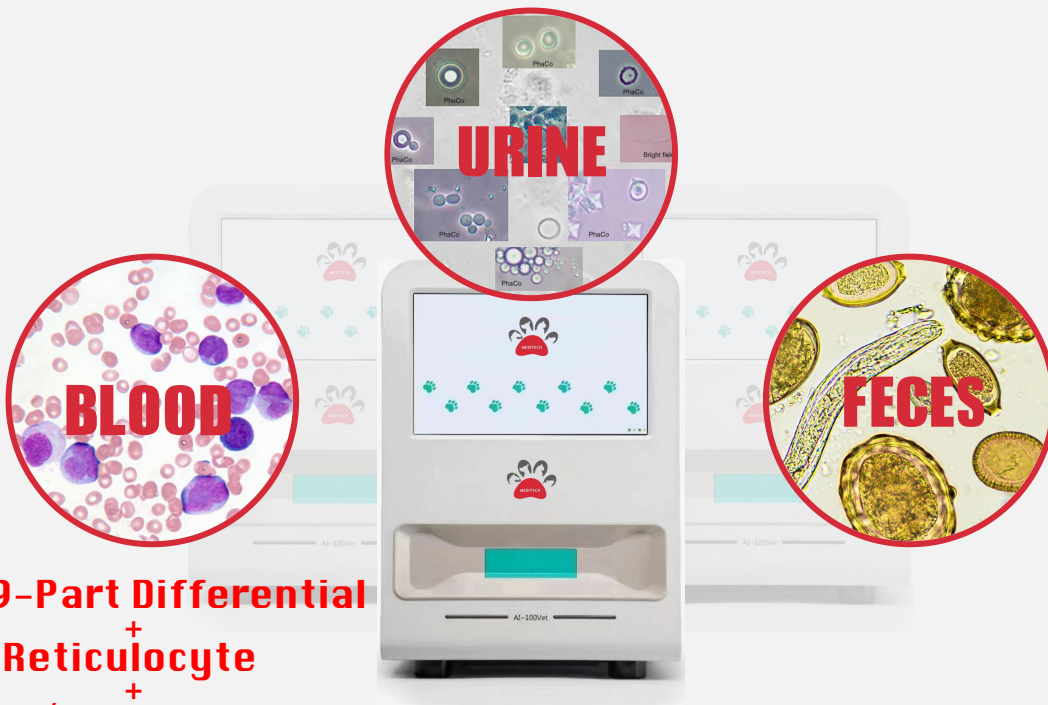




3-in-1 Innovation!



The first morphology and hematology analyzer with —



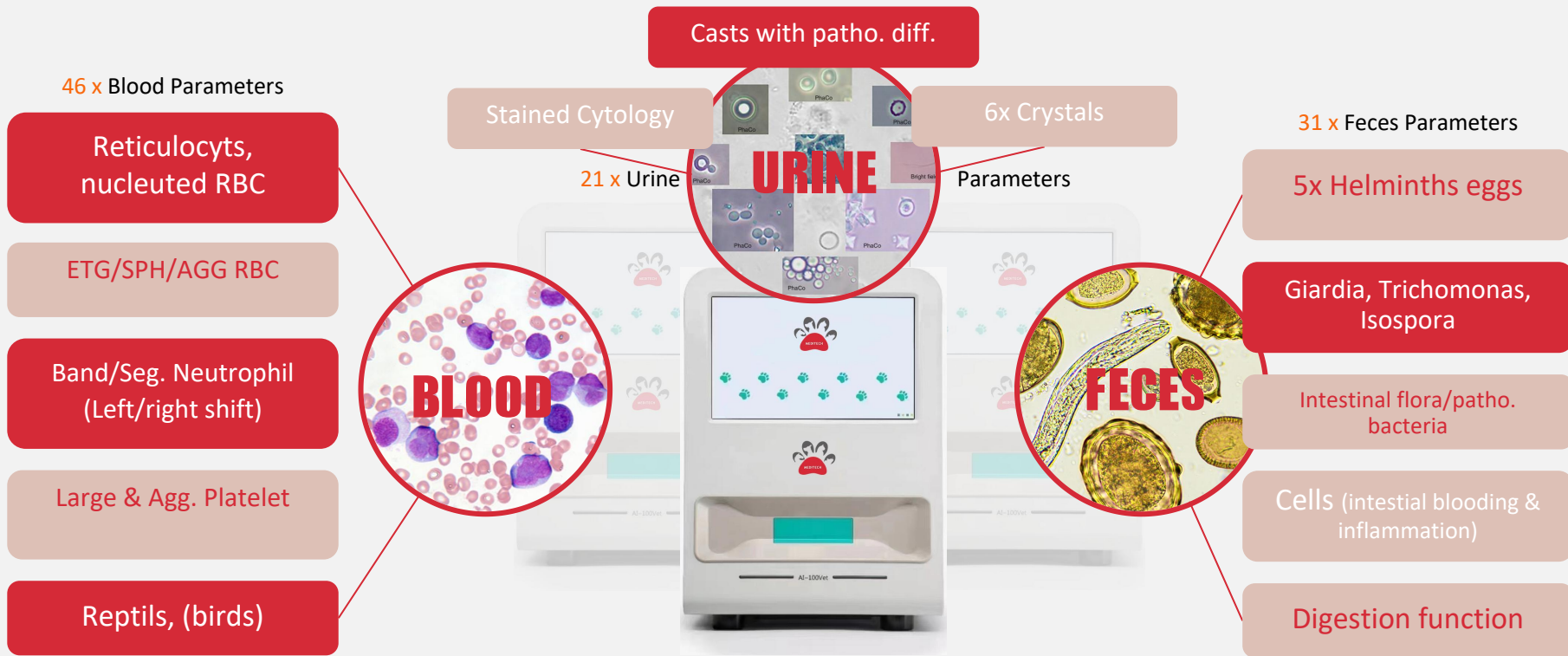
WBC 9-Part Differential
+
Reticulocyte
+
RBC U/H Differential





