

# A. Magnetic / MRI-Related (Non-Invasive)

1. **Functional Connectivity MRI (fcMRI)**  
Not the same as BOLD; uses correlation structure across voxels. (Probably implicit, but useful to explicitly list.)
2. **Resting-State fMRI (rs-fMRI)**  
Now its own category of technique with unique analytical methodology.
3. **Magnetic Particle Imaging (MPI)**  
A new imaging modality using superparamagnetic nanoparticles for high temporal resolution neurovascular imaging.
4. **Susceptibility-Weighted Imaging (SWI)**  
Useful for venous blood mapping and hemodynamic activity in vivo.
5. **Quantitative Susceptibility Mapping (QSM)**  
Reconstructs iron content and hemodynamic changes related to function.
6. **Functional MRS (fMRS)**  
Measures neurotransmitter fluctuations (Glu, GABA) in vivo in real time.
7. **Magnetic Resonance Elastography (MRE)**  
Measures brain tissue stiffness changes associated with neural activity and neurovascular coupling.
8. **Chemical Exchange Saturation Transfer (CEST) fMRI**  
Detects metabolites and neurotransmitters via exchangeable protons, providing molecular functional imaging.
9. **Microvascular Volumetric Pulsatility Mapping**  
Combines VASO and ASL at ultra-high field to map pulsatile flow in cerebral microvessels linked to activity.
10. **Neurite Orientation Dispersion and Density Imaging (NODDI)**  
Multicompartment diffusion MRI model to estimate neuronal microstructure and density in vivo.
11. **Neuromelanin-Sensitive MRI**  
Detects neuromelanin in substantia nigra and locus caeruleus as a proxy for catecholaminergic neuron function.
12. **Free-Water Diffusion Imaging**  
Removes extracellular free water contributions from diffusion signals to assess
13. **Optically Pumped Magnetometer MEG (OPM-MEG)**  
Room-temperature, scalp-proximate optically pumped magnetometer arrays for high-sensitivity, flexible MEG recordings in naturalistic and mobile settings.
14. **Hyperpolarized 13C Metabolic MRI**  
Real-time metabolic flux imaging using hyperpolarized 13C tracers to measure rapid changes in brain energy metabolism linked to neural activity.
15. **Magnetic Resonance Fingerprinting (MRF)**  
Rapid, quantitative multiparametric tissue mapping for assessing dynamic physiological changes during brain activity.
16. **VASO (Cerebral Blood Volume fMRI)**  
Non-BOLD fMRI technique sensitive to cerebral blood volume changes for layer-specific functional imaging.

### **17. Q-ball Imaging**

High angular resolution diffusion imaging (HARDI) technique for resolving complex fiber crossings in white matter.

### **18. Continuous Arterial Spin Labeling (CASL)**

Non-invasive perfusion MRI method using continuous RF inversion of arterial water for quantitative CBF mapping.

### **19. Pulsed Arterial Spin Labeling (PASL)**

ASL variant using short RF pulses to label arterial blood, offering different trade-offs for perfusion imaging.

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## **B. PET / Radiotracer Advances**

### **20. Fiber-Coupled PET Detectors (minimally invasive)**

Enables localized in vivo PET from freely moving animals.

### **21. Neuroimmune PET Ligands (TSPO, CSF1R etc.)**

For studying microglial/astrocyte activation dynamically. (Not purely activity, but increasingly part of functional neuroimaging.)

### **22. Total-Body PET Imaging**

Enables whole-brain dynamic imaging with unprecedented temporal resolution and sensitivity.

### **23. Positronium Lifetime Imaging**

Emerging PET technique providing information about tissue microenvironment during neural activity.

### **24. Synaptic Vesicle Glycoprotein 2A (SV2A) PET Imaging**

Uses tracers like [<sup>11</sup>C]UCB-J to quantify synaptic density in vivo.

### **25. Mitochondrial Complex I PET Imaging**

Employs [<sup>12</sup>F]BCPP-EF to assess mitochondrial function and energy metabolism related to neural activity.

### **26. CSF1R PET for Neuroinflammation**

Tracers like [<sup>11</sup>C]CPPC to image microglial activation during brain processes.

### **27. Positron Emission Metabolic Tracing with Short-Lived Isotopes (rapid kinetic PET)**

High-temporal-resolution PET protocols using short-lived radiotracers and fast kinetic modeling to track sub-minute metabolic changes associated with neural events.

