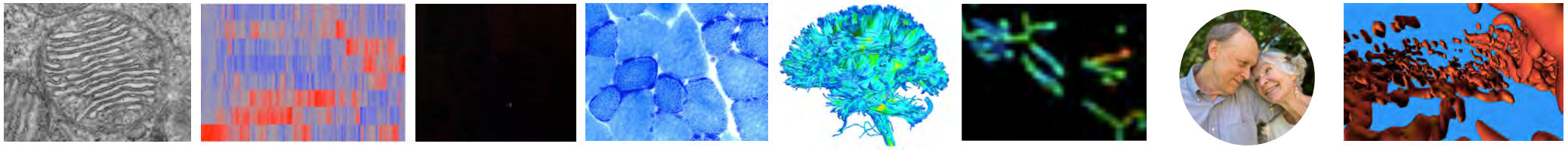
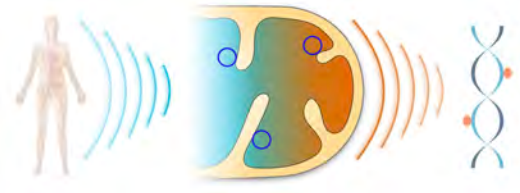


Profiling multifunctional mitochondria in the brain and beyond



Martin Picard, Ph.D.
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Department of Neurology, H. Houston Merritt Center
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New York State Psychiatric Institute (NYSPI)

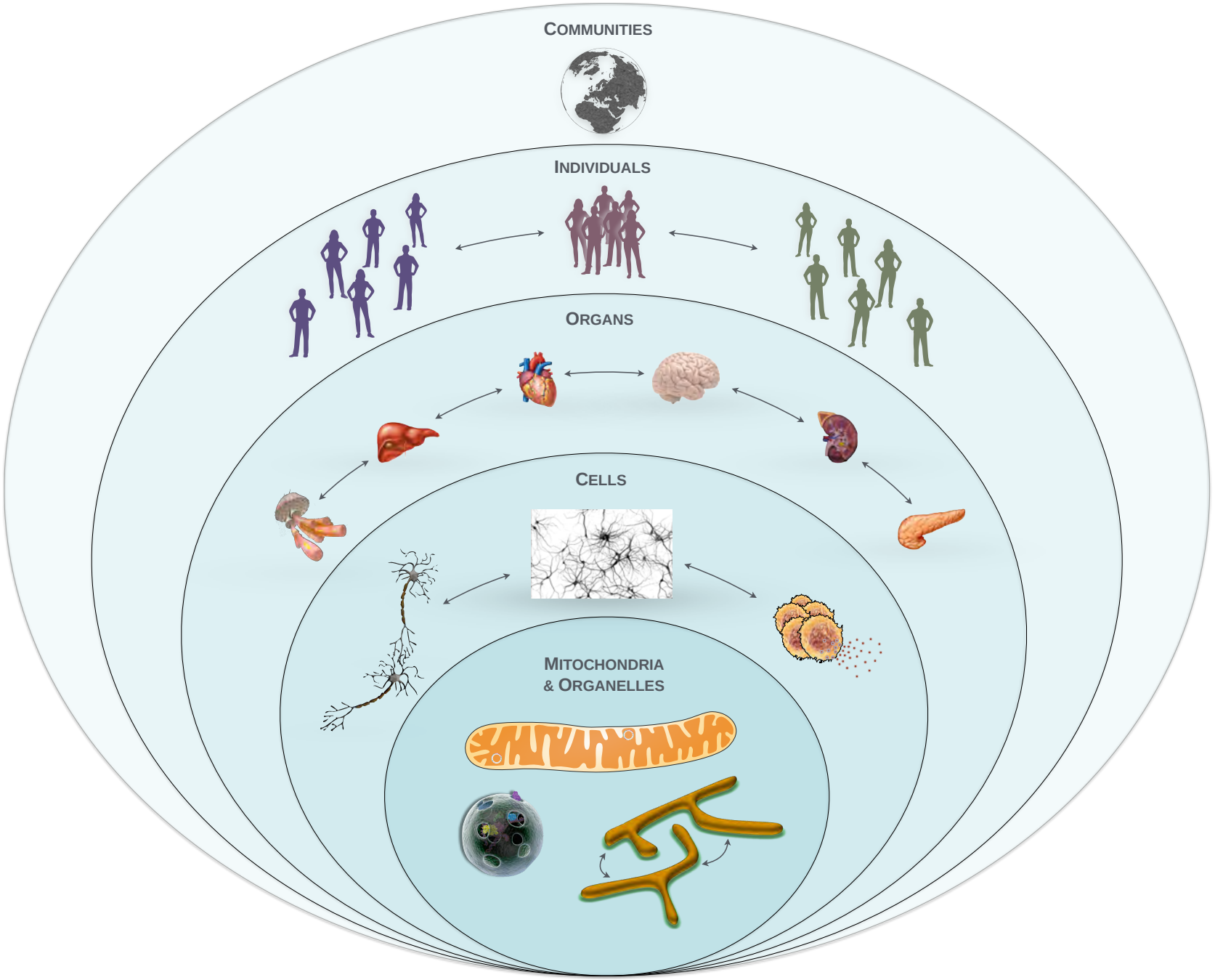
COLUMBIA
COLUMBIA UNIVERSITY
IRVING MEDICAL CENTER

NEW YORK STATE OF OPPORTUNITY. New York State Psychiatric Institute

1. Role of mitochondria in organisms

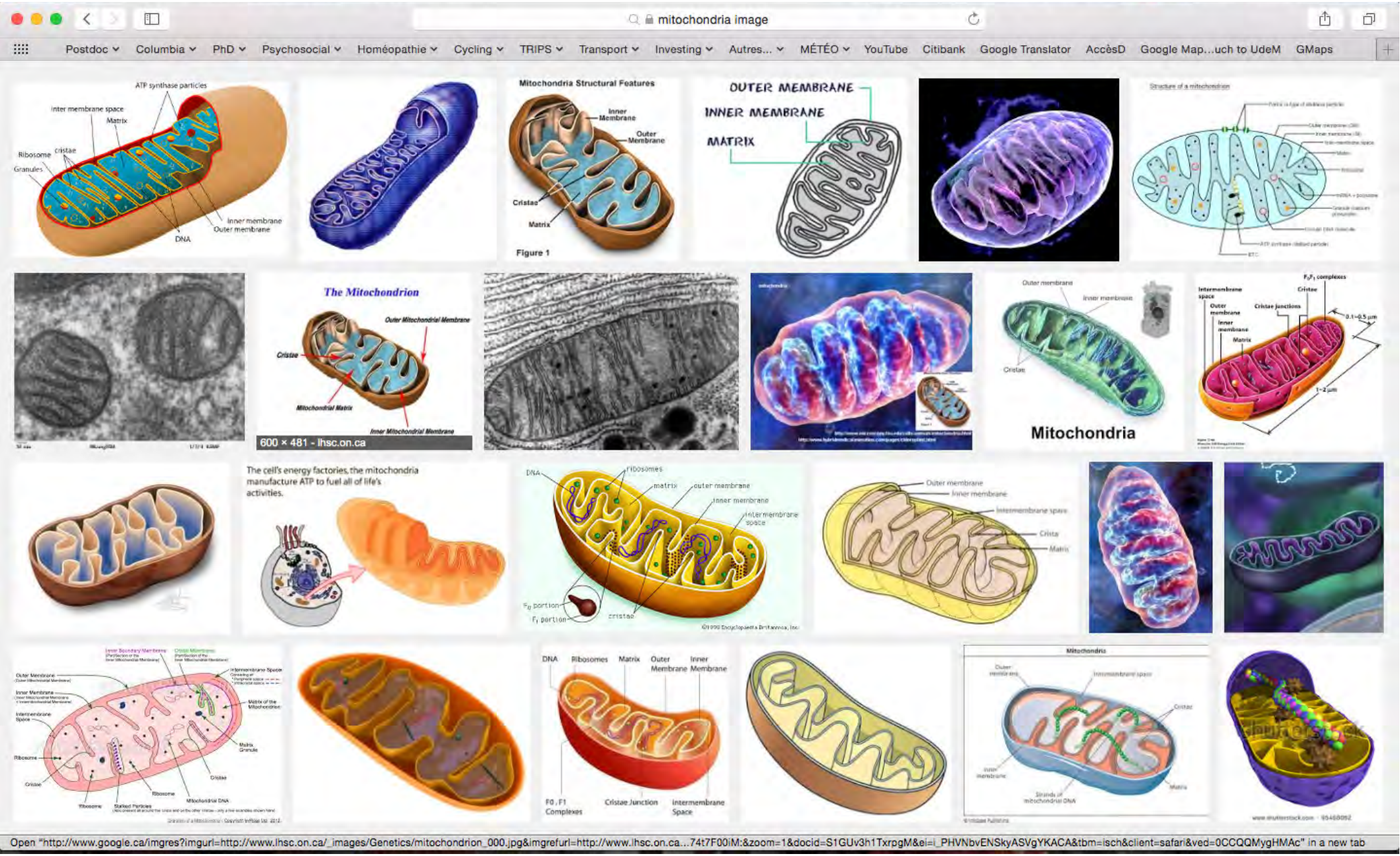
2. Celebrating diversity of mitochondria

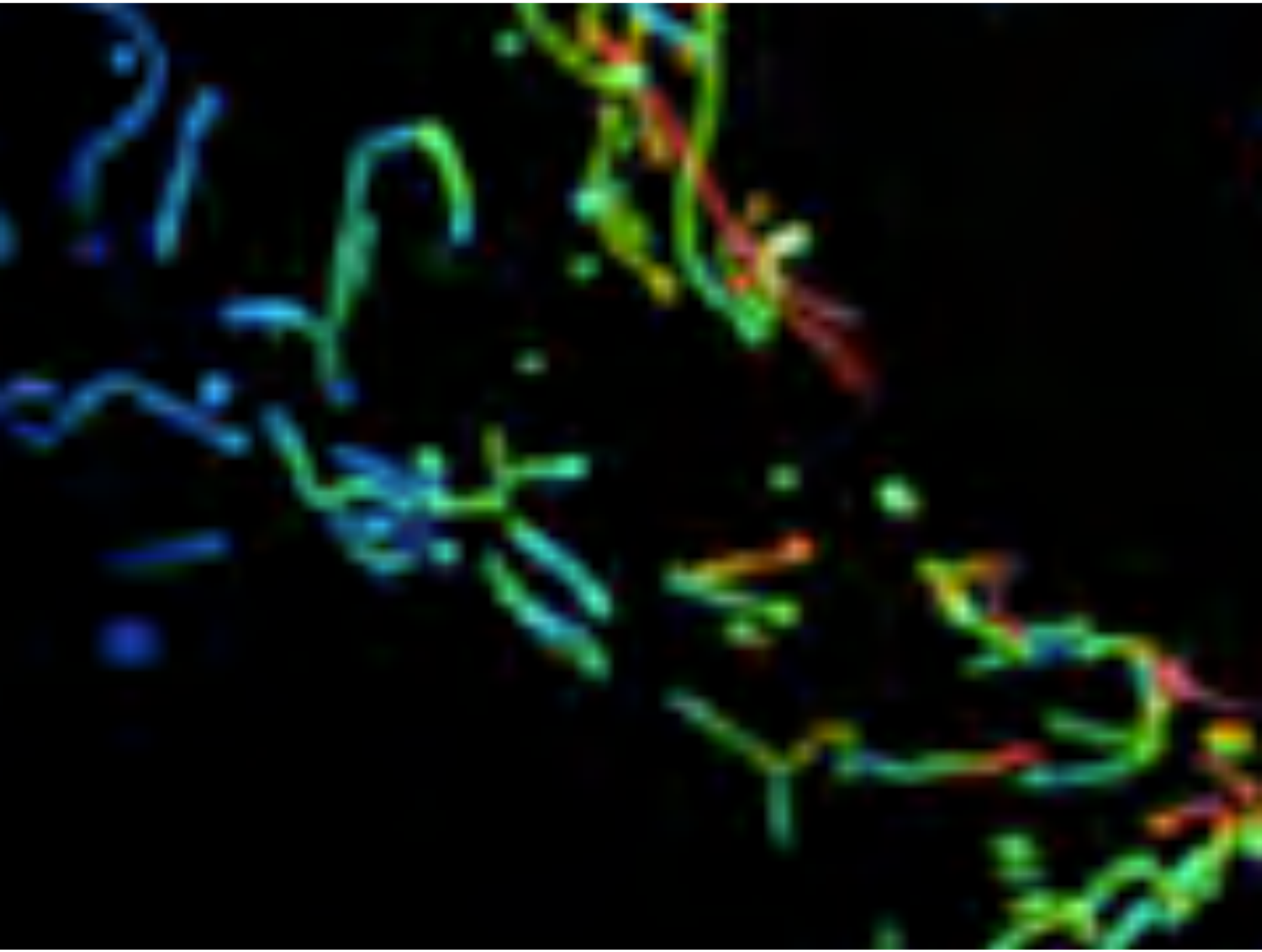
3. Human brain mitochondria



Picard and Sandi. *Neurosci Biobehav Rev* (2021)

What do mitochondria look like?