Special features of the 3-in-1 Innovation

What AI-100Vet Can ‘SEE’:





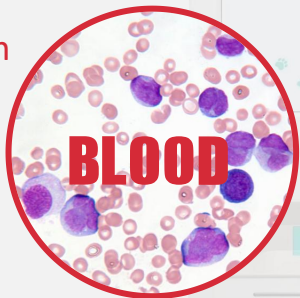
Special features of the 3-in-1 Innovation What AI-100Vet Can HELP:



Save Time
Save Labor
Save Money



- Anemia classification
- Haemolysis/IMHA screening
- Left shift (infection monitoring)
- Thrombopenia confirming
- Exotic animals



- Acute Kidney cystitis, uroliths...



- Parasites infection Dx and Tx monitoring
- Dysbiosis and/or intestinal bact. infection
- EPI screening





Convenient & Efficient!



Simple Operation



No preanalysis of samples required
 Liquid staining and loading can be done in
 3 simple steps
 Fully automated AI detection



In 7-10 minutes

Qualitative and Quantitative report
 Visual results with morphology images
 AI diagnostics interpretation available
 Review on display instead of manual

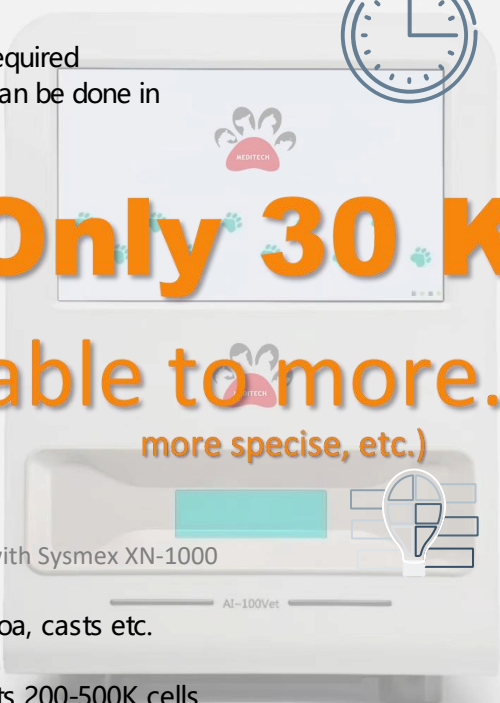
Only 30 KG

AI extendable to more... (body fluid, dermatology, more specise, etc.)