### **28. μ-opioid Receptor PET (<sup>[11</sup>C]carfentanil)**

In vivo imaging of endogenous opioid release during various behavioral states.

### **29. Dopamine D<sub>2</sub>/D<sub>3</sub> Receptor Occupancy PET**

Measures dynamic changes in dopamine transmission using radioligands like [<sup>11</sup>C]raclopride.

### **30. Cerebral Metabolic Rate of Oxygen (CMRO<sub>2</sub>) PET**

Quantitative imaging of oxygen metabolism coupled to neural activity using <sup>15</sup>O-labeled tracers.

### **31. Astrocyte-Specific PET Tracers (<sup>11</sup>C-deuterium-L-deprenyl)**

Targets monoamine oxidase B in astrocytes for glial activity mapping.

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# C. Electrophysiology & Implant Technologies

## 32. Neuropixels Probes (latest gen)

Not just "multielectrode"; Neuropixels constitute a distinct class due to ultra-high channel count (1,000+), used *in vivo* widely.

## 33. Silicon Probe Laminar Recordings

For cortical column laminar activity (V1, CA1).

## 34. High-Density “ECoG Grids” (Neuropixels–ECoG hybrids)

New grids with thousands of micro-electrodes recordings across cortex.

## 35. EMG-Assisted Brain–Body Coupled Recording

Used in behaving animals to integrate motor output with neural activity.

## 36. Flexible Bioelectronic Neural Interfaces

Conformable electrode arrays that minimize tissue damage and enable stable long-term recordings.

## 37. Transparent Graphene Microelectrode Arrays

Allow simultaneous electrical recording and optical imaging/optogenetics.

## 38. Ultrasonic Neural Dust Motes

Wireless, millimeter-scale implants for chronic neural recording.

## 39. Optetrode Recordings

Hybrid probes combining electrical electrodes with optical fibers for simultaneous recording and stimulation.

## 40. CMOS-Integrated Neural Probes

On-chip amplified high-density probes for low-noise *in vivo* recordings.

## 41. Sharp Electrode Intracellular Recordings

Historical method using fine glass micropipettes for high-impedance intracellular potential measurements.

## 42. Wireless High-Density Neural Probes (wireless Neuropixels variants)

Fully wireless, implantable high-channel-count probe systems enabling untethered large-scale neural recordings in freely moving animals.

## 43. Autonomous Robotic In Vivo Patch-Clamp (AutoPatch / Robopatcher)

Automated robotics-enabled intracellular patch-clamp in awake or anesthetized

## 44. Graphene Field-Effect Transistor Neurochemical Sensors (gFETs)

Implantable gFET biosensors for direct, high-sensitivity, real-time detection of neurotransmitters and neuromodulators *in vivo* with electrical readout.

## 45. Tetrode Recordings

Historical method using four-wire bundles for extracellular recording with improved single-unit isolation.

## 46. Juxtacellular Recording and Labeling

Technique for extracellular recording followed by intracellular labeling of recorded neurons.

#### **47. Carbon Fiber Microelectrodes**

Miniaturized electrodes for electrochemical detection of neurotransmitters with fast-scan cyclic voltammetry.

#### **48. Floating Microelectrode Arrays**

Untethered microelectrodes that move with brain tissue for stable chronic recordings.

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## **D. Optical / Imaging-Based Techniques**

#### **49. Light-Sheet Fluorescence Microscopy (LSFM) In Vivo**

Now used in small transparent animals (zebrafish, larval models). Technically in vivo and provides whole-brain fast activity imaging.

#### **50. Swept Confocally-Aligned Planar Excitation (SCAPE) Microscopy**

High-speed volumetric neural imaging in freely moving animals.

#### **51. Structured-Light 3D Imaging of Cortical Hemodynamics**

Used to map neural activity via intrinsic signals & hemodynamic changes.

#### **52. Mesoscopic Calcium Imaging (NOT miniscope)**

Widefield mesoscopic  $\text{Ca}^{2+}$  imaging in vivo (distinct from fiber photometry).

#### **53. Adaptive Optics for In Vivo Neural Imaging**

Corrects deep-tissue optical distortion; rapidly emerging technique.

#### **54. Oblique Plane Microscopy (OPM)**

High-resolution, high-speed volumetric imaging for large-scale neural activity monitoring.

#### **55. Multifocal Two-Photon Microscopy**

Simultaneous imaging of multiple planes for 3D functional imaging.

#### **56. Line-Scanning Temporal Focusing Microscopy**

High-speed volumetric imaging with reduced out-of-focus excitation.

