

Epidemiology

تلخيص محاضرات د.أشرف
Lectures (20 , 21 , 22)
instrument , screening &
validity



التلخيص هو عبارة عن كتابة كلام الدكتور في شرحه
للمحاضرات ، يعني معلومات المحاضرة اللي في السلايدات
ولكن بالطريقة والترتيب اللي عرضهم فيها الدكتور

هذه الأداة لهذا البحث (هذا المقياس) ⇒ Validity

نتيجة يوثق عليها ⇒ reliability

ثبات النتيجة والقرارة على الأداة والتي هي أهم من عدد الاستجابات شرطية أن لا تتغير العوائل. ⇒ Consistency

* Validity and its reading = reliability are the 2 roles in instrument in researchs, so you have to describe instrument carefully in your established research.

* Gold standard are instrument which do 100% every time it ~~meas~~ in measurements, which are valid and reliable.

* Gold standard are ~~so~~ highly cost & time consuming, complicated to use, so in screening & survey it is not good.

* in screening we use another instrument that are not gold standard, its validity are less, so it will produce

1. True Positive
2. False negative } ⇒ in already ^{Positive} ~~negative~~ readings

1. True negative
2. False ~~negative~~ Positive } ⇒ in already negative readings

* Screening are related to detection of disease or condition in Preclinical Phase, Patient looks healthy and that is a difficult in screening

* screening for diseases are:

1. important disease

2. Dangerous ⇒ which has dangerous complications if occur

3. has well-defined stages ⇒ mainly Preclinical Phase.

4. high prevalence ⇒ ~~the~~ main aim of screening to low the prevalence.
↳ So you don't make screening for rare diseases.

* if you measure by questionnaire without instrument, it is **survey**.

* screening done by instrument.

* screening elicit disease ~~during~~ⁱⁿ Preclinical phase, not during disease phase.

* screening are not measure incidence because it doesn't ~~measure~~ related to cases, it applied at preclinical phase.

* survey will measure incidence, it ~~is~~ applied on diseased.

* we don't use gold standard instrument except in research centers, some times it will be dangerous as well as it doesn't used except by expire, also it is expensive

* Validity = 100% Positive

* reliability = 100% Negative

* True Positive = sensitivity

~~* specificity =~~

* false negative = specificity

describes
instrument
which are
not gold
standard.

* series

* Sensitivity = validity in non gold standard

specificity = reliability in non gold standard

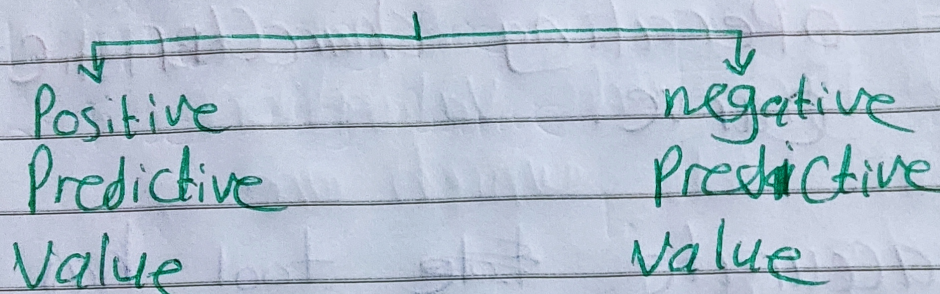
* most of screenings done by
non gold standard.

$$\text{Sensitivity} = \frac{\text{Number who test positive with disease (a)}}{\text{Number with disease (a+c)}}$$

$$\text{Specificity} = \frac{\text{Number who test negative without disease (d)}}{\text{Number without disease (b+d)}}$$

* any instrument will give result, the important thing for you to ensure the result collected by instrument other than gold standard are correct result, this is known as

Predictive Value



$$PPV = \frac{\text{Number of +ve with disease}}{\text{Number who +ve test}} \quad NPV = \frac{\text{no. who -ve test without disease}}{\text{No. who tested -ve}}$$

* when you establish your research, you have to know that the journals has degrees according to degree of methodology in research, mainly tool which has been used

$$\text{Positive predictive value} = \frac{\text{Number who test positive with disease (a)}}{\text{Number who test positive (a+b)}}$$

$$\text{Negative predictive value} = \frac{\text{Number who test negative without disease (d)}}{\text{Number who test negative (c+d)}}$$

* accuracy \Rightarrow دقة النتائج (في حساب True Positive & True Negative)

* the plot on which the accuracy = relation between specificity & sensitivity are known as Receiver operating characteristic (ROC) curve.