What do mitochondria do?

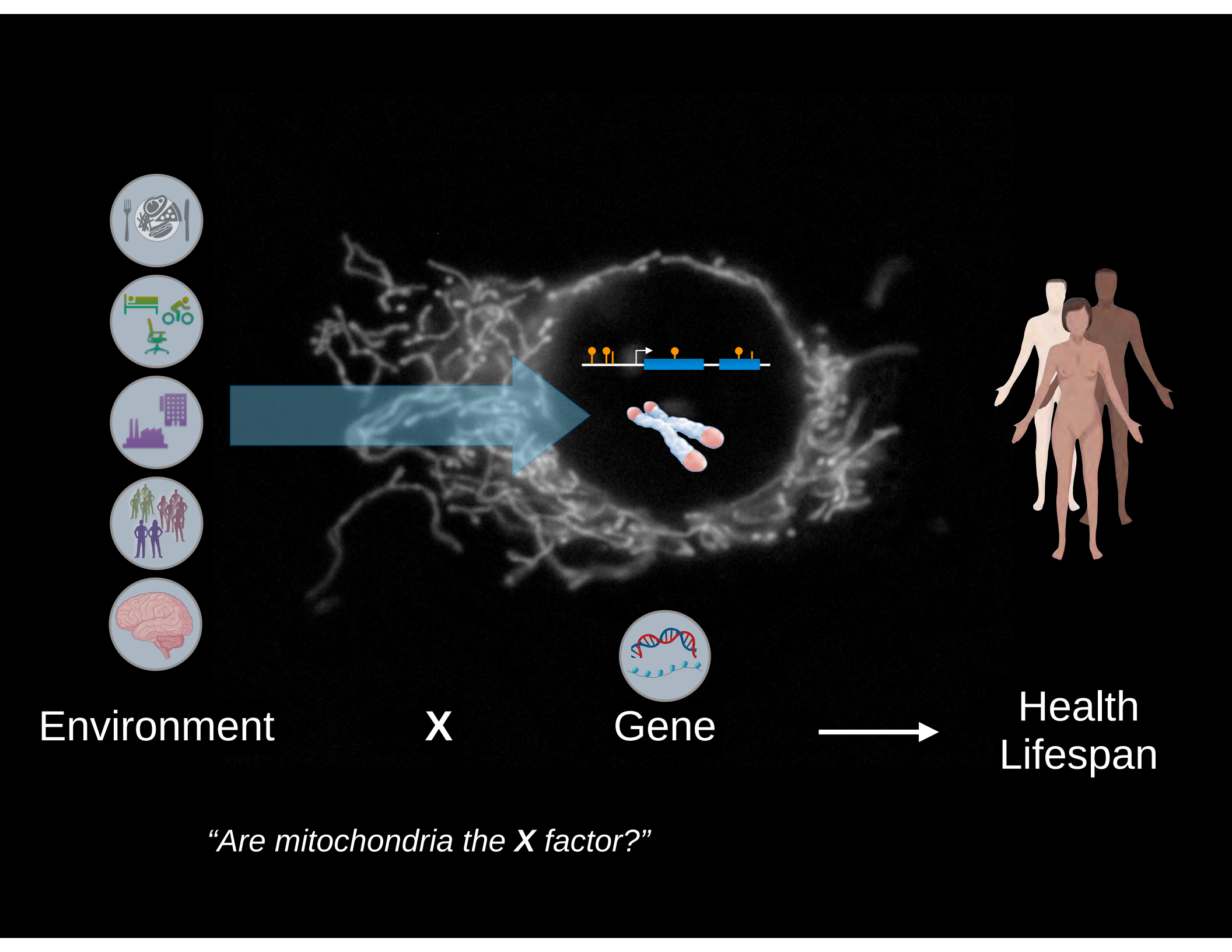
SCIENTIFIC AMERICAN®

HEALTH

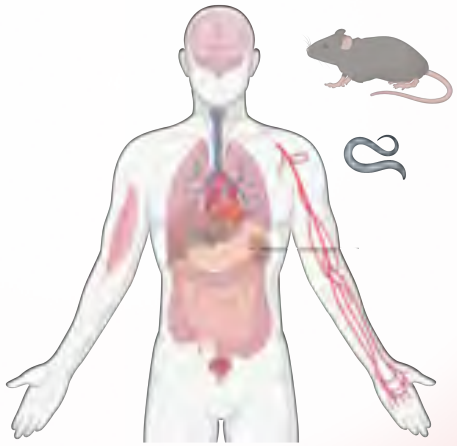
Powerhouse of the Cell

It is the mitochondrion, a small body which appears to play a central role in the oxidation of foodstuff. Its structure, as revealed by the electron microscope, mirrors its function

By Philip Siekevitz on July 1, 1957


Environment
X
Gene
Health
Lifespan
"Are mitochondria the X factor?"

Multicellular organisms

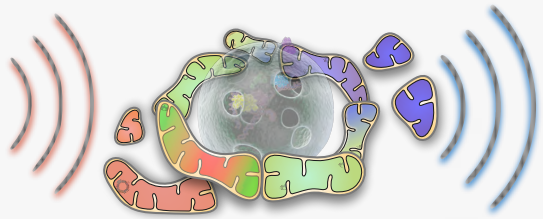


INPUTS
- Peptide hormones
- Steroid & other hormones
- Nutrients levels
- Metabolites
- Ions
- Gases (e.g., O_2, NO)
- ATP/ADP (ΔGp)
- $NAD(P)^+$/NAD(P)H ratio
- mtDNA variations
- Others

OUTPUTS (mitochondrial, DIRECT)
- Metabolites
- Lipids
- DNA and RNA
- cf-mtDNA (whole, fragments)
- ATP (ΔGp)
- Ions
- ROS
- Gases
- Heat
- Steroid hormones
- Small peptides
- Others

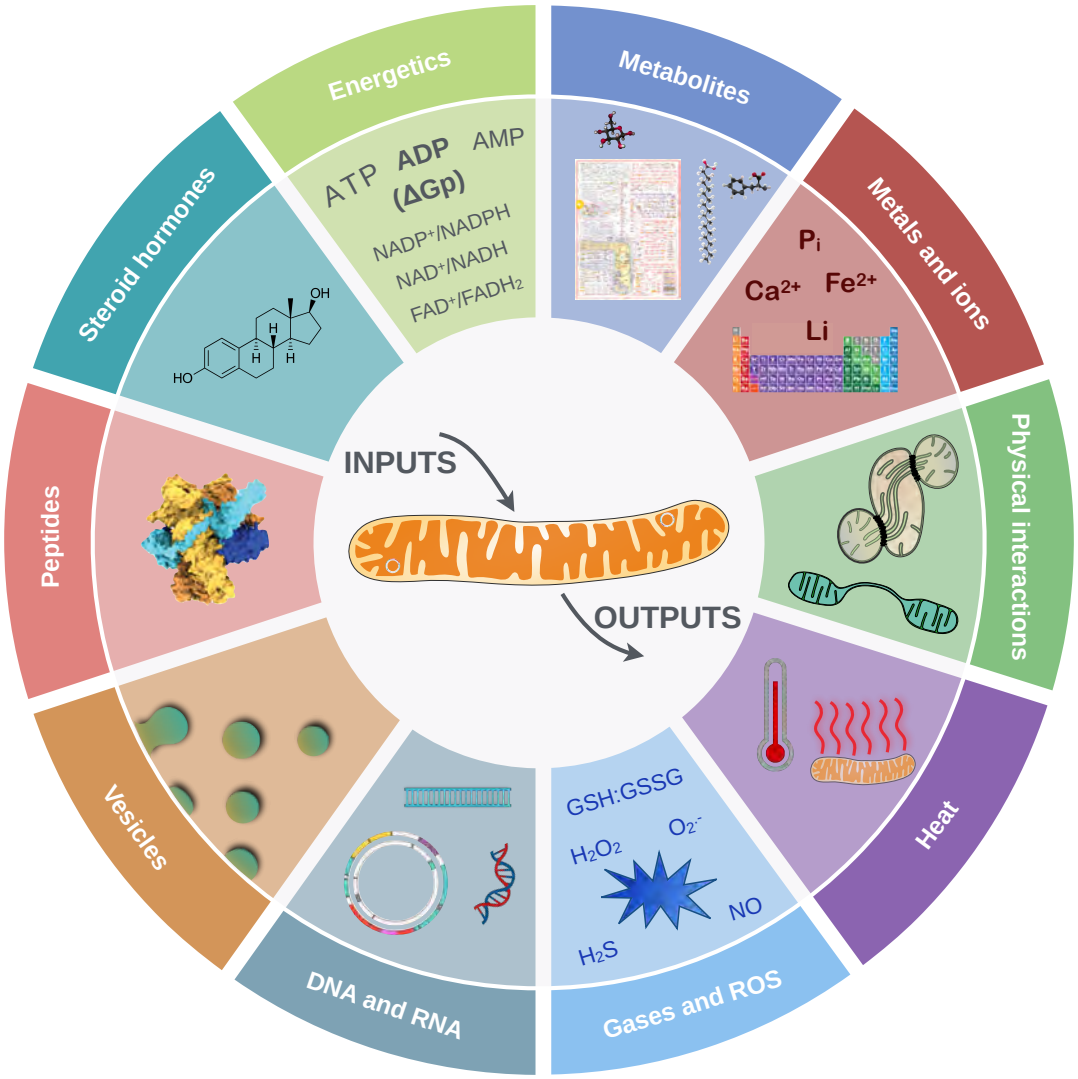
OUTPUTS (via nucleus, INDIRECT)
- Peptide hormones

