Postanalytical phase



POSTANALYTICAL PHASE

Test request





Black box: the lab





And the RESULT is created

The technician approves the result; it is transferred to the lab informatics system

Graduated staff monitors and compares the lab results to each other and other clinical data (delta check: relationship to prior results).

(S)he validates, if they are ok.

Validated results = the finding.

ONLY VALIDATED RESULTS CAN BE USED FOR CLINICAL DECISION MAKING

And then the findings are created

- Sent via the informatics system
- Printed form

Data	NOTE
Patient's name and identifier	Items required depends on lab. Finding without name / identifier cannot be used for clinical decision
Name of the analyte measured	There are several analytes on one finding; these are listed consequtively
Result	Most often a number, rarely a note (positive, negative)
Parameter (unit)	Result without parameter cannot be used
Healthy reference range	or 'normal value'.
Comment	Any comment on test or test specimen (eg. hemolyzed, or few amount), any interpretation of result
Lab performing the test	Private or state
Name and identifier of validating staff	In general a doctor or clinical biochemist

The question of units

- The units are not universal
- There are conventional and SI units
- UK, AUS, NZL, CAN, HU, NL SI units (mmol/L)
- USA, D, ISR, JA conventional units (mg/dL)
- The value is meanless without the unit.

Some examples: the electrolytes

analyte	SI	conventional
se Na, se K, se CI,	1 mmol/l	1 maeq/L
se HCO3-		
se Ca	1 mmol/L	4 mg/dL or 2
		maeq/L
se Mg	1 mmol/L	2.4 mg/dL
se P	1.2 mmol/L	3 mg/dL

A characteristic example: the calcium

Let's say: serum calcium is 3.05.

Is it high, normal or low?

Depends on the unit used:

Total calcium, reference range:

 $2.3 - 2.6 \, \text{mmol/L H}$

4.6 - 5.2 maeq/L L

9.2 - 10.4 mg/dL LL

Some examples: chemicstry & haematology

SI (System	International)	Conventional units
Blood Hgb	148 g/L	14.8 g/dL
Total protein	66 g/L	6.6 g/dL
Se Glucose	4.3 mmol/L	77 mg/dL
BUN	25 mmol/L	70 mg/dL
Se Chol.	5.5 mmol/L	212 mg/dL
Se TG	3.3 mmol/L	292 mg/dL
Creatinine	88 <i>μ</i> mol/L	0,99 mg/dL

Some examples: chemicstry & haematology

SI (System International)	Conventional units
Se Bilirubin 50 µmol/L	2,9 mg/dL
Lactate 2 mmol/L	18 mg/dL
Ammonia 40 μmol/L	68 μg/dL
Se T4 total 113 nmol/L	8,8 μg/dL
Se T4 free 12 pmol/L	0,9 ng/dL
Bacterial CFU 109/L	10 ⁶ /mL

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Interpretation of test results

Common interpretations:

- 'negative'
- 'Bad / good'
- 'Normals'
- XY is increased

Problem:

- Healthy reference value? (may be age, labor and population specific!)
- Preanalitical problems are not considered ('the labor works ugly')
- The meaning of the test is not taken in account

First question: compare to with

- Matched age?
- Same gender?
- Documentedly healthy?
- Not affected from the investigated condition (but still unhealthy)
- What conditions are allowed to have a control subject?
- Size of control population?
 (answers depend on a number of factors)

Interpretation of test result

Important terms:

SENSITIVITY =

Positive / Total number of patients

$$sensitivity = \frac{number\ of\ True\ Positives}{number\ of\ True\ Positives + number\ of\ False\ Negatives}$$

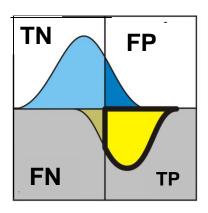
SPECIFICITY =

Negative / Total number of healthy

$$specificity = \frac{number\ of\ True\ Negatives}{number\ of\ True\ Negatives + number\ of\ False\ Positives}$$

sensitivity

Rate of patients with true positive result (se)