#### **57. Light Field Microscopy**

Captures volumetric data in a single exposure for fast 3D neural activity imaging.

#### **58. Kilohertz Two-Photon Fluorescence Microscopy**

Ultrafast scanning rates for capturing rapid neuronal dynamics in vivo.

#### **59. NIRE Cranial Window Method**

Uses nanosheet-resin windows for large-scale, chronic high-resolution imaging in

#### **60. Volumetric Fluorescence Lifetime Imaging Microscopy (FLIM) for In Vivo Metabolic Readouts**

High-speed volumetric FLIM approaches enabling in vivo mapping of metabolic state (NADH, FAD) and oxygen-consumption dynamics across populations of cells.

#### **61. TIRF Microscopy In Vivo**

Total internal reflection fluorescence microscopy adapted for superficial cortical imaging in live animals.

#### **62. Random Access Microscopy**

Acousto-optic deflector-based rapid laser positioning for imaging distributed neurons.

#### **63. Reflectance Imaging**

Measures intrinsic optical signals from cortical surface without exogenous labels.

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**64. Second Harmonic Generation (SHG) Microscopy**

Label-free imaging of membrane potential and structural proteins in live tissue.

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## E. Optical Indicators / Novel Sensors

**65. Genetically Encoded Dopamine Indicators (GRAB-DA, dLight1)**

Used widely in vivo with fiber photometry and two-photon microscopy.

**66. Genetically Encoded Glutamate Indicators (IGluSnFR)**

High temporal resolution; used to study excitatory transmission.

**67. Genetically Encoded Acetylcholine Indicators (GAcH)**

Maps cholinergic dynamics in vivo.

**68. Genetically Encoded cAMP/PKA/Second-Messenger Sensors**

(e.g., Pink Flamingo, G-Flamp) reflect intracellular signaling during neural activity.

**69. Genetically Encoded Serotonin Sensors (GRAB5-HT)**

Monitor serotonergic transmission in behaving animals.

**70. Genetically Encoded Norepinephrine Sensors (GRAB-NE)**

Track noradrenergic activity during behavior and cognitive tasks.

**71. Fluorescent False Neurotransmitters (FFNs)**

Visualize neurotransmitter release and recycling in vivo.

**72. Genetically Encoded Chloride Indicators (Cl-Sensor)**

Monitor chloride dynamics relevant for inhibitory transmission.

**73. Genetically Encoded ATP Indicators (IATPSnFR)**

Track cellular energy status and metabolic activity in neurons.

**74. Genetically Encoded Lactate Sensors**

Monitor lactate shuttling and metabolic coupling between neurons and glia.

**75. Genetically Encoded Redox Indicators (roGFP)**

Detect oxidative stress and redox-state changes during neural activity.

**76. Genetically Encoded Nitric Oxide Sensors (geNOps and related probes)**

Fluorescent genetically encoded reporters for nitric oxide dynamics to monitor NO signaling in vivo during neural activity.

**77. Genetically Encoded Potassium Indicators (GEPIs)**

Fluorescent sensors for monitoring potassium dynamics in extracellular space.

**78. pHluorins (pH-sensitive GFPs)**

pH-sensitive fluorescent proteins for tracking synaptic vesicle recycling.

**79. SypHy**

Synaptophysin-pHluorin fusion protein for imaging synaptic vesicle exocytosis.

**80. GCamp Variants (GCamp6f, GCamp7f, GCamp8)**

Successive generations of genetically encoded calcium indicators with improved kinetics and sensitivity.

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## F. Photoacoustic & Hybrid Techniques

81. **Voltage-Sensitive Photoacoustic Imaging**  
A new area combining voltage dyes with photoacoustics.
  82. **Nanodiamond Magnetometry in Vivo**  
Nitrogen-vacancy (NV) diamond sensors used for in vivo magnetic-field measurements of neuronal activity.
  83. **Multispectral Optoacoustic Tomography (MSOT)**  
Provides spectral unmixing of multiple chromophores for functional brain imaging.
  84. **Photoacoustic Computed Tomography (PACT)**  
Deep-tissue functional imaging with optical contrast and ultrasound resolution.
  85. **Functional Photoacoustic Microscopy (fPAM)**  
High-resolution imaging of hemodynamic responses to neural activity.
  86. **Photoacoustic Lifetime Imaging Microscopy (PALM)**  
Measures oxygen consumption and metabolic rates in brain tissue.
  87. **Microbubble-Enhanced Functional Ultrasound (Contrast-Enhanced fUS)**  
Use of intravascular microbubble contrast agents to boost sensitivity and spatial resolution of functional ultrasound hemodynamic measurements linked to neural activity.
  88. **Photoacoustic Neurotransmitter Sensing**  
Molecular imaging of neurotransmitter release using photoacoustic contrast agents.
  89. **Granger Causality Photoacoustic Imaging**  
Combines photoacoustic imaging with Granger causality analysis for functional connectivity mapping.
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## G. Neural Activity Through Blood Flow & Oxygenation