INPUTS → 1 **SENSING** → 2 **INTEGRATION** → 3 **SIGNALING** → **OUTPUTS**

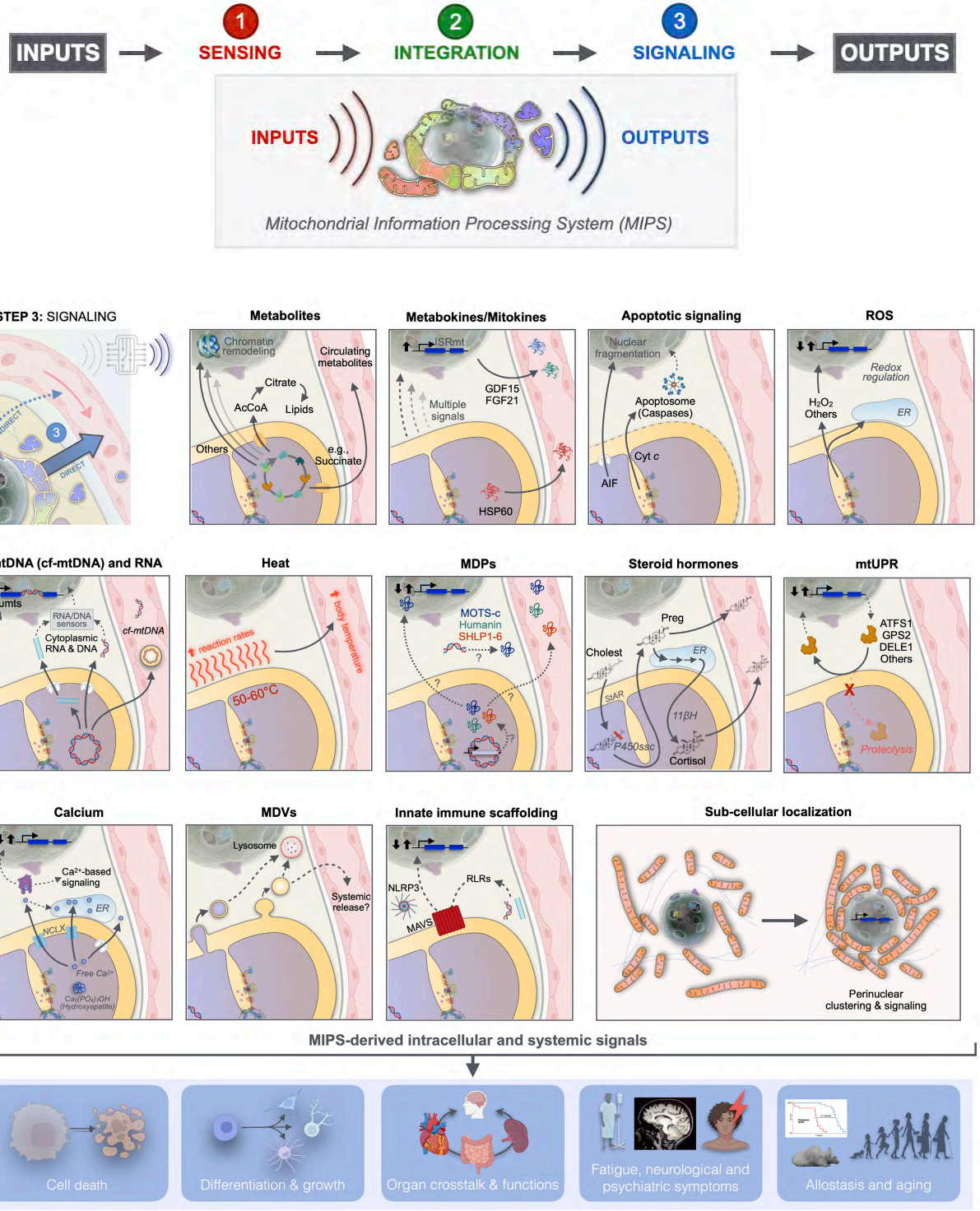


Mitochondrial Information Processing System (MIPS)

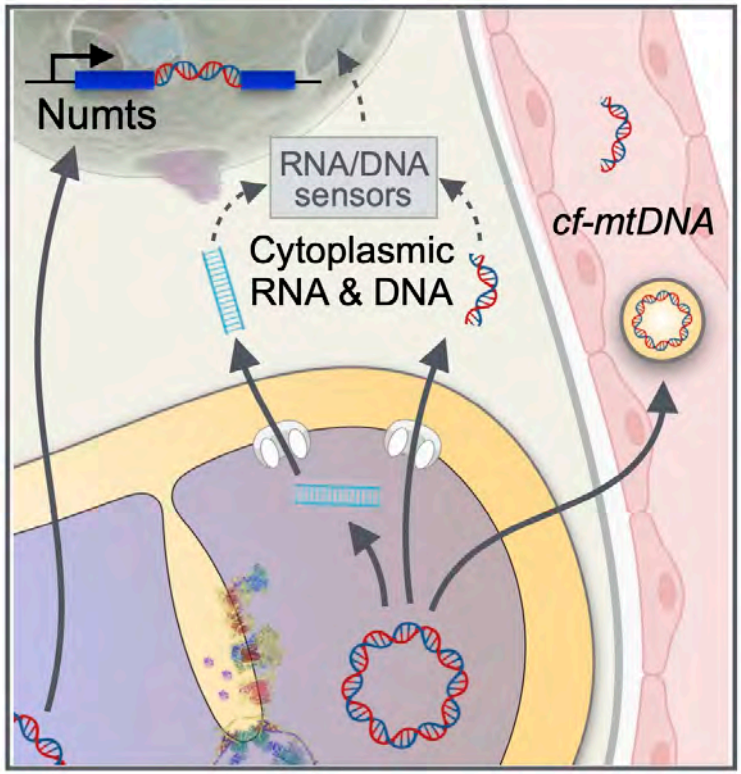
The hallmarks of mitochondrial signal transduction

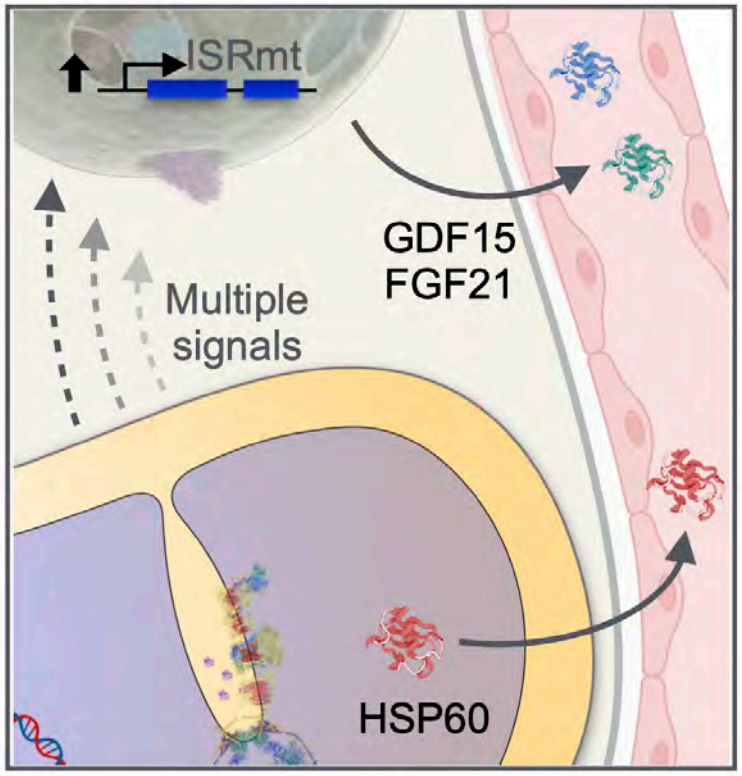


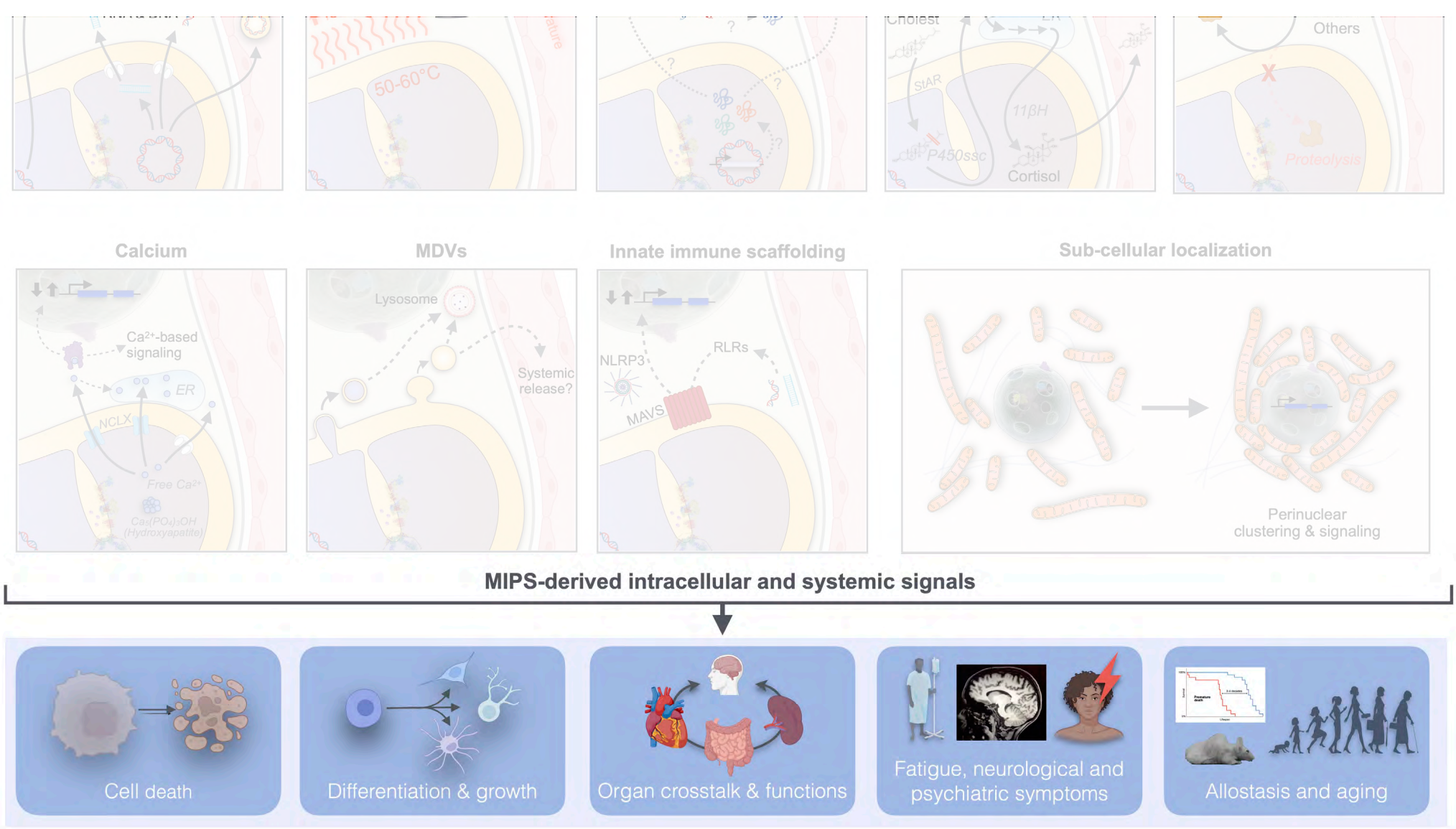
Picard and Shirihai. *Cell Metab* 2022


INPUTS
1
SENSING
2
INTEGRATION
3
SIGNALING
OUTPUTS
INPUTS
OUTPUTS
Mitochondrial Information Processing System (MIPS)
STEP 3: SIGNALING
INDIRECT
DIRECT
Metabolites
Chromatin remodeling
Circulating metabolites
Citrate
AcCoA
Lipids
Others
e.g., Succinate
Metabokines/Mitokines
ISRmt
GDF15
FGF21
Multiple signals
HSP60
Apoptotic signaling
Nuclear fragmentation
Apoptosome (Caspases)
Cyt c
AIF
ROS
Redox regulation
H_2O_2
Others
ER
mtDNA (cf-mtDNA) and RNA
Numts
RNA/DNA sensors
Cytoplasmic RNA & DNA
cf-mtDNA
Heat
reaction rates
body temperature
50-60°C
MDPs
MOTS-c
Humanin
SHLP1-6
Steroid hormones
Preg
Cholest
ER
StAR
11βH
P450ssc
Cortisol
mtUPR
ATFS1
GPS2
DELE1
Others
Proteolysis
Calcium
Ca^{2+}-based signaling
ER
NCLX
Free Ca^{2+}
$Ca_5(PO_4)_3OH$ (Hydroxyapatite)
MDVs
Lysosome
Systemic release?
Innate immune scaffolding
NLRP3
RLRs
MAVS
Sub-cellular localization
Perinuclear clustering & signaling
MIPS-derived intracellular and systemic signals
Cell death
Differentiation & growth
Organ crosstalk & functions
Fatigue, neurological and psychiatric symptoms
Allostasis and aging

