Absolute Risk Reduction

Reported effects of a medical treatment are most directly expressed as **absolute risk reduction**.

In a controlled medical study, patients are assigned to one of two groups: control group (no treatment) or treated group. At the conclusion of the study, the <u>adverse event rate (%) for the control group (CER)</u> and <u>adverse event rate (%) for the</u> <u>treated group (TER)</u> are observed. The difference between the CER and the TER is the absolute risk reduction (ARR). Reporting these three numbers - CER, TER, ARR - most directly communicates the <u>treatment effect</u>, allowing viewers to assess their <u>risk of adverse event without treatment</u> (CER), <u>risk of adverse event with</u> <u>treatment</u> (TER), and <u>risk reduction conferred by treatment</u> (ARR).

In publications and in advertisements, effects of a medical treatment are often expressed by performing an additional mathematical calculation: *dividing the* absolute risk reduction (**ARR**) by the **CER** to generate a <u>percentage of a percentage</u>, termed the relative risk reduction (**RRR**). The **RRR** is a **larger number** which communicates neither the **baseline risk** nor the **absolute risk reduction** conferred by treatment.

The value of a medical treatment is clearly communicated by reporting:

- 1) adverse event rate (%) in the control group (CER),
- 2) adverse event rate (%) in the treatment group (TER), and
- 3) absolute risk reduction (%) conferred by treatment (ARR)

With these three numbers, the reader can directly assess how individuals similar to those in the study will fare, in terms of <u>*risk* of an adverse event without treatment</u> and the <u>benefit to be expected by treatment</u>.