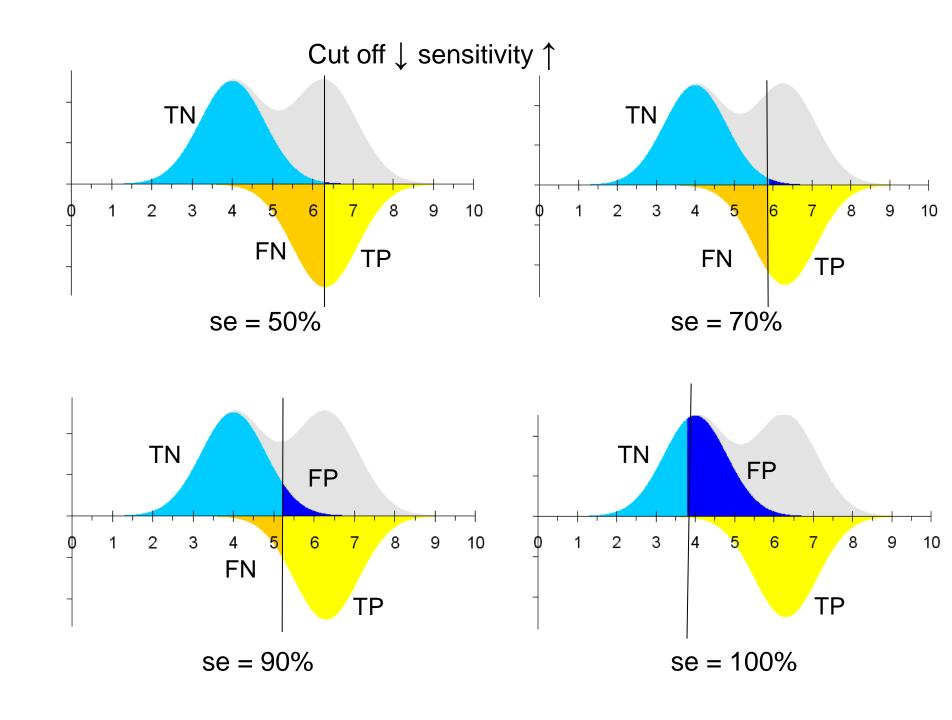


= the probability of positive test result in a patient

$$= TP / (TP + FN)$$
Se

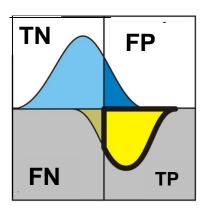
For screening high sensitivity test is required (i.e. to miss just a few patients)

(Sensitivity should be lowered if there is no sufficient resource for treatment of identified cases)



specificity

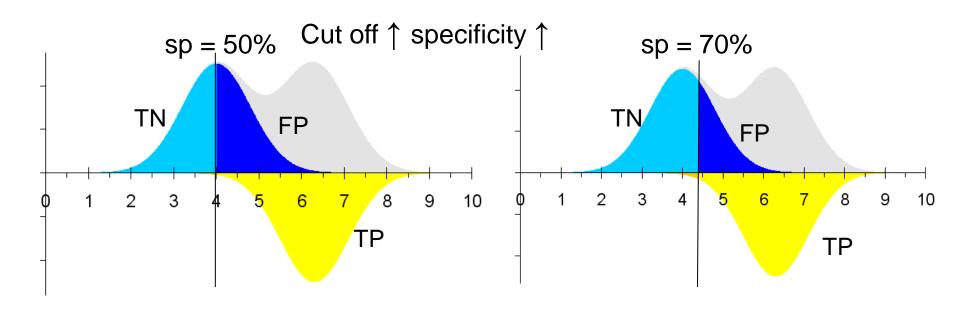
Rate of healthy subjects presenting true negative results