mtDNA (cf-mtDNA) and RNA



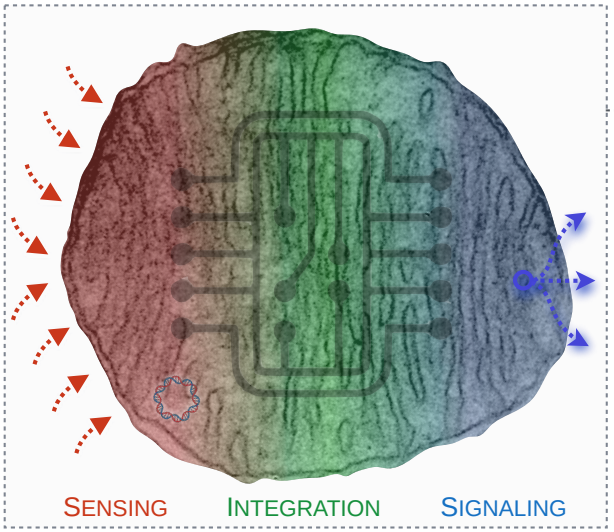

Metabokines/Mitokines
ISRmt
GDF15
FGF21
Multiple
signals
HSP60



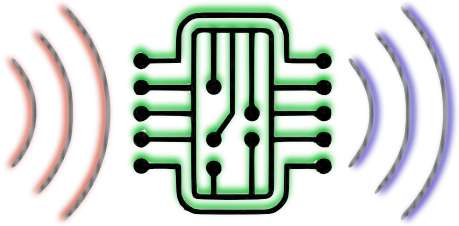
Picard and Shirihai. *Cell Metab* 2022

Mitochondrial **I**nformation **P**rocessing **S**ystem — **MIPS**

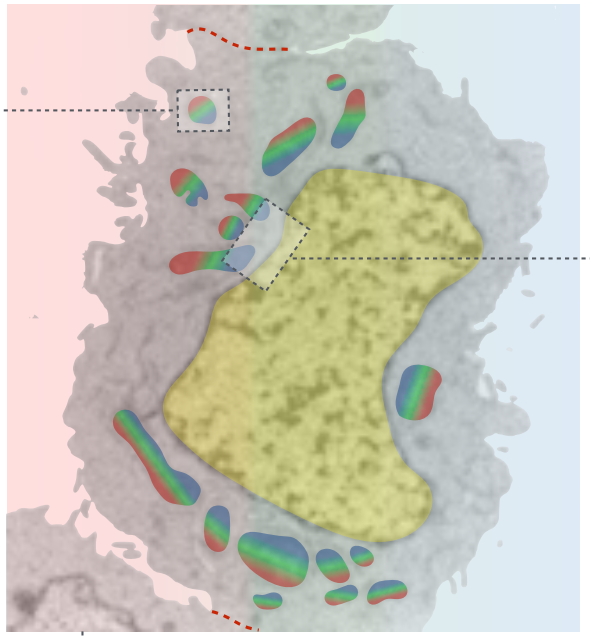
Signal transducing mitochondrion

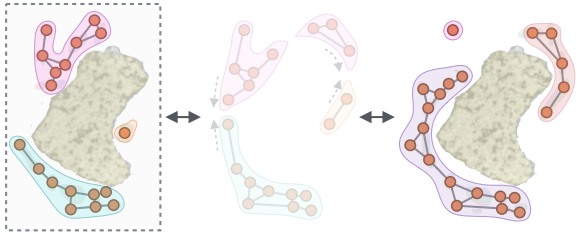


Incoming data *Outgoing data*



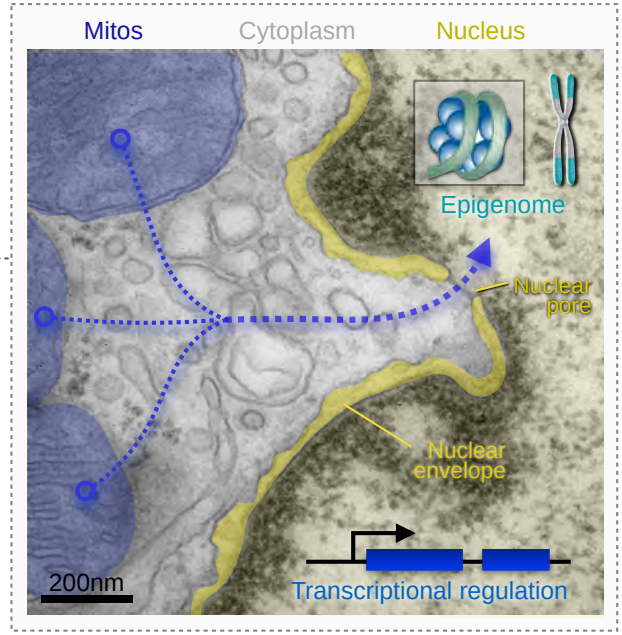
"Mitochondria are the processor of the cell"

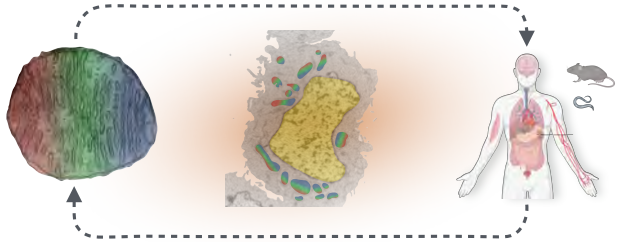


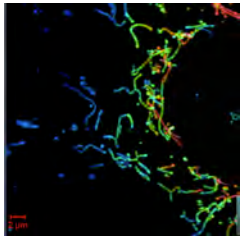


Dynamic remodeling of mito networks

Mito-nuclear unit

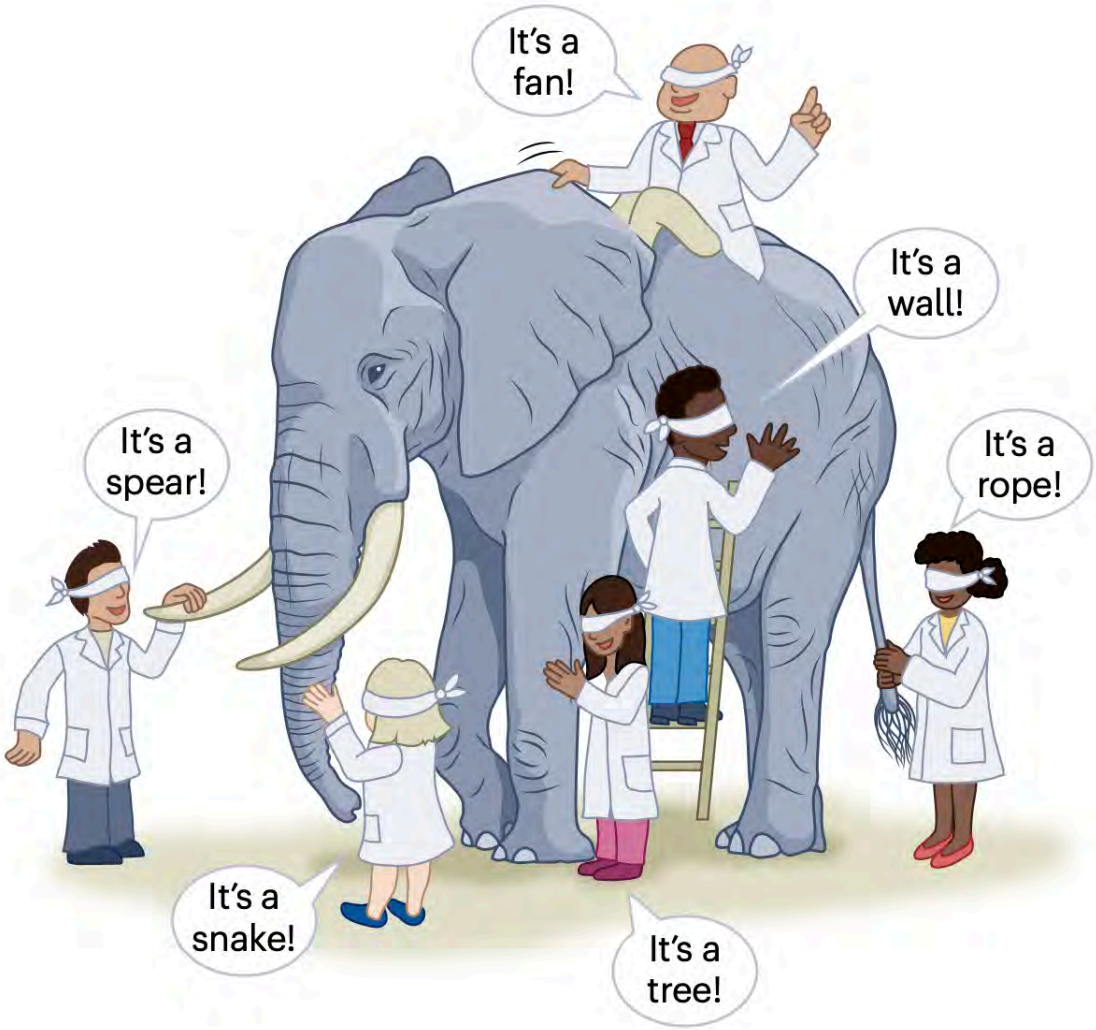




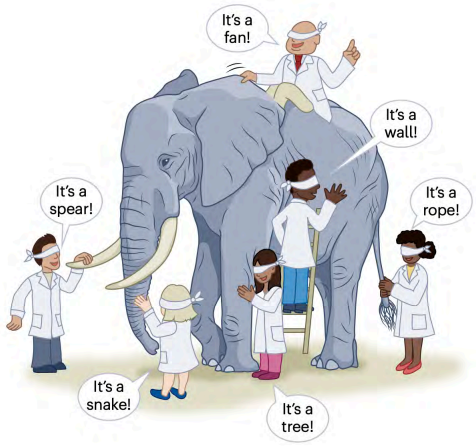


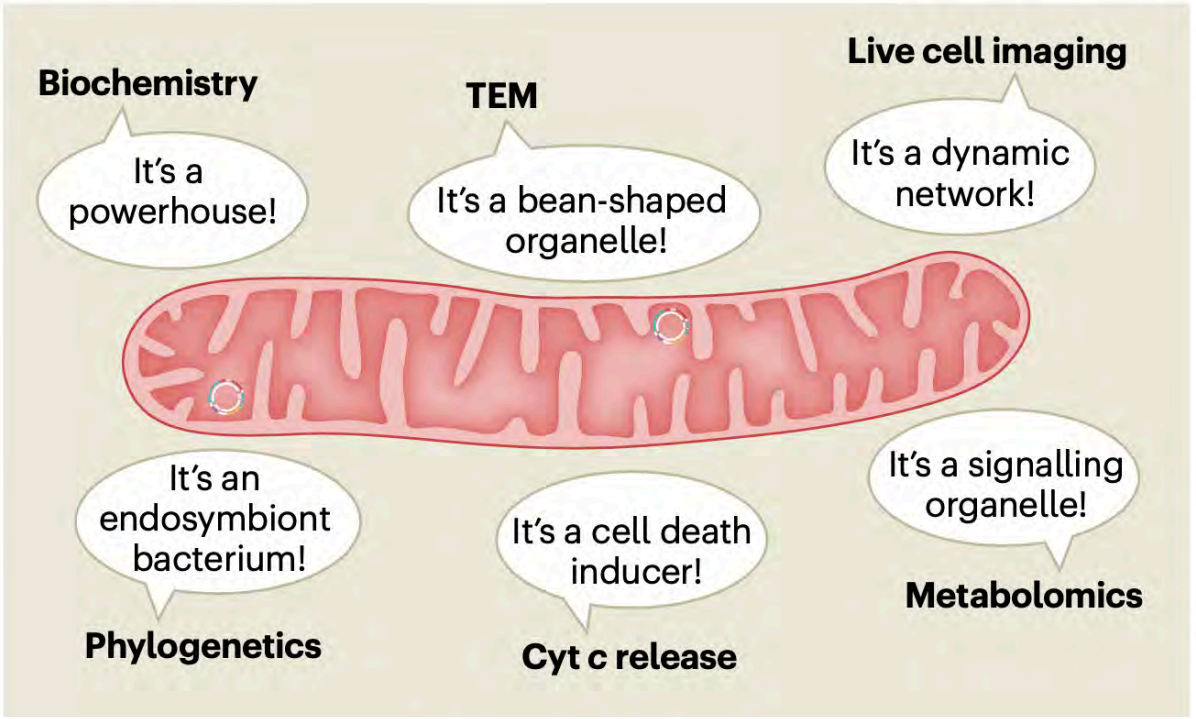
Picard and Shirihai. *Cell Metab* 2022

How do we research mitochondria?