* The accurate tool has ROC value ≥ 0.8 (equal or more 0.8)

* before using any tool in research for data collection to calculate incidence or prevalence for any condition, you have to know five informations about your tool:

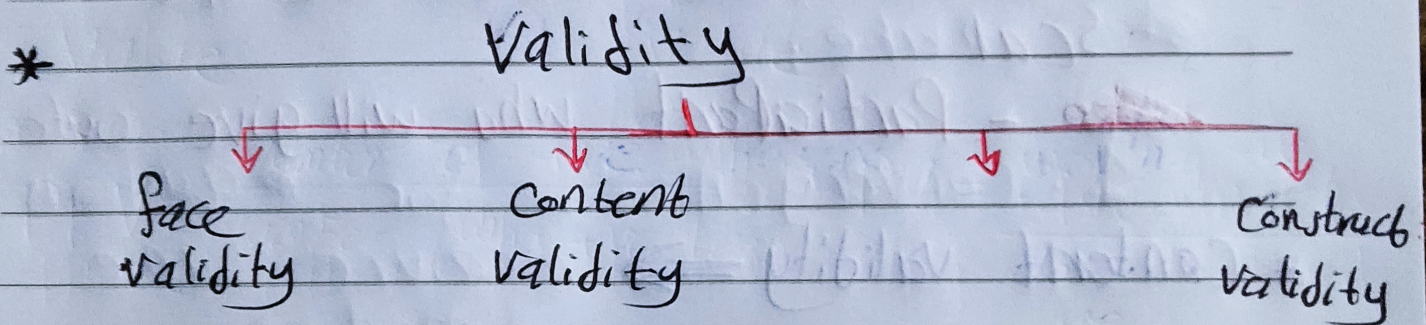
~~validity~~

~~reliability~~

الدكتور ما ذكر ال 5

* one of most important steps in research to get correct results and relation between case and other factors, is to get good sample.

- getting sample and measurement some thing related to tool which used.



* tool which you use may be instrument or other as questionnaire

* Types of validity are done according to the type of the instrument you are going to choose the root in order to validate your questionnaire.

* Face validity →

بتجيب أسئلة معينة كالأمر تكون
من مصادر أو يختار مجموعة من الفئة التي
يدرك خبرتها، أما بما وجوا عليها وسألهم
أنت هذي الأسئلة بتقدر أنها كانت
بتقريباً إرتى بالضبط؟؟

- lowest type of validation
- used in tools which are as
Scale
- Participant who will give outcomes

* Content validity →

- used in tools and instruments
which as scale

* نخبرنا التي قبلها كبيت من صحتها
واعتمد فيها على نفس الموضوع (subject)
هذي واحد خبر يراعى ال feedback
تعتبر الأسئلة وهو يوطئ نسجته

- expert who will give out comes

* Kappa Statistic →

- used to make statistical results from questionnaire by ~~effect~~ content validity.

* Criterion-related validity →

Criterion → القياس الذي نقيسه
what are you going to measure by this instrument.

Criterion

Predictive

Concurrent

↓
The ability of questionnaire or instrument to make prediction related to state which measure, but at different times

↓
The ability of questionnaire or instrument to make prediction related to state which measured when it measured by new & old tools at same time

* Construct validity →

- Purely statistics for questionnaire
- Not for instruments, questionnaire only
- Questions as same for samples that measure same dimension
- Each Psychiatric concept has degrees and dimensions.

~~make~~ make:

1. Convergen → for # questions measure same dimension

2. Divergen → separate each dimension from another.

مثلاً (بيني) أنت مثلا السعادة فيها تفاصيل مثلا
سعادة رقيقة وسعادة هائلة وسعادة
حياتية --- الخ ك أنت تعتبر كل جزء منهم
dimension وبتقسيمه بسؤال مفصل أسأل
مفصلة بتقسيمه ك بعد بيتي بسؤال تفصيلي
واجهاتيات عديدة لكل جزء من أجزاء السعادة مفصلة.