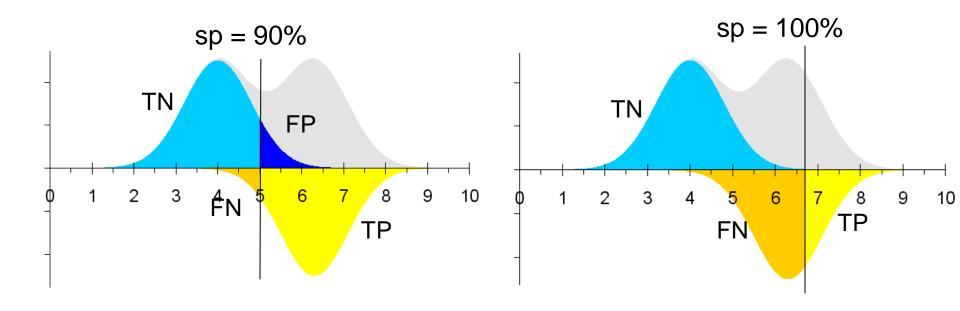


= probability of having a negative test result in a healthy subject

High specificity is required when false positive results should be disclosed.

(Usually required for confirmatory tests)





Predictive values

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Prevalence of predictive value of a lab test depends on prevalence of Analogies

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( Screening for severe congenital metabolic disorders

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 Negative predictive value: if test result is negative, then what is the probability that the tested subject is not affected?

negativ predictive value (NPV): TN/(TN+FN)

Depends on prevalence in population

Predictive value

- Importance of PREVALENCE
- The rate of healthy and diseased subjects have an impact on test PPV and NPV values.
- That means: the information provided by a given test depends on characteristics of tested population
- If I use even a very high sensitivity test (but there is no patient in the tested population), then I will get false positive results only

An example

- HIV-test: both specificity and sensitivity are around 0,99
- If 20000 HIV infected patients are in Hungary, the prevalence of HIV infection is 0.002.
- Let's suppose that prevalence of HIV positivity among blood donators is the same (in fact it is lower)
- Question: what will be the rate of false positive results amon blood donators?
- SP:0,99, SN:0,99, PR:0,0020,

The PPV

$$\frac{0,99*0,0020}{(0,99*0,0020) + (0,01*0,9980)} = \frac{0,00198}{0,01196} = 0,165$$

That means that out of 100 positive (reactive) samples just 17 confers to HIV seropositive blood donators, and 83 results are false positive (In fact this rate is much more higher).

The real NPV

$$\frac{0.99*0.998}{(0.99*0,002) + (0.99*0,998)} = \frac{0.988}{0.99} = 0,998$$

Therefore, the risk of obtaining a false negative result is very-very low.

Receiver Operating Characteristics (ROC) curve

In case of dichotomic results the cut-off value is clear (usually). In case of continuous results the cut-off calue should be established carefully.

The callenge: to identify cut-off value to discriminate positive and negative with the best specificity and sensitivity (ROC curve)

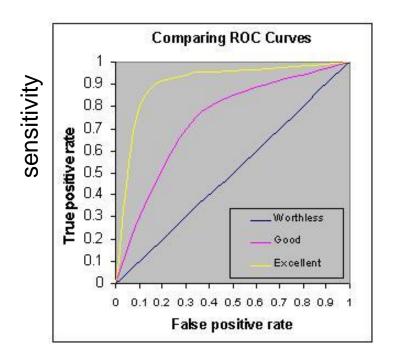
ROC curve:

 Indicates the true positive rate and the false-positive rate at difference cut-off values

(i.e.: compares sensitivity to 1- specificity values)

- Appropriate to assess clinical utility of a given test
- Supports the selection of the most appropriate cut-off (reference) value

ROC curve



Area under the curve 0.9-0.99 = excellent 0.8-0.9 = good 0.7-0.8 = moderate 0.6-0.7 = fair < 0.6 = failed

1-specificity

The test is more accurate whe the curve tends to follow left and up border. The clinical utility of test is lower if the curve's slope is nearer to 45°.

Finally: I have a reference range

- The referencia range has too (lower and upper) limits.
- In some cases data are normally distributed
- In other cases they are skewed to the left or the right
- Abnormal = any value outside the range.