Monzel et al. *Nat Metab* 2023

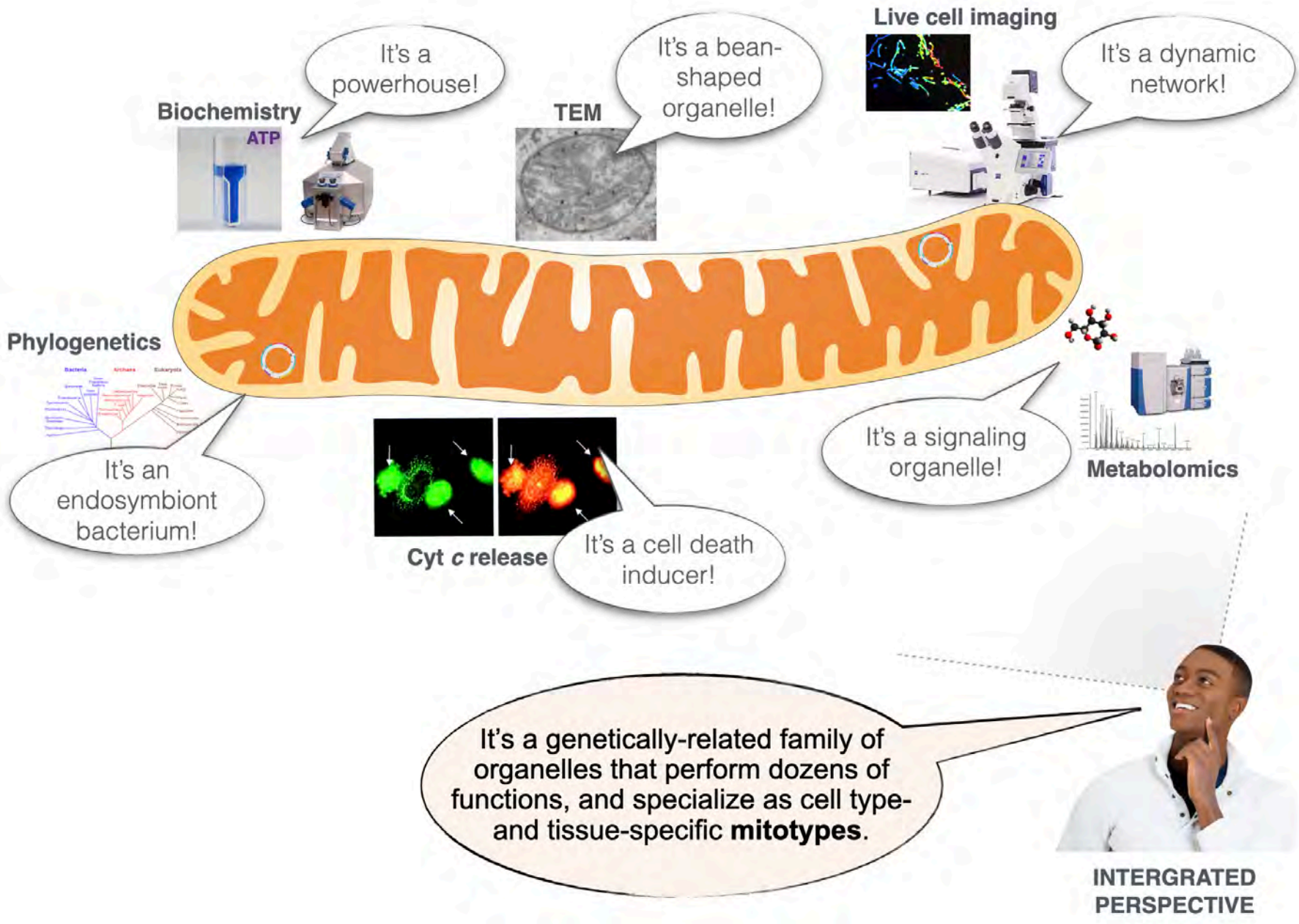




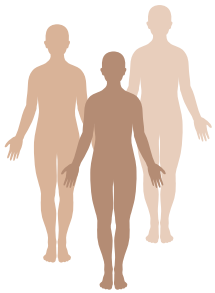
Integrated perspective

It's a family of organelles that exist as distinct mitochondrial phenotypes, defined by their molecular and morphological features, activities, functions and behaviours

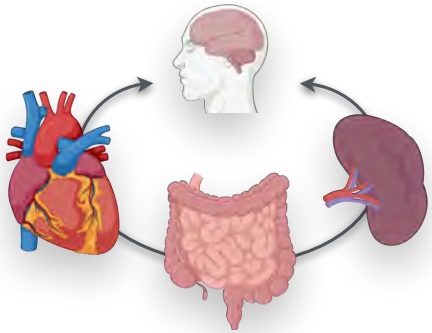
Monzel et al. *Nat Metab* 2023


Biochemistry
ATP
It's a powerhouse!
TEM
It's a bean-shaped organelle!
Live cell imaging
It's a dynamic network!
Phylogenetics
Bacteria
Archaea
Eukaryota
It's an endosymbiont bacterium!
Cyt c release
It's a cell death inducer!
It's a signaling organelle!
Metabolomics
It's a genetically-related family of organelles that perform dozens of functions, and specialize as cell type- and tissue-specific mitotypes.
INTERGRATED PERSPECTIVE

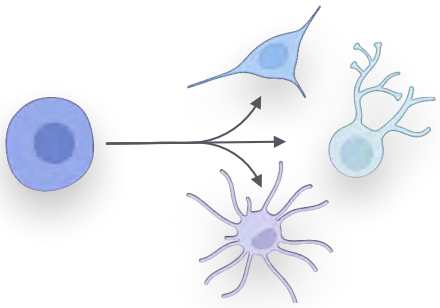
a **Domains of human health**



- Development and growth
- Physical activity
- Wound healing
- Immunity
- Cardiovascular fitness
- Locomotion
- Digestion
- Sleep
- Cognition
- Learning and memory
- Social interactions
- Others...