High Accuracy

Over 95% for celles (compare with Sysmex XN-1000
 and manual microscopy),
 Over 90% for parasites, protozoa, casts etc.
 (compare with microscopy),
 Observe 1000+ HPF and counts 200-500K cells



Simple Maintenance

No liquid-way
 No calibration, washing, puffering required
 Consumables in room temprature 15 months safe





Co

1 2 3 4 5 6 7 8 9 10 11



Fluid Morphology Report

No:	LIS:	Doctor:	Sample:	Owner:
Pet name:	Species:	Gender:	Pet age:	Weight:

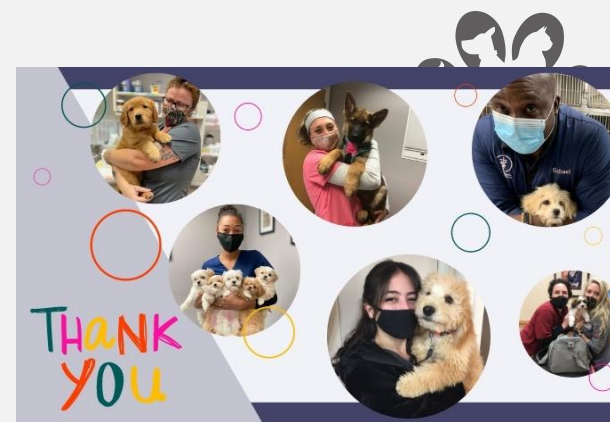
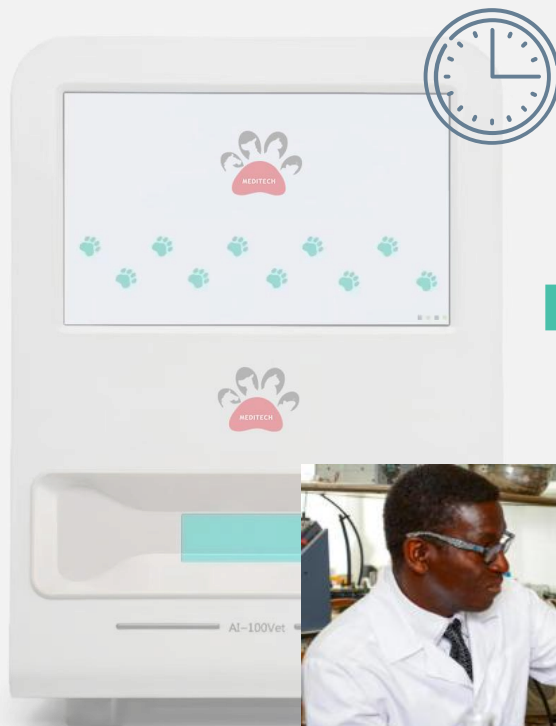
AI extendable to more... (body fluid, dermatology, more specise, etc.)



Convenient & Efficient!

Advantage Summary:

1. More parameters: more clues for vets
2. Shorter turn-around-time of diagnostics: early start of therapy
3. Saving time and labor: better hospital management and financial efficiency!



**Happy Clients
Happy Staff**





THANK YOU!



TOOTOO Meditech Co., Ltd.

Address: 3A, Phase II, Smart Home, Baolong Street,
Longgang District, Shenzhen, Guangdong, China

Tel: +86-755-28306385

Email: info@newtootoo.com

Web: www.newtootoo.com





Technical Specification of AI-100Vet - 1



Product Name	AI-100Vet		
Methology	Chamber-slide-based automated microscopy and AI morphology on liquid staining samples		
Sample type	Blood, urine, feces		
Species	Dog, cat, rabbit, chinchilla, mouse, rat, hamster, ferretm turtle		
Sample volume	Blood: 10 μ L Urine: 500 μ L Feces: 150 μ L	Testing time	Blood: 7-9min Urine: 9min Feces: 9min
Parameters	Blood(x46): WBC, NEU, NST, NSG, NSH, LYM, MON, EOS, BAS, RBC, RET, ETG, SPH, % of above, NRBC, NRBC/WBC%, AGG, HGB, HCT, MCV, MCH, MCHC, RDW/HDW-SD/CV, PLT, PCT, MPV, LPLT, P-LCR, APLT, PDW-SD/CV Urine (x21): HYA, CEC, GRA, WAC, MAP, COC, COMC, COD, CP, UAC, CYSC, RBC, WBC, RTE, SEC, TEC, SPE, COS, BAC, SAC, FAT, PHL Feces (x30): ALE, ANE, CEE, DIP, SPI, TRE, TRI, FLA, COD(0-2), COS, BACI, SBAC, CBAC, TBAC, C/B, CAM, BAC, SS1, SS2, YEA, RBC, WBC, EPC, STA, LFAT, PLA, AF		