Examples when abnormal is below the ref range:

- Vitamine levels
- Protein levels (non-acute phase)
- CBC

Examples when abnormal is above ref. range:

- Tumor markers
- Markers of tissue necrosis
- Some hormones
- CBC

Some specific reference ranges

- Pediatric values
- Gender-specific values
- Cycle-dependent ranges
- Diurnal variations
- In some cases reference values are changing dramatically and just relative values can be given

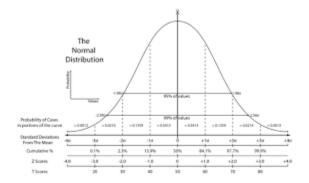
Percentile

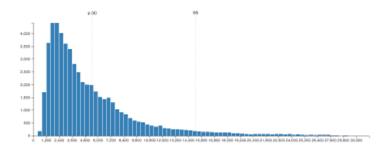
MoM

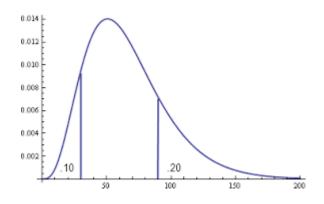
T-score és Z-score

Special reference ranges

Percentile

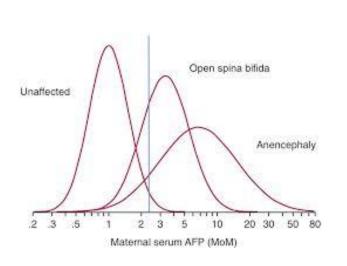


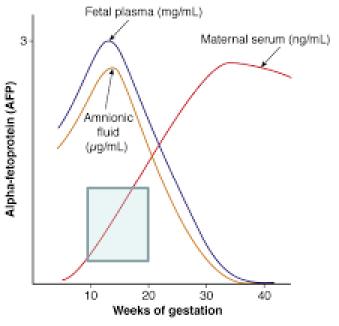




Special reference ranges

MoM





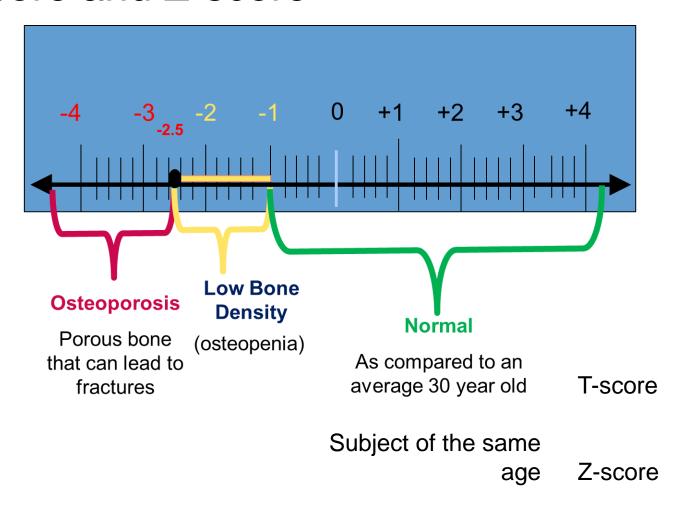
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MoM = multiples of median

For those cases when reference range changes very quickly in time (e.g pregnancy or cancer)

Special reference ranges

T-score and Z-score



Reference values

Cut-off value

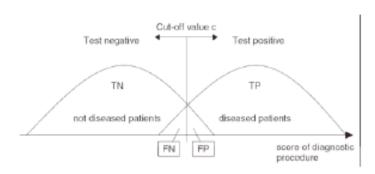
General principle: measured values in 95 (90 – 99)% of healthy subjects should be within the healthy reference range.

The probability is 95% that any analyte is within the healthy reference range.

If 20 analytes are measured in a (healthy) subject, just $0.95^{20} = 0.36 \rightarrow 36\%$ is the probability that each analyte is normal.