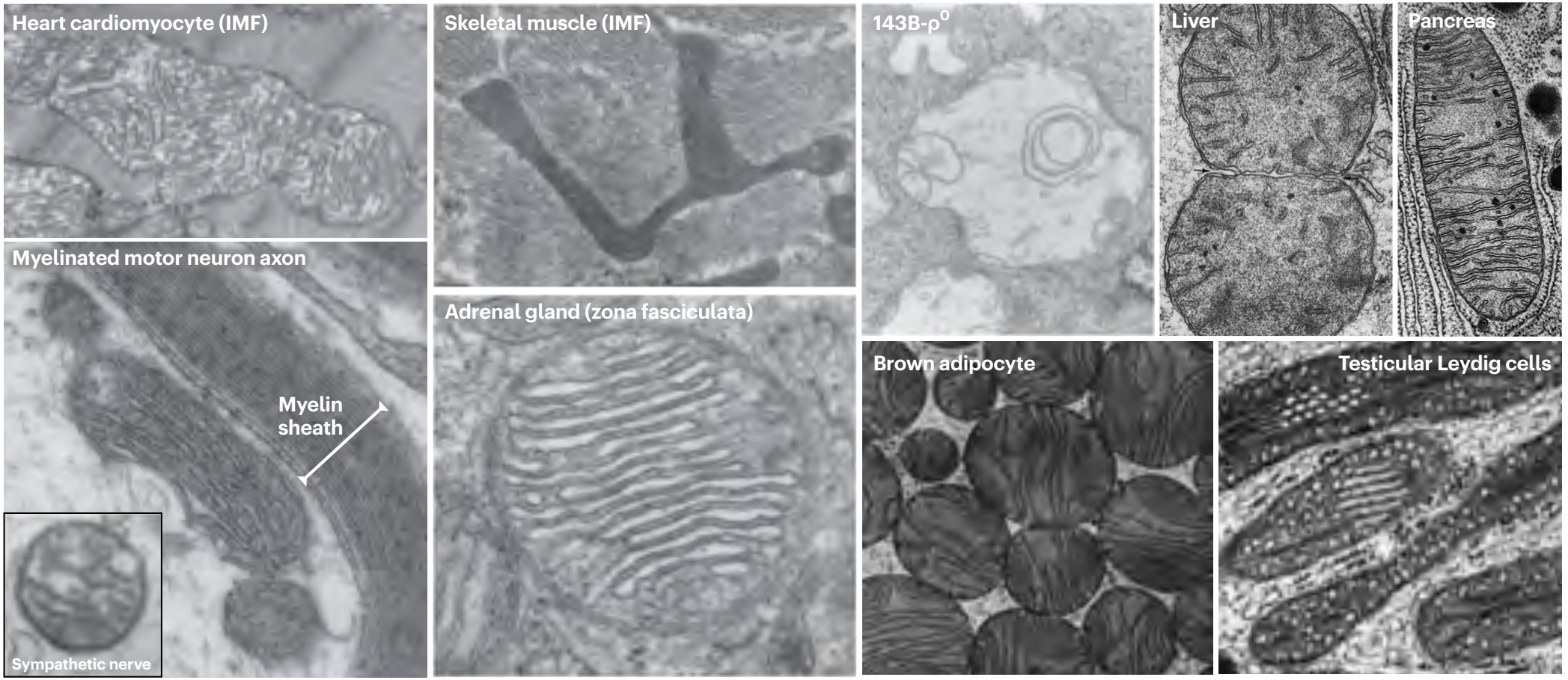


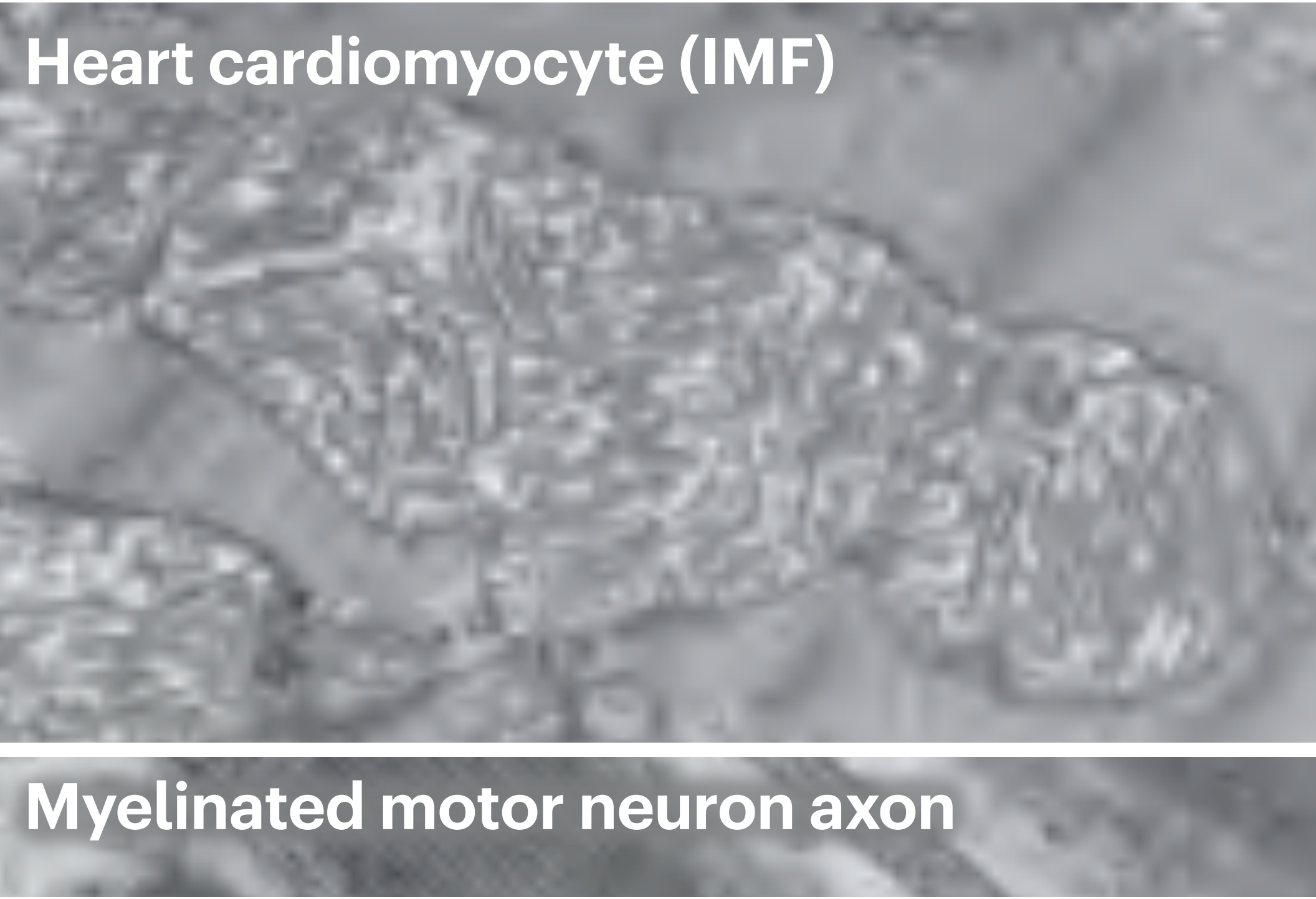
Organ systems

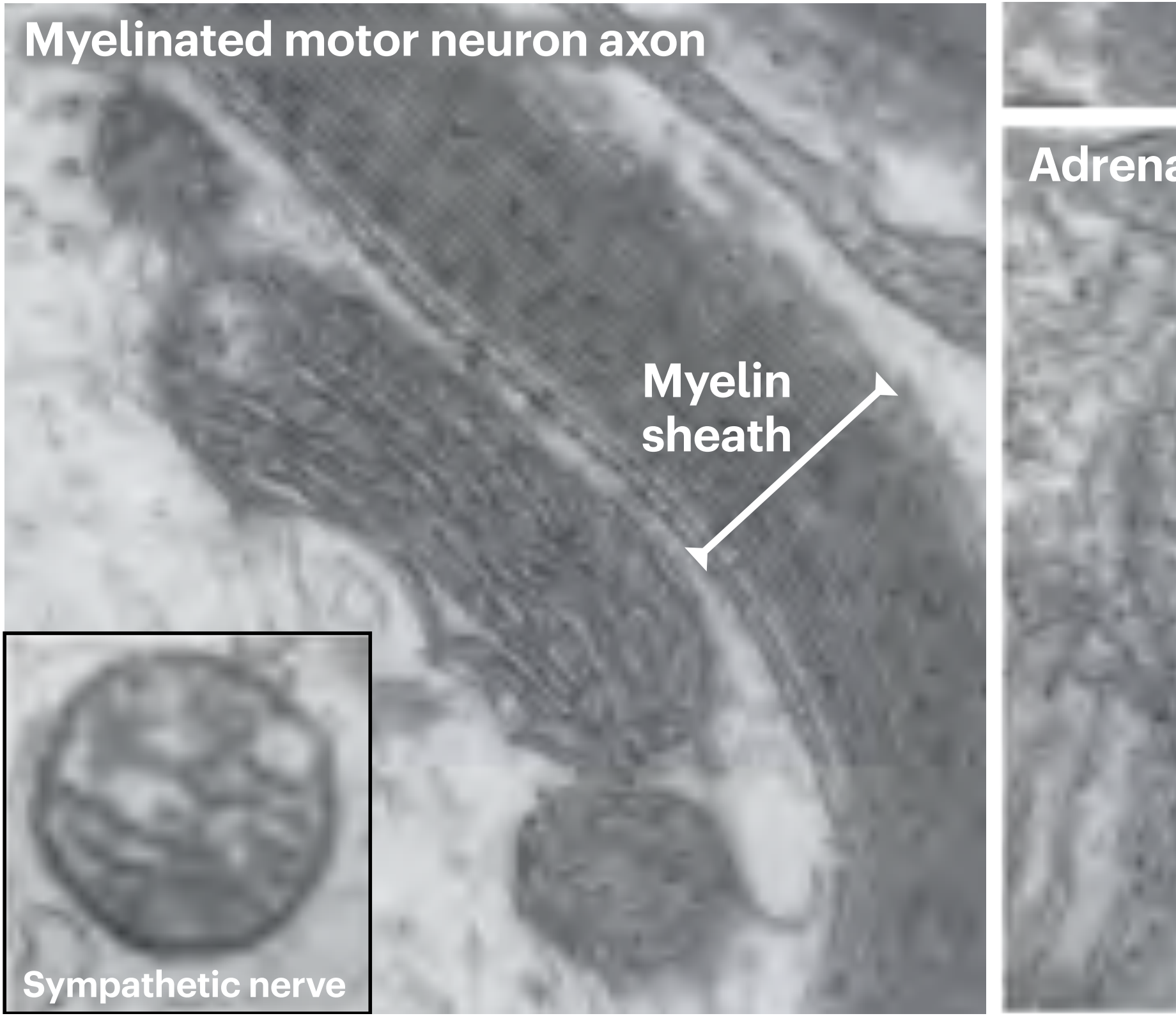


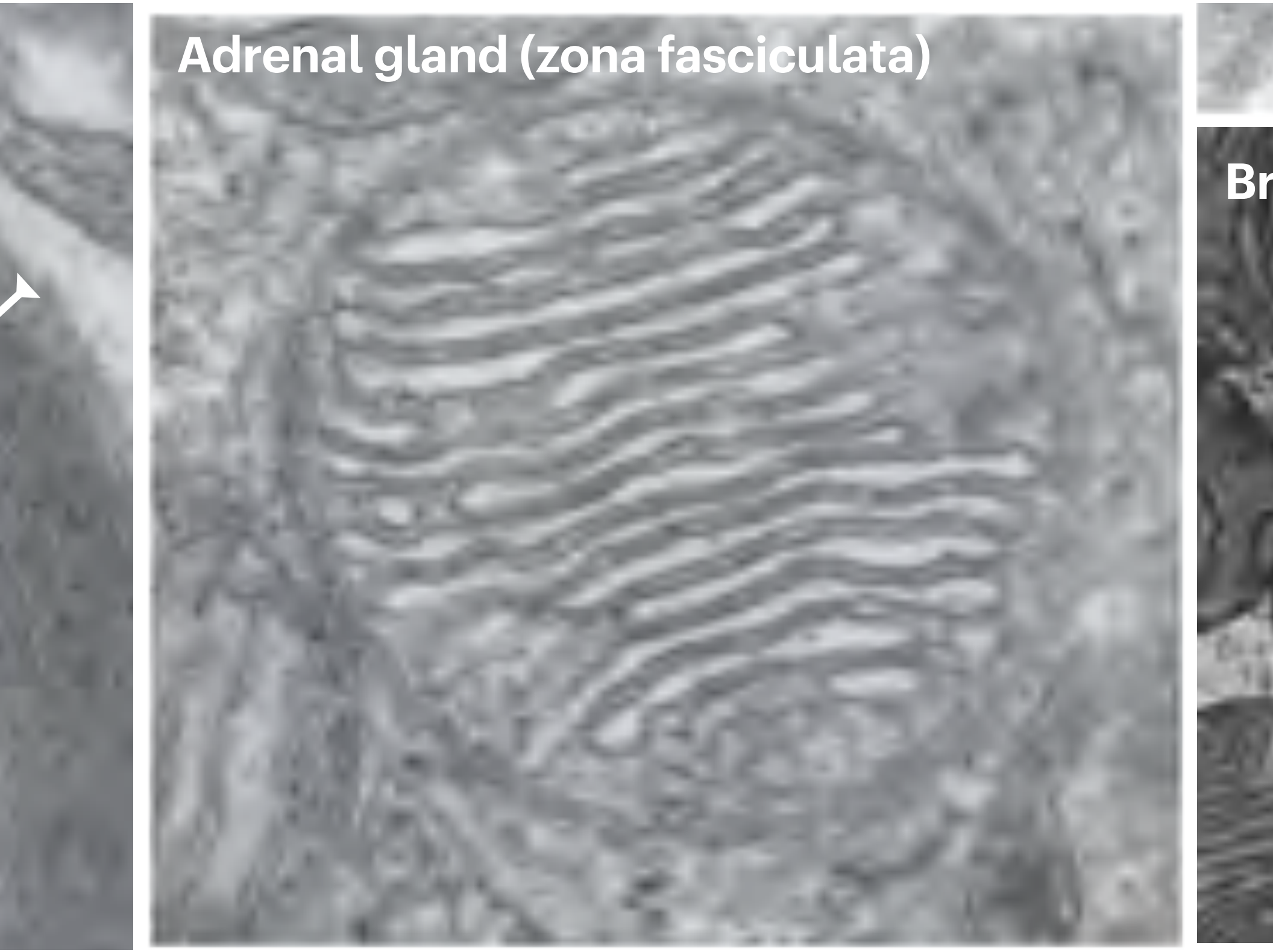
Cell types

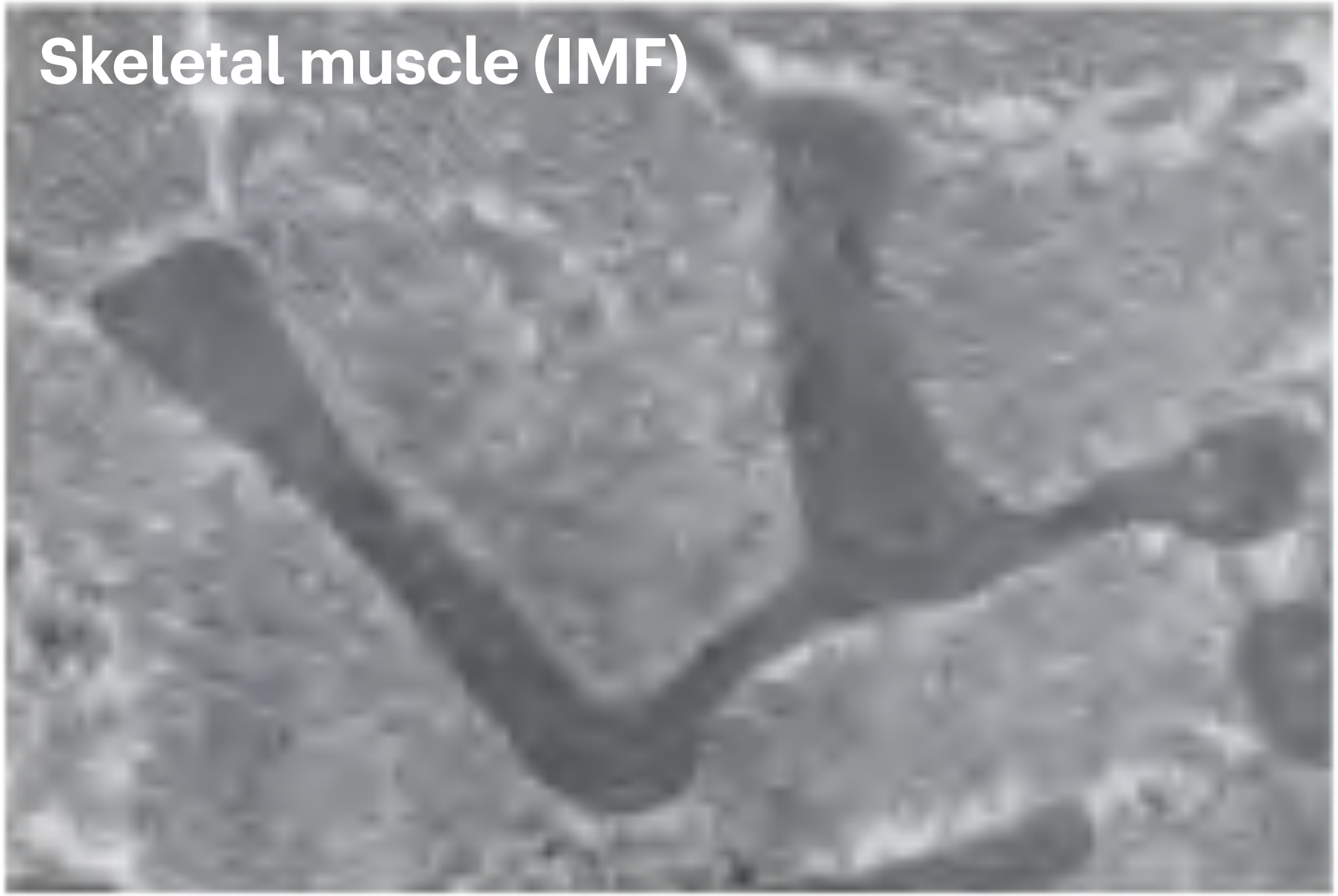
Mitotypes?