90. **Thermal Infrared Functional Imaging (ITIRI)**  
Detects activity-induced microvascular heating.
91. **Speckle-Modulated Optical Coherence Tomography (OCT)**  
OCT-based neuronal activity mapping (beyond standard laser speckle imaging).
92. **Doppler Optical Coherence Tomography (D-OCT)**  
For real-time blood-flow-linked neural-activity measurements.
93. **Visible Light Optical Coherence Tomography (vis-OCT)**  
Provides oximetry and flow measurements with higher resolution than NIR-OCT.
94. **Hyperspectral Imaging of Intrinsic Signals**  
Spectral unmixing of hemoglobin, cytochrome oxidase, and other chromophores.
95. **Time-Domain Near-Infrared Spectroscopy (TD-NIRS)**  
Offers depth-resolved hemodynamic monitoring with improved accuracy.
96. **Functional Diffuse Optical Tomography (fDOT)**  
3D reconstruction of brain activation through light scattering and absorption.
97. **In Vivo Electron Paramagnetic Resonance (EPR) Oximetry**  
Direct in vivo EPR/Electron Spin Resonance oximetry for quantitative tissue pO<sub>2</sub> mapping and dynamic oxygen-consumption measurements coupled to neural activity.

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**98. Laser Doppler Flowmetry**

Continuous measurement of cerebral blood flow using laser Doppler shifts.

**99. Oxygen-15 Water PET ( $[^{15}\text{O}]\text{H}_2\text{O}$  PET)**

Gold standard for quantitative cerebral blood flow measurement using positron emission tomography.

**100. Hydrogen Clearance CBF Measurement**

Historical method using hydrogen electrodes to measure local cerebral blood flow.

**101. Autoradiographic CBF Measurement ( $[^{14}\text{C}]$ iodoantipyrine)**

Ex vivo quantitative mapping of cerebral blood flow using radiotracers.

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## H. Behavioral + Neural Integrated Techniques

**102. Neuromorphic Cameras Linked With Neural Imaging**

High-speed event-driven sensors capturing animal behavior synchronized with brain activity (not a brain technique per se, but part of modern *in vivo* pipelines).

**103. Deep Label-Free Microscopy (DLFM)**

Uses deep learning to extract neural activity from label-free imaging of scattering changes.

**104. Acoustic Recording of Neural-Related Behavior**

Ultrasonic vocalization recording synchronized with neural activity measurements.

**105. Eye-Tracking Integrated Neural Recording**

Combines pupil tracking with brain activity to study visual attention and processing.

**106. Pose Estimation with Neural-Activity Synchronization**

Uses AI-based body tracking (e.g., DeepLabCut) linked to real-time neural signals.

**107. Real-Time Closed-Loop Neural Decoding & Stimulation Systems**

Integrated pipelines that decode ongoing neural activity in real time and deliver contingent stimulation (optogenetic, electrical, ultrasonic) to probe causality during behavior.

**108. Whisker Tracking with Neural Recording**

High-speed videography of whisker movements synchronized with neural data.

**109. Operant Conditioning Chambers with Neural Interfaces**

Behavioral boxes integrated with neural recording/stimulation for learning studies.

**110. Sleep-Wake Monitoring with EEG/EMG**

Polysomnography combined with neural activity measurements across sleep stages.

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## I. Interference / Modulation Techniques

**111. Focused Ultrasound Blood–Brain Barrier Opening (FUS-BBBO)**

Used *in vivo* to modulate circuits or allow entry of neuromodulators.