Technical Specification of AI-100Vet - 2



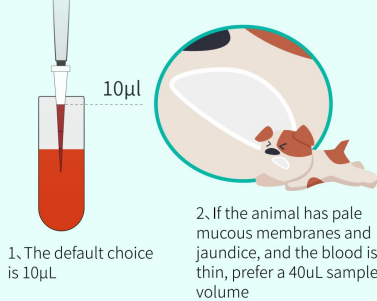
Product Name	AI-100Vet
PC	Integrated, Windows 10
Display	Integrated touch screen, 12 inch LCD (1280x800dpi)
Data connection	USB(3.0), Wlan, network cabel
Power supply	110-240 VAC, 50-60Hz, 650VA
Language	English, Spanish, Chinese
Size	485x485x335 (mm)
Weight	30.8kg



Correct Pipetting of Blood Samples

Step1

Before starting the test, confirm the blood sample volume as follows:

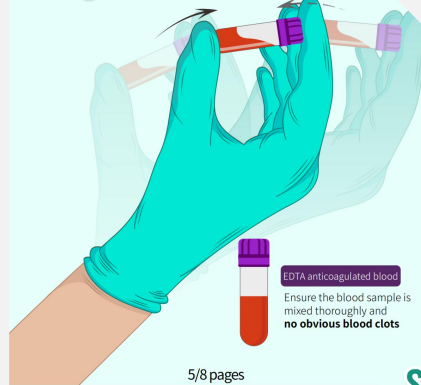


1. The default choice is 10 μ L

2. If the animal has pale mucous membranes and jaundice, and the blood is thin, prefer a 40 μ L sample volume

4/8 pages

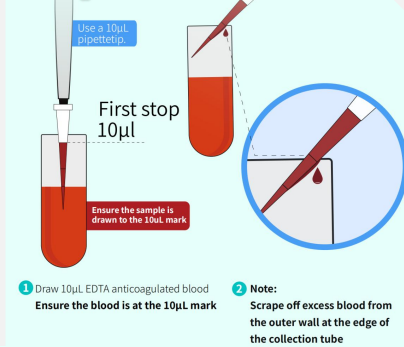
Step2



EDTA anticoagulated blood
Ensure the blood sample is mixed thoroughly and no obvious blood clots

5/8 pages

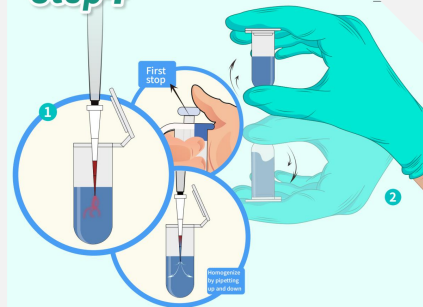
Step3



1 Draw 10 μ L EDTA anticoagulated blood
Ensure the blood is at the 10 μ L mark

2 **Note:**
Scrape off excess blood from the outer wall at the edge of the collection tube

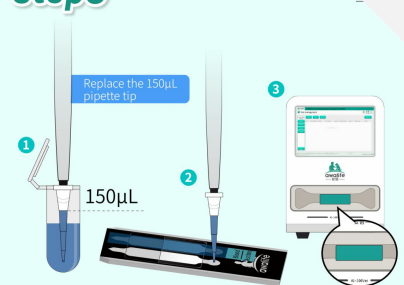
Step4



Add the blood sample to the staining solution. Dip the pipette tip into the liquid surface to dispense the sample. Then, gently pipette up and down 5-10 times to homogenize before inverting and mixing.

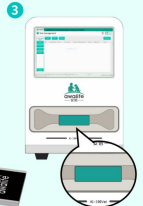
7/8 pages

Step5



1 Replace the 150 μ L pipette tip

2 150 μ L



After mixing thoroughly, immediately draw 150 μ L of the stained sample, quickly dispense it into the chip, then proceed to machine testing

3 **Note:**
If the mixed solution is left for more than 15s before adding, it needs to be removed before adding to the chip.