Heart cardiomyocyte (IMF)
Skeletal muscle (IMF)
143B-ρ^0
Liver
Pancreas
Myelinated motor neuron axon
Myelin sheath
Sympathetic nerve
Adrenal gland (zona fasciculata)
Brown adipocyte
Testicular Leydig cells


Heart cardiomyocyte (IMF)
Myelinated motor neuron axon

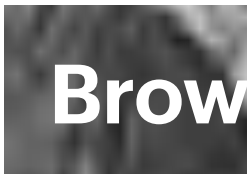

Myelinated motor neuron axon
Myelin sheath
Sympathetic nerve
Adrena

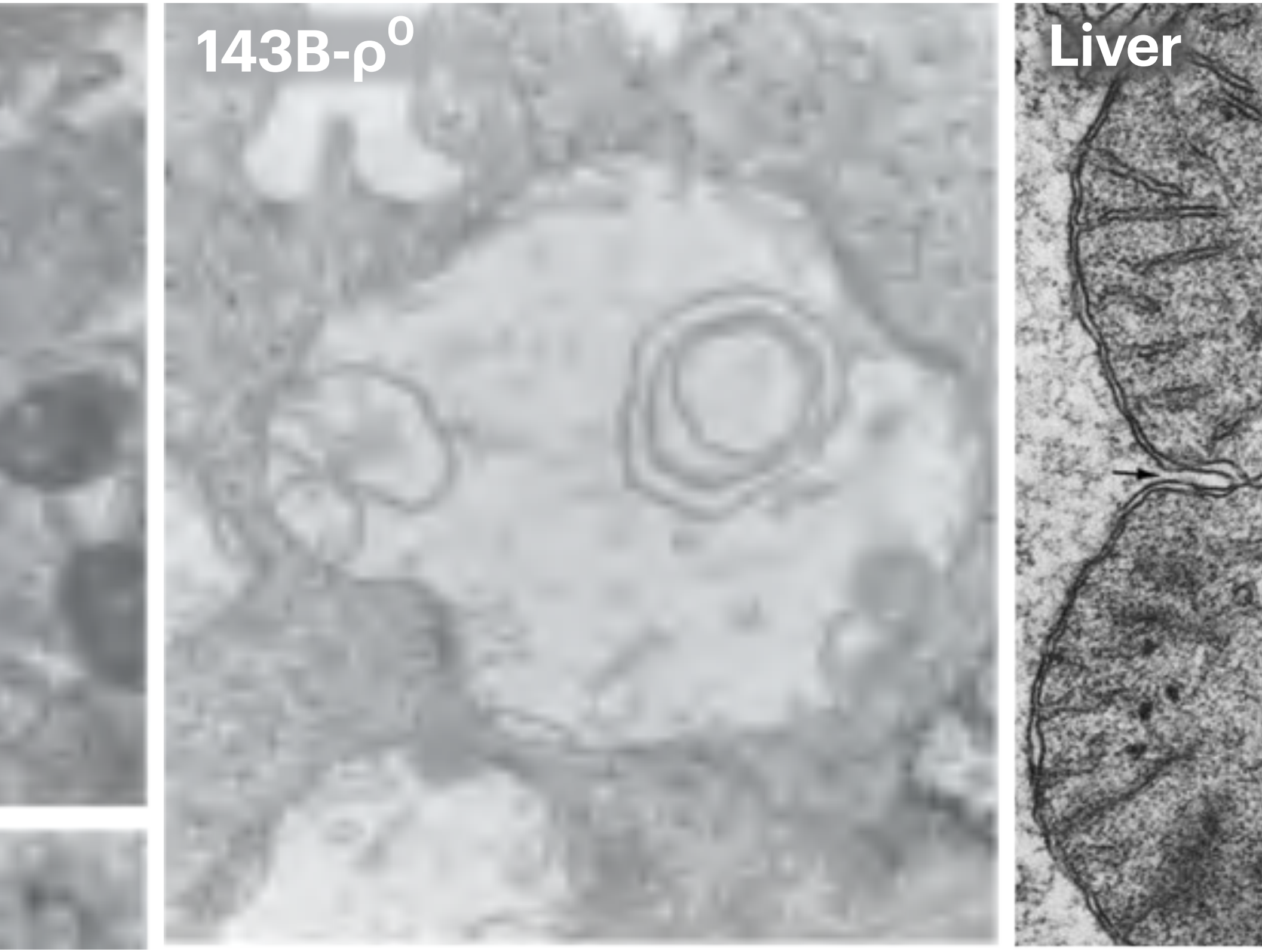

Adrenal gland (zona fasciculata)

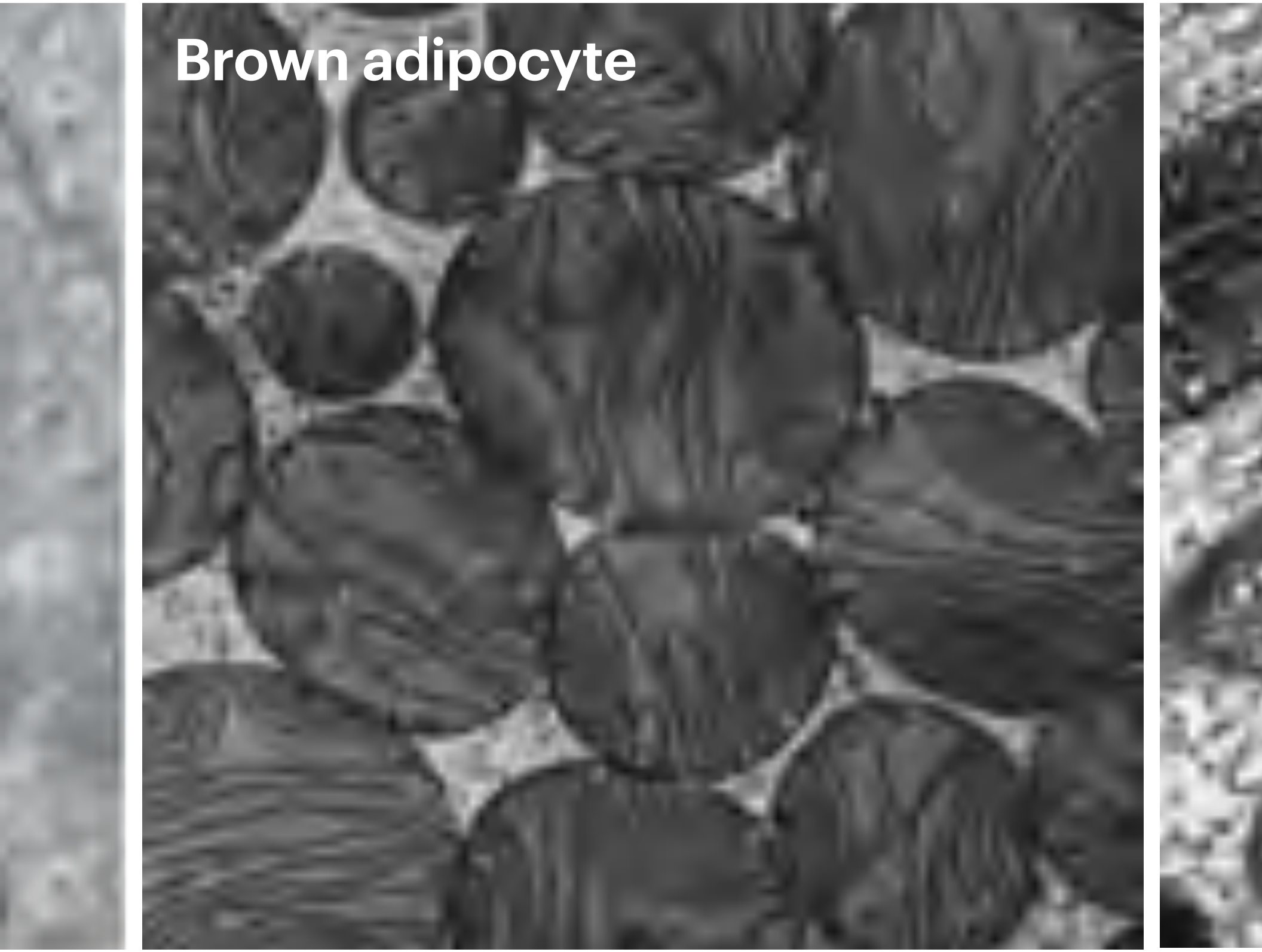

Skeletal muscle (IMF)

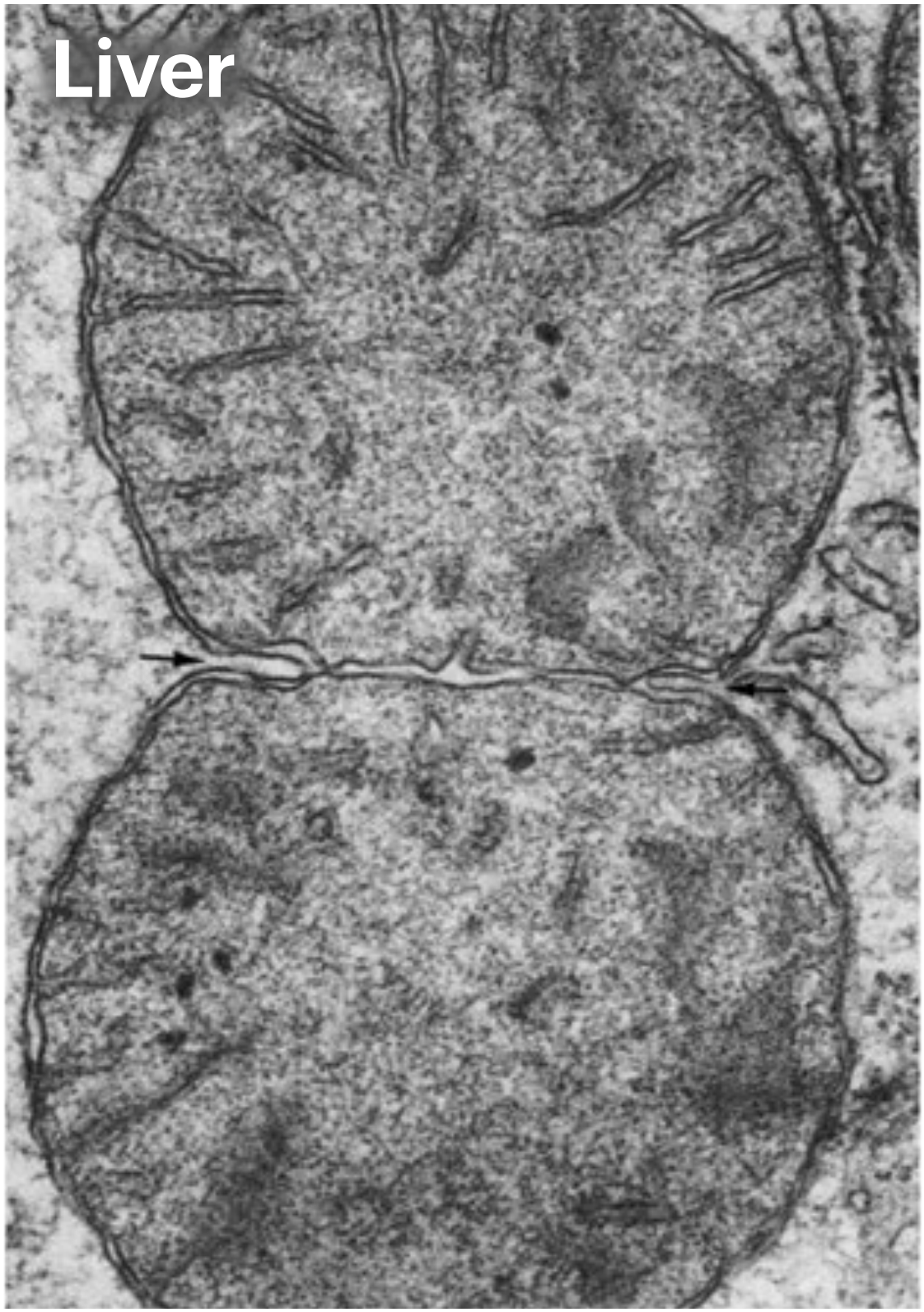

Adrenal gland (zona fasciculata)