112. **Temporal Interference Stimulation (TI Stimulation)**  
Non-invasive deep-brain electromagnetic stimulation using intersecting high-frequency currents.
  113. **Transcranial Random Noise Stimulation (RNS)**  
Another electrical non-invasive modulation technique widely used.
  114. **Photothermal Neuromodulation (non-genetic)**  
Uses nanoparticles or infrared light to activate neurons.
  115. **Infrared Neural Stimulation (INS)**  
Pulsed IR light drives neural activity—distinct from optogenetics.
  116. **Scanning Ultrasound Neuromodulation (SUN)**  
Focused ultrasound for precise spatiotemporal neural control.
  117. **Optoacoustic Neuromodulation**  
Uses laser-generated ultrasound for deep-brain stimulation.
  118. **Ion-CeMST (Ion Current–Controlled Microfluidic Stimulation)**  
Microfluidic-based chemical stimulation with spatiotemporal precision.
  119. **Transcranial Photobiomodulation (tPBM)**  
Low-level near-infrared light to modulate brain metabolism and activity.
  120. **Vagus Nerve Stimulation (VNS)**  
Electrical stimulation of vagus nerve to indirectly influence brain circuits.
  121. **Deep Transcranial Magnetic Stimulation (dTMS)**  
Uses H-coil for deeper penetration than standard TMS.
  122. **Closed-Loop Responsive Neurostimulation (RNS) for Research**  
Implantable or external systems that detect electrophysiological biomarkers and
  123. **Chemical-Genetic Actuation (PSAM/PSEM)**  
Pharmacologically selective actuator modules for remote neural control.
  124. **Cortical Cooling**  
Focal cooling for reversible neural inactivation to study functional localization.
  125. **Lidocaine Inactivation**  
Local pharmacological blockade of neural activity for connectivity mapping.
  126. **Muscimol Inactivation**  
GABA<sub>A</sub> receptor agonist for reversible cortical silencing in behavioral studies.
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## J. Emerging & Frontier Approaches

127. **Bioluminescent Voltage Imaging (e.g., LOTUS-V)**  
No excitation light; extremely low phototoxicity for *in vivo* activity imaging.
128. **Upconversion Nanoparticle-Based Neural Imaging**  
Allows deeper brain optical readout using NIR-to-visible conversion.
129. **Magnetothermal Neural Recording & Readout**  
Not just thermogenetic stimulation—actual readout emerging via nanomaterials.
130. **Molecular fMRI (m-fMRI)**  
Probes coupling specific cellular signaling events to MRI contrast.
131. **Functional Ultrasound Localization Microscopy (ULM-fUS)**  
Super-resolution vascular mapping during activity.

132. **"Neuromodulation via Magnetic Nanodiscs"**  
A newer mechanogenetic-like frontier method.
133. **Quantum Diamond Microscopy**  
Uses NV centers for magnetic imaging of neural activity at nanoscale.
134. **X-ray-Induced Acoustic Computed Tomography (XACT)**  
Combines X-ray absorption with ultrasound detection for functional imaging.
135. **Magnetic Resonance Spectroscopic Imaging (MRSI)**  
Spatially resolved spectroscopy for mapping neurotransmitter distributions.
136. **Neutron Stimulated Emission Computed Tomography (NSECT)**  
Emerging nuclear technique for elemental mapping of neural tissue.
137. **Holographic Optogenetic Stimulation**  
Uses holography for multi-site optical neural control and readout.
138. **Nanowire Intracellular Recordings**  
Nanoscale wires for minimally invasive intracellular potential measurements.
139. **AI-Augmented Functional Connectivity Analysis**  
Machine-learning-enhanced mapping of dynamic brain networks from imaging data.
140. **Cryogenic Electron Microscopy for In Vivo Snapshots**  
Adapted for rapid freezing and imaging of activity states (though borderline *in vivo*).
141. **In Vivo Quantum Diamond Scalp Magnetometry (NV-MEG)**  
Scalp-mounted or near-scalp arrays of diamond NV sensors for high-spatial-resolution magnetic field mapping of population neural currents, bridging nanoscale NV sensing advances with whole-head MEG-style recordings.
142. **CLARITY with In Vivo Applications**  
Tissue transformation for post-mortem structural mapping of functionally identified circuits.
143. **Expansion Microscopy In Vivo**  
Physical tissue expansion for super-resolution imaging of activated circuits.
144. **Magnetic Resonance Phased Array Microscopy**  
Ultra-high field MRI with phased array coils for microscopic resolution in live animals.
145. **X-ray Optogenetics**  
Combining X-ray stimulation with optogenetic actuators for deep brain modulation.
146. **Neutron Scattering Tomography**  
Emerging technique for mapping light element distributions in neural tissue.
147. **Diamond Quantum Sensing of Neural Magnetic Fields**  
NV center magnetometry for direct detection of action potentials and synaptic currents.