If the specificity 99% for each, just 81% is the probability that each analyte is normal.

(i.e. the principle 'how to make a diseased person from a healthy one')



And now, let's see what is happening in real life

- Emergency lab tests
- Alarm values

Some remarks on emergency lab tests

Sürgős lapon feltüntetendő:

Beteg adatai (név, szül. dátum, TAJ szám):

Diagnózis: BNO-kód:

Vizsgálatkérő intézet	Vizsgálatkérő osztály	
Vizsgálatkérő orvos:	Telefonszám (ha nincs, nem kerül az krizis	
	eredmény bemondásra):	

Mintavétel dátum & időpont (óra, perc): Vizsgálati minta tipusa:

Nativ vér	EDTA-s vér	Vizelet	
NaF-os vér	Citrátos vér	egyéb	

Glüköz	T.bilirubin	iCa **	VIZELETVIZSGÁLAT TESZTCSÍKKAL
Na, K, Cl	D.Billirubin	Ammónia*	
Kalcium	ALP	Laktát*	ÁLTALÁNOS VÉRKÉP
Foszfát	GOT	Béta-HCG**	
Osszfehérje	GPT	Troponin T	VÉRCSOPORT
Albumin	GGT		
Kreatinin	LDH	haptoglobin**	VERGAZ*
Karbamid	Szérum amiláz		SZÉKLET VÉR
Húgysav	Vizelet amiláz		
Szérum ozmolalitás	CRP		LIQUOR fehérje, glükóz, sejtszám
Vizelet ozmolalitás	Prokalcitonin		

GYÓGYSZERSZINTEK SZÉRUMBAN

Metotrexit	Digoxin	Fenobarbitál	Valproát	Litium	
Vankomicin	Karbamazepin	Fenitoin	Drog- teszr**		

HEMOSZTÁZIS VIZSGÁLATOK Kitöltendő, hogy milyen specifikus kezelést kap a beteg

K-vitamin antagonista:	Nem frakcionált heparin:	Klopidogrel:	Direkt trombingátló:	
LMWH	Aszpirin:	Fibrinolizis:	Rivaroxaban:	
Egyéb kezelés, azaz:				

Protrombin idő	Trombin idő	D-dimer	H	eparinszine***	
APTI	Fibrinogén	Antithrombin III.	R	livaroxaban***	

^{*} speciális csőbe veendő; ** külön telefonos egyeztetés alapján; *** csak abban az esetben, ha összhangban áll a beteg gyógyszerelésével



Some remarks on emergency lab tests

- Vital lab tests
- These include: toxicity, electrolyte levels, metabolic disturbances, tissue necrosis
- TAT : < 1 hour
- Technicians provide them without validation by graduated staff ('intermediate')
- Restricted lists (depends on institution)
- More expensive and need more efforts

The findings are generated

Uncle Pete's findings

Lab findings asked as emergency

Available after 43 min of TAT

- ALP: 1100 U/L
- Bilirubin: 80 micromol/L
- Direct bilirubin: 54 micromol/L
- LDH: 340 U/L
- Not increased: amilase, troponin, GOT, GPT
- WBC: 10.4 G/L
- Urine: Ubg negative, bilirubin positive

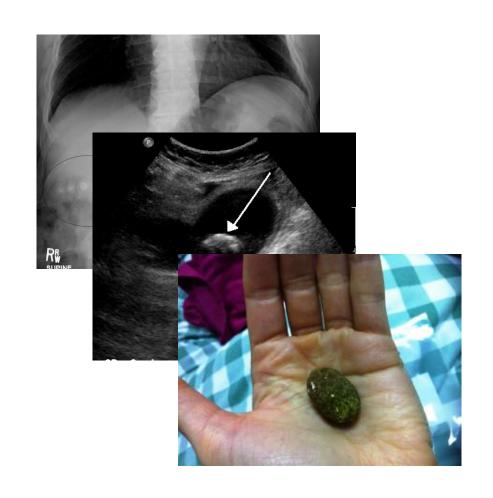


Imaging findings

Diagnosis:

