

	<b>CROSS SECTIONAL</b>	<b>CASE CONTROL</b>	<b>Cohort Studies</b>
<b>Advantages</b>	<ul style="list-style-type: none"> <li>-primarily used to determine <b>prevalence</b></li> <li>-useful at identifying associations and generating hypotheses about the <b>cause of disease</b></li> <li>-useful to study chronic conditions</li> <li>- Relatively easy, quick and inexpensive.</li> <li>- Minimal ethical problems</li> <li>- Can be used to estimate the risk by calculating the <b>odds ratio</b>.</li> <li>-allows study of several diseases / exposures</li> <li>-useful for estimation of the population burden, health planning and priority setting of health problems</li> <li>-relatively common conditions; allows for stratification; different from surveillance / registers</li> </ul>	<ul style="list-style-type: none"> <li>-Study of <b>rare diseases (NCDs)</b></li> <li>-Study of diseases with long latency period</li> <li>-Evaluate <b>all possible factors</b> associated with the disease</li> <li>-Quantification of the risk associated with exposure (s)</li> <li>-Save cost and time (least expensive; least time-consuming)</li> <li>-No value in the study of rare exposure</li> <li>-<b>Not for study of several diseases associated with a single exposure</b></li> <li>-effect measure: <b>odds ratio(past)</b></li> </ul>	<p><b>Retrospective</b></p> <ul style="list-style-type: none"> <li>-In retrospective there is <b>less time consumed</b> for the study (the data is already there)</li> <li>-Retrospective is <b>cheaper</b> than prospective</li> <li>-Retrospective is suitable for diseases that take a long time to develop (e.g. cancers, Parkinson's,.)</li> </ul> <hr/> <p><b>Cohort studies</b></p> <ul style="list-style-type: none"> <li>-Useful in <b>rare exposures</b></li> <li>-Can <b>study multiple outcomes of a single exposure / risk factor</b></li> <li>-<b>Certain about the temporality of exposure and disease</b> (disease occurs after exposure)</li> <li>-We can calculate incidence proportion and rate</li> <li>-Can quantify Risk Ratio and Risk Difference</li> <li>-<b>Provides better evidence</b> than case-control study, and cross-sectional study</li> <li>-Can establish a <b>natural history of disease</b> when not known</li> <li>-suitable for incidence estimation</li> <li>- effect measure: <b>relative risk (follow up)</b></li> </ul>
<b>Disadvantages</b>	<ul style="list-style-type: none"> <li>- do not differentiate between cause and effect or the sequence of events</li> <li>- Rare conditions cannot efficiently be studied</li> <li>- It deals with <b>survivors</b></li> <li>- Not useful for establishing causal relationships</li> </ul>	<ul style="list-style-type: none"> <li>-Cases don't represent cases in the general population</li> <li>-Selection bias</li> <li>- Recall bias</li> <li>- Inability to define the temporal sequence between the disease and the exposure</li> </ul>	<p><b>Retrospective</b></p> <ul style="list-style-type: none"> <li>-Sometimes difficult to determine the accuracy of the historical data in retrospective studies</li> </ul>

	<p>- Confounding is difficult to control</p>	<p>-not suitable for calculation of frequency measures.</p>	<p><b>Cohort studies</b></p> <ul style="list-style-type: none"> <li>-There is potential for loss to follow up, especially in diseases that take a long time to develop</li> <li>-Measurement errors, multiple interviews, tests</li> <li>-Not suitable for evaluation of rare diseases</li> <li>-Takes a long time (if prospective)</li> <li>-More expensive than case-control and cross-sectional studies</li> <li>-Provides weaker evidence than RCTs</li> <li>-Non response</li> </ul>
	<p><b>*observational (analytic) research could be descriptive</b></p> <p><b>Uses:</b></p> <ol style="list-style-type: none"> <li>1. Describe the state of health</li> <li>2. Describe the distribution of risk factors &amp; other attributes.</li> <li>3. Factors associated with diseases</li> <li>4. Factors associated with use of health services</li> <li>5. Determine the association of various factors and diseases.</li> <li>6. Make comparisons within and among various communities to determine if services are allocated according to needs</li> </ol> <p><b>Examples:</b></p> <ol style="list-style-type: none"> <li>1. National Surveys; (NHANES) in USA</li> <li>2. Patient satisfaction in primary care clinics</li> <li>3. CHD in relation to physical exercises.</li> </ol>	<p>-overall aim is to identify and quantify the risk factor(s) associated with the occurrence of a health problem</p> <p><b>-Analytic research</b></p>	<p><b>Examples of Famous Cohort Studies</b></p> <ul style="list-style-type: none"> <li>-The Framingham Study (1948)</li> <li>-British Physicians Cohort UK</li> <li>-Nurses Health Study USA (1976)</li> <li>-Women Health Initiative (WHI)</li> <li>-Study of women across the nation (SWAN) in USA</li> </ul> <p><b>What can we measure in a Cohort study:</b></p> <ul style="list-style-type: none"> <li>-Risk (incidence proportion)</li> <li>-Rate (incidence rate)</li> <li>-Prevalence</li> <li>-Risk Ratio (relative risk)</li> <li>-Risk Difference</li> <li>-Attributable Risk Fraction</li> </ul>

	4. Obesity in relation to diabetes mellitus 5. KAP 6. A census		
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<b>Factors affecting incidence rate</b>	<b>Factors affecting Prevalence</b>
<ul style="list-style-type: none"> <li>-New risk factor</li> <li>-Changing habits</li> <li>-Changes in virulence of causative organisms</li> <li>-Changes from intervention programs</li> <li>-Selective migration of susceptible persons</li> <li>-Population pattern (aging)</li> <li>-Reporting</li> <li>-Screening</li> <li>-New diagnostic tools</li> </ul>	<ul style="list-style-type: none"> <li>-Changes in incidence</li> <li>-Changes in disease duration and chronicity</li> <li>-Intervention programs</li> <li>-Selective attrition</li> <li>-Changing classifications</li> </ul>

<b>Advantages of longitudinal study design</b>	Allow the researcher to measure pattern of change & obtain factual information, requiring collection on a regular or continuing basis
<b>Disadvantages of longitudinal study design</b>	Conditioning effect

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