8/8 pages

**Fresh
Dog/Cat
Blood**



Correct Preparing of Urine Sample



Step 1



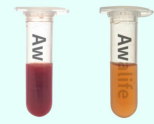
Over-Concentrated Samples Ideal Concentration

1 High-Concentration Samples (Bilirubinuria Sample, Highly Turbid Urine)

Manual microscopy is recommended. If using AI-100Vet, dilute the urine with saline to below the threshold concentration, discard the supernatant, and use the bottom sample, though results may not be reliable.

2 High-Concentration Samples (Hematuria Samples)

Manual microscopy is recommended. If using the AI 100 VET, dilute the urine with saline to below the threshold concentration, discard the supernatant, and use the bottom sample, though results may not be reliable.

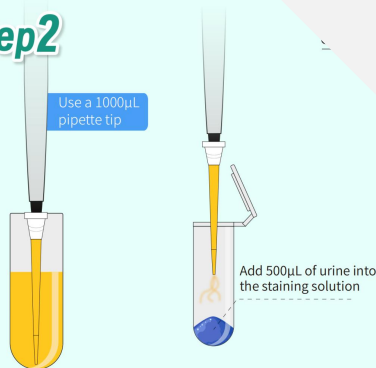


Over-Concentrated Samples Ideal Concentration

3 Normal Urine Samples proceed to the next step

2/5pages

Step 2



Add 500µL of urine into the staining solution

Draw 500µL of urine from the bottom of the test tube and add it to the staining solution

3/5pages

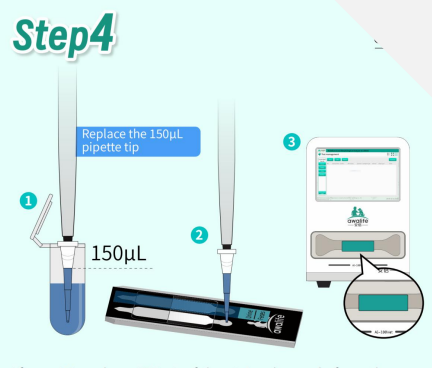
Step 3



Thoroughly mix the urine and staining solution by inverting until there is no visible dye residue at the bottom of the tube

4/5pages

Step 4



After mixing, draw 150 µL of the stained sample from the bottom of the tube, quickly dispense it into the chip channel in one go, and then proceed with the machine detection.

5/5pages





Correct Preparing of Feces Sample



Fecal Turbidity Card

Place the Tube Here

Highest Turbidity Lowest Turbidity

Check the transparency of the sample to confirm if it is between the highest and lowest turbidities

Place the fecal dilution tube in the turbidity card sample placement area, observe the transparency, and confirm whether the sample concentration is between the highest and lowest concentrations

2/7pages

Use a 150 μ L pipette tip

150 μ L

1

2

Draw 150 μ L of the fecal sample and add it to the staining solution, invert and mix 5-10 times

5/7pages

Use a 150 μ L pipette tip

150 μ L

1

2

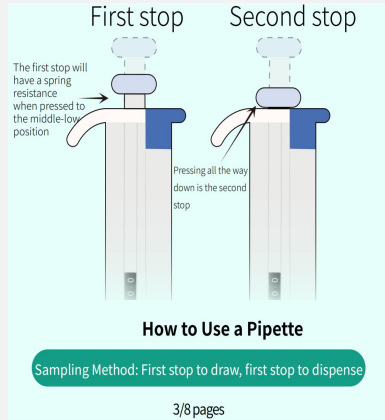
3

After settle down for 1 minute, immediately draw 150 μ L of the bottom stained sample, quickly inject it into the chip in one go and then proceed with the machine testing

7/7pages



Attention



Common Error:

HCT too high:

- too much sample is added (pipette tip attached, pipette pressed too much)
- sediment RBC are aspirated (not well mixed anemia blood)
- aspirated and dissociated blood clots

HCT too low:

- not enough sample (bubbles, blocks, tip cleaning error)
- supernatant are aspirated (not well mixed anemia blood)
- blood clots

Solution: repeat AI-testing → repeat operation repeat → repeat with heparin blood

Anemia blood:



Sample Volume: 10ul 40ul 80ul

Mixing and immediately testing!!!



Attention



Step 1



Over-Concentrated Samples Ideal Concentration

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Manual microscopy is recommended. If using AI-100Vet, dilute the urine with saline to below the threshold concentration, discard the supernatant, and use the bottom sample, though results may not be reliable.

2 High-Concentration Samples (Hematuria Samples)

Manual microscopy is recommended. If using the AI 100 VET, dilute the urine with saline to below the threshold concentration, discard the supernatant, and use the bottom sample, though results may not be reliable.



Over-Concentrated Samples Ideal Concentration

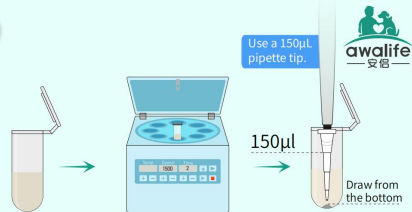
3 Normal Urine Samples proceed to the next step

2/5pages

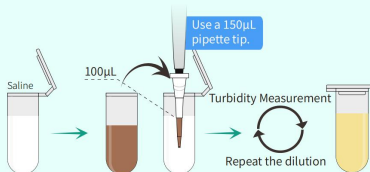


Step 2

Compare the concentration



If the concentration is too low, continue to add samples to about 0.5mL, centrifuge at 1500 rpm for 2 minutes, and take 150µL of the sediment for staining

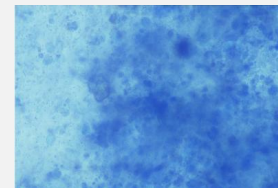


If the concentration is too high, dilute the sample in the fecal turbidity tube with physiological saline and repeatedly compare the transparency until the sample concentration is between the highest and lowest concentrations

3/7pages



Filling Status



Focusing Images



18-min Enhance Mode for Urine and Feces

sample volume 150ul

Standard mode (approximately 9 minutes)

Enhanced mode (approximately 18 minutes)

Texture

Dry feces Formed stool Soft stool

Loose stools Watery stools

Smell

Smelly Fishy odour Foul smelling

Color

Brown Yellow Red

Black Milky white

- ✓ Enhanced mode takes double so many FOV images as standard mode
- ✓ It takes photo from the edge of the chamber for parasites eggs

More important question for solid fecal sample:

is the **sampling** homogeneous??

➤ total sample dilution
or

➤ 5-10 different position sampling

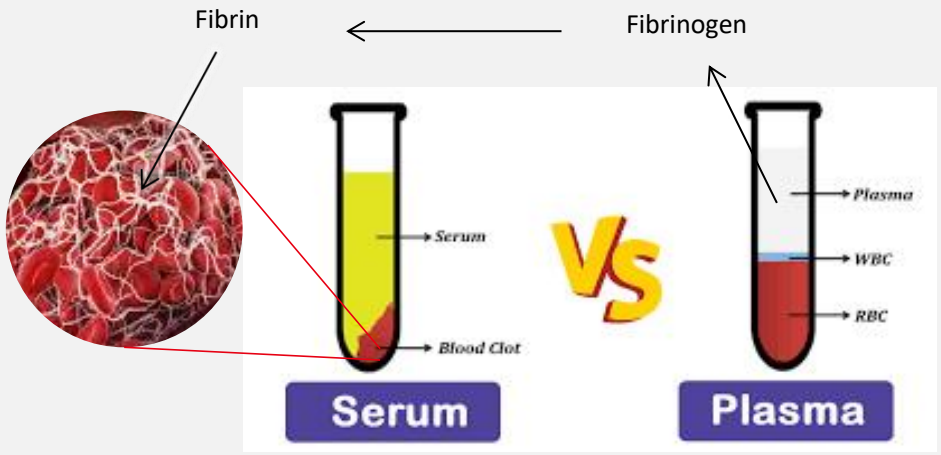
Cave: pus inclusive parasites eggs

→ shake!





• Cave: Plasma VS Serum



In-vivo: Whole blood = plasma+blood cells
 In-vitro:
 With anticoagulant: plasma+blood cells → hematology, PCR, coagulation
 Without anticoagulant: serum+blood clot → biochemistry, immunoassay

ANTICOAGULANTS:



EDTA-blood:
 Mammals' hematology
 Blood type
 Blood smear
 Coomb's test



Heparin-blood:
 Reptiles & birds' hematology
 Small mammal animals



Sodium-citrate-blood:
 Coagulation test
 (blood:citrate 1:9 mixing)



Sodium-fluoride-blood:
 Glucose and lactose test