Gallstones
(with a still opened pancreas duct)

ERCP



Findings of repeated testing

- ALP: 540 U/L
- Bilirubin: 50 micromol/L
- Direct bilirubin: 24 micromol/L
- Normal: amilase
- WBC: 7.5 G/L

- Critical values that should be communicated to the doctor immediately
- Require immediate clinical intervention
- There is no unique guide, depends on environment (lab & department)

CBC				
		less than	more than	
WBC	G/I	1,5	20,0	
RBC	T/I	2,0	6,0	
Haemoglobin	g/l	65	200	
Hematocrit	1/1	0,25	0,6	
Platelet count	G/I	50	1000	

Hemostasis tests				
		less than	more than	
APTT	sec.		100	
			3,5 (ambulatory)	
Prothrombin time	INR		4 (admitted)	
Fibrinogen	g/l	0,5	6	
D-dimer	mg/l		20	
Antithrombin-III activity	%	50		

		less than	more than	comment
Amilase	U/I		1000	Pancreas
Ammonia	µmol/l		60	metabolism
Digoxin	μg/l		2,5	drug
Phenytoin	μg/ml		25	drug
Phenobarbital	μg/ml		40	drug
Phosphate	mmol/l	0,5	3,0	metabolism
Glucose	mmol/l	2,0	20,0	metabolism

Clinical chemistry and drug levels II				
		less than	more than	comment
GOT (ASAT)	U/I		1000	liver
GPT (ALAT)	U/I		1000	liver
Uric acid	µmol/l		750	metabolism
calcium	mmol/l	1,5	3,3	electrolyte
Potassium	mmol/l	3,0	6,0	electrolyte
Carbamazepine	μg/ml		12	drug
BUN	mmol/l		20	kidney
Creatinine	µmol/l		400	kidney
Creatine kinase (CK)	U/I		1000	heart

Clinical chemistry and drug levels II

		less than	more than	comment
LDH	U/I		1000	lysis
Lipase	U/I		700	Pancreas
Lithium	mmol/l		1,5	drug
Magnesium	mmol/l	0,5	1,5	electrolyte
Sodium	mmol/l	120	155	electrolyte
Procalcitonin	μg/l		10	Szepszis
Osmolality	mosm/kg	250	325	electrolyte
Troponin T	ng/l		300	Szív
Valproate	μg/ml		150	drug
Vancomycin	μg/ml		80	drug
Blood pH		7,1	7,6	metabolism

Alarm values – do remember

- Value reported verbally is of no legal power
- The reporter's name and position should be documented
- The basis of clinical decision should be the written findings
- Legal issue: as lab report has just partial impact on clinical decision making, the lab's responsibility is limited

Data and remnant samples

Issues:

- Ownership (sample and data)
- Who and how could treat the results?
- Issues of data saving
- How to destroy (or store) samples

What can I do with the sample?

- Thumb's rule: diagnostic samples can be used only for treatment purposes
- The use of sample for other purposes requires ethical permission (exception: method development on anonymous samples)
- Biobanks can be established when patients' consent is obtained

A few words about costs

- Lab testing costs money (reagent, salary, consumables, maintenance of lab)
- Costs of lab are calculated using a central list in Hungary
- In general, costs are paid by the physician / hospital
- These may be quite high, but just a small portion of total health care costs (2-4 per cent)
- The most expensive tests are those that are done in an unjustified manner

Summary I

- Postanalytical phase: the phase when the lab test results are used
- Lab tests have their specificity and sensitivity; these are tests' caharacteristics
- The positive and negative predictive value of the test depends on prevalence either
- The selection of the appropriate reference group is of critical importance
- The higher the number of lab tests the higher the risk of false positivity

Summary II

- Emergency tests are available 24/7; the list is restricted
- Alarm values: depend on environment.
 This case the lab notifies the health care staff immediately
- One should pay for lab tests
- Most expensive tests are those that are ordered / interpreted improperly