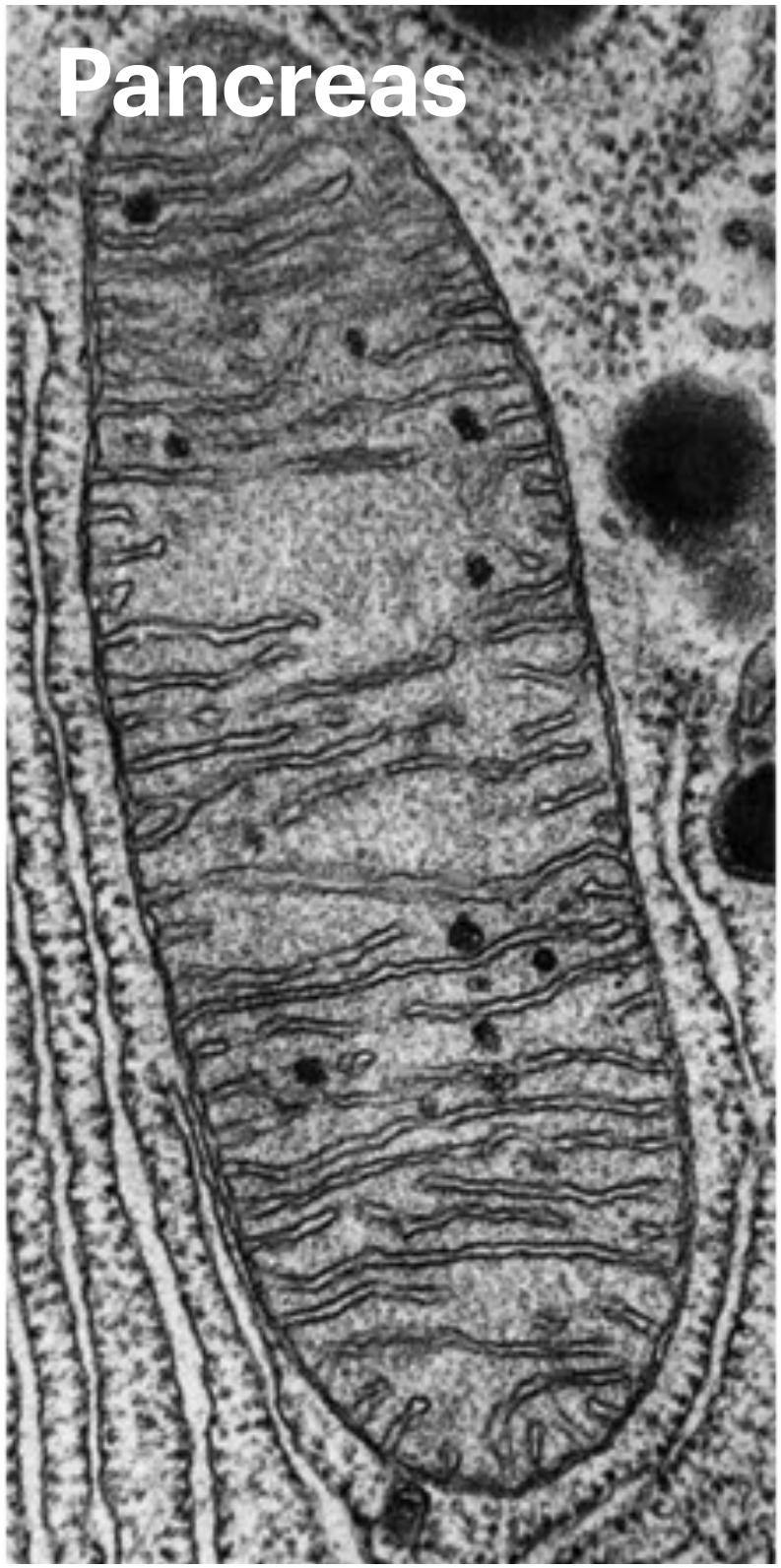

143B-

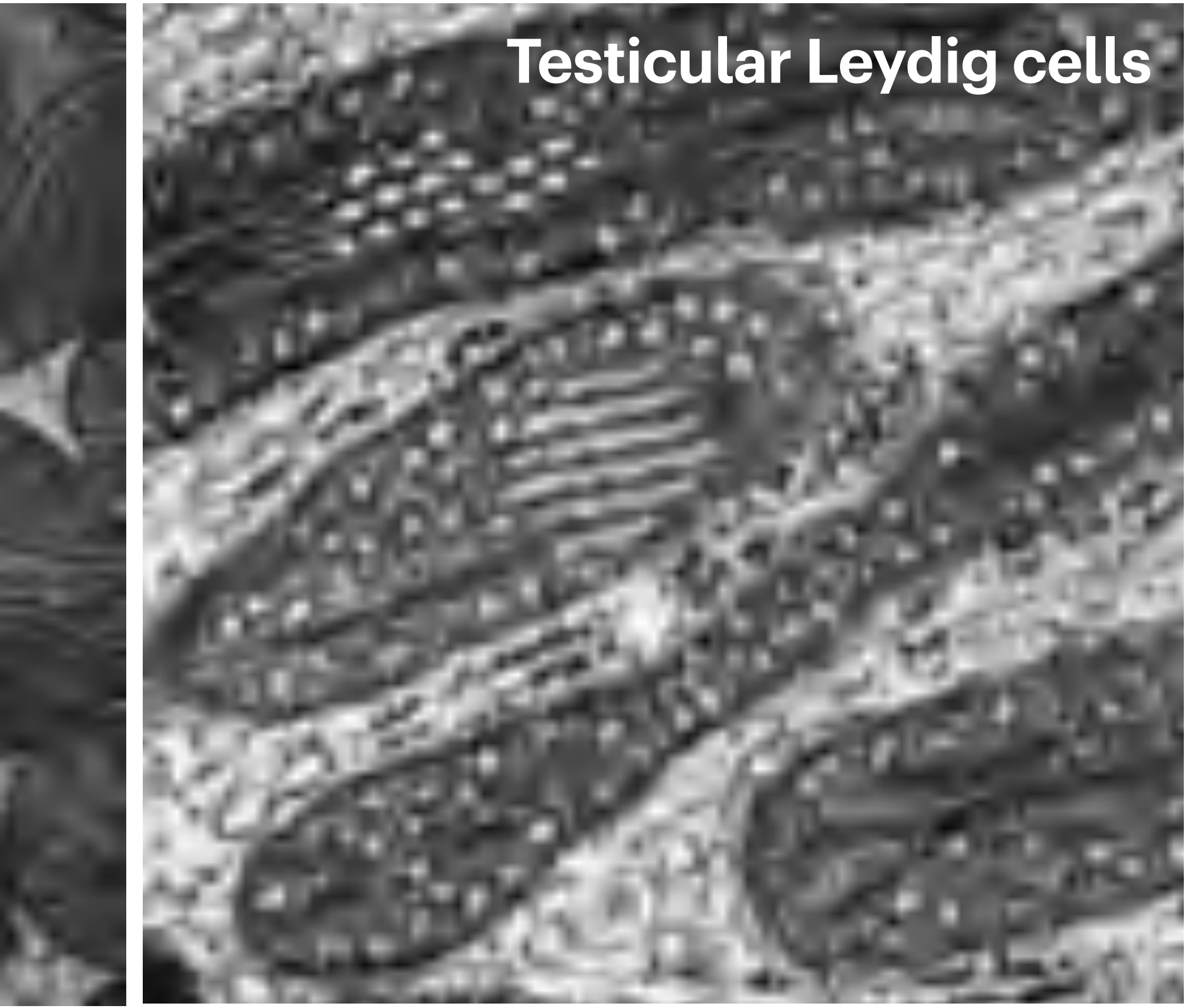

Brow


143B-ρ0
Liver

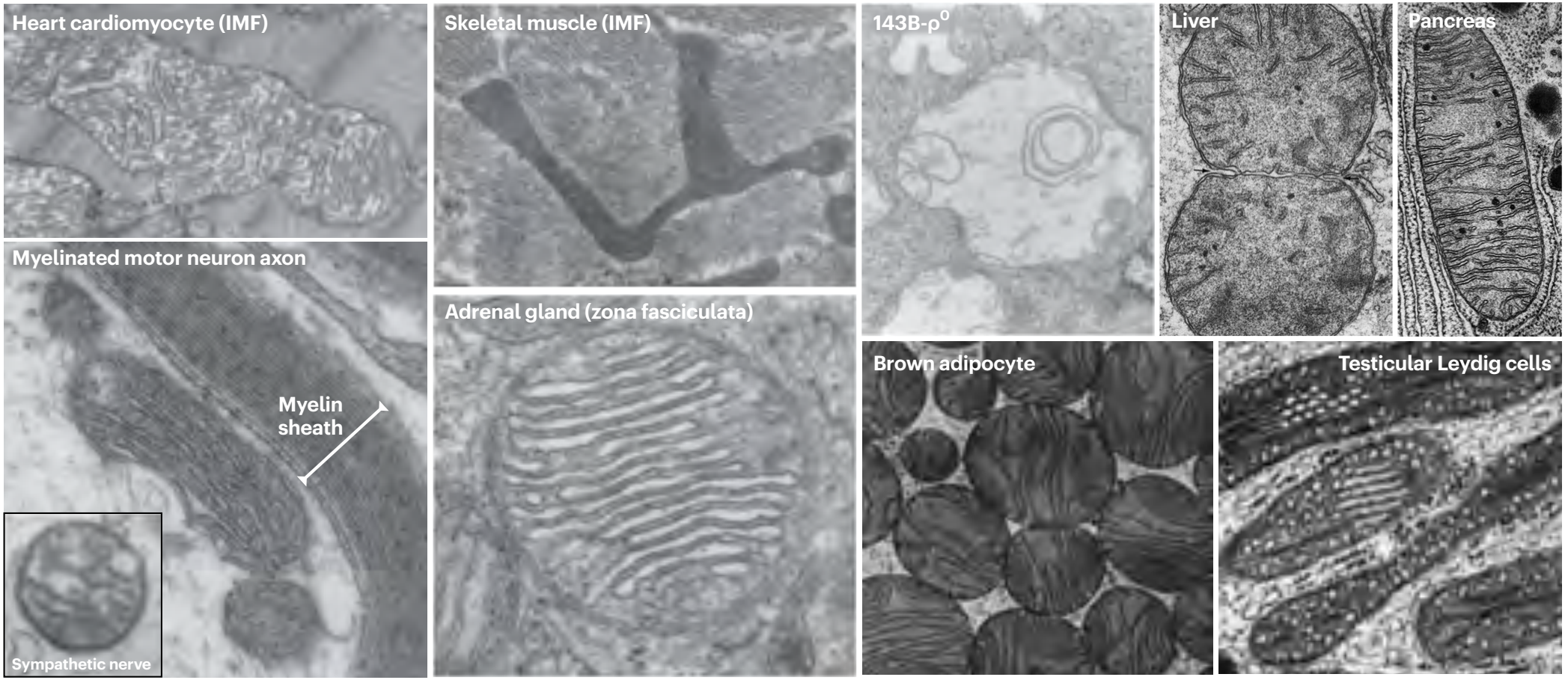

Brown adipocyte


Liver


Pancreas

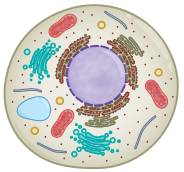

Testicular Leydig cells

The many faces of mitochondria

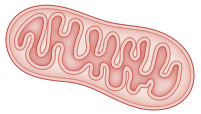


Analogous levels of biology

Organism

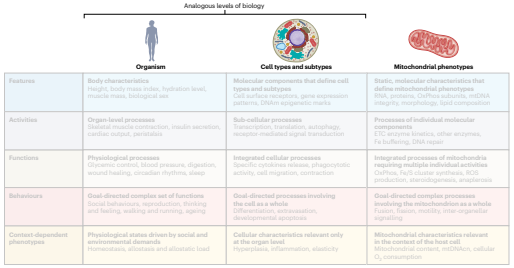


Cell types and subtypes

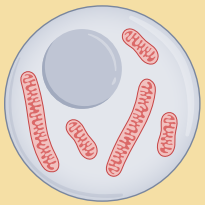


Mitochondrial phenotypes

Features	**Body characteristics** Height, body mass index, hydration level, muscle mass, biological sex	**Molecular components that define cell types and subtypes** Cell surface receptors, gene expression patterns, DNAm epigenetic marks	**Static, molecular characteristics that define mitochondrial phenotypes** RNA, proteins, OxPhos subunits, mtDNA integrity, morphology, lipid composition
Activities	**Organ-level processes** Skeletal muscle contraction, insulin secretion, cardiac output, peristalsis	**Sub-cellular processes** Transcription, translation, autophagy, receptor-mediated signal transduction	**Processes of individual molecular components** ETC enzyme kinetics, other enzymes, Fe buffering, DNA repair
Functions	**Physiological processes** Glycemic control, blood pressure, digestion, wound healing, circadian rhythms, sleep	**Integrated cellular processes** Specific cytokines release, phagocytotic activity, cell migration, contraction	**Integrated processes of mitochondria requiring multiple individual activities** OxPhos, Fe/S cluster synthesis, ROS production, steroidogenesis, anaplerosis
Behaviours	**Goal-directed complex set of functions** Social behaviours, reproduction, thinking and feeling, walking and running, ageing	**Goal-directed processes involving the cell as a whole** Differentiation, extravasation, developmental apoptosis	**Goal-directed complex processes involving the mitochondrion as a whole** Fusion