

See also PDF online from Purves et al.

Method	Pros	Cons
Computational Modeling	<ul style="list-style-type: none"> • Forces researcher to be explicit about mechanism • Allows for direct, testable predictions • Simple but powerful (small changes, big effects; validity of a hypothesis) 	<ul style="list-style-type: none"> • Simplification of Nervous System • Sometimes at odds with biology (e.g., “all knowing”) • Catastrophic interference • Relatively narrow (re:, generalization) • Research sometimes in isolation
Behavior (RT/accuracy/self report/etc.)	<ul style="list-style-type: none"> • Most simple method & underlies all other methods • Flexible 	<ul style="list-style-type: none"> • Gives incomplete picture of mechanism • Only as good as your design
Single Cell Recording (not discussing in class, except Perception paper)	<ul style="list-style-type: none"> • Records at the level of individual neurons (usually) • Direct measure of neuronal activity to expt manipulation 	<ul style="list-style-type: none"> • May record extracellularly; unclear then if activity is of single neuron • Aggregate behavior might be more complicated (e.g., multiunit)
Lesions	<ul style="list-style-type: none"> • Convergence across humans & animals for particular brain region fxn • How necessary is a brain region for a particular function 	<ul style="list-style-type: none"> • Don't know if effect isolated to region or its connection to other regions • Compensatory strategy to minimize effects of lesion • Difficulty in precision of area affected; hard to generalize • In animals, training is much more difficult than in humans • In humans, not under control of exper • Ethical concerns for animal treatment
Genetic manipulations (optogenetics, epigenetics)	<ul style="list-style-type: none"> • Identify risk factors for diseases • Which cognitive fxns are heritable (knockout) • GxE interactions 	<ul style="list-style-type: none"> • Genes can have many downstream effects, so hard to isolate specific mechanism of action • Often need a lot of people to make anything of GxE, and knockouts tend to be really specific (less generalizable)
Structural imaging (MRI, CT scans)	<ul style="list-style-type: none"> • Identify brain regions impacted in disorder, how disorder & healthy individuals vary as a fxn of damage 	<ul style="list-style-type: none"> • Has little to do with a particular experimental manipulation (usually, re: temporal scale), only general abilities
DTI	<ul style="list-style-type: none"> • Discover the flow of information within the brain for white matter tracks 	<ul style="list-style-type: none"> • Same as structural imaging above
TMS (newer things like tDCS, tACS)	<ul style="list-style-type: none"> • Can either impair or improve task performance • Researchers are now looking at how stimulation can improve brain fxn • Noninvasive virtual lesion 	<ul style="list-style-type: none"> • Effects of TMS usually brief • Only works for superficial cortical regions • Affects large area, limiting anatomical resolution • Sometimes adverse effects
fMRI	<ul style="list-style-type: none"> • High spatial resolution underlying 	<ul style="list-style-type: none"> • Indirect measure of neuronal activity

	<p>the regions impacted by task manipulations</p> <ul style="list-style-type: none"> • Noninvasive 	<ul style="list-style-type: none"> • OK/poor temporal resolution • Not cheap
EEG (ERPs)	<ul style="list-style-type: none"> • High temporal resolution underlying cognitive processes • Direct measure of neuronal activity on scalp • Cheap, noninvasive 	<ul style="list-style-type: none"> • Poor spatial resolution • Needs a lot of trials to average over •
MEG	<ul style="list-style-type: none"> • Similar to EEG, but affects sulci, not gyri • Less affected by distortions in skull than EEG • Has simpler source estimation 	<ul style="list-style-type: none"> • Same problems & benefits as EEG
PET	<ul style="list-style-type: none"> • Figuring out the concentration of particular neuromodulators in the brain (e.g., dopamine) 	<ul style="list-style-type: none"> • Short half life of reagents • Radioactive materials • Expensive • Poor temporal resolution (block designs)
Pharmacological perturbations (not in textbook)	<ul style="list-style-type: none"> • Drug use on cognitive processes • Experimental control setting, effects monitored 	<ul style="list-style-type: none"> • Lack of specificity in the effects (don't know the actual mechanism)

Also not covered in textbook: eye-tracking (see: https://en.wikipedia.org/wiki/Eye_tracking for brief summary), EcoG (see: <https://en.wikipedia.org/wiki/Electrocorticography> for brief summary of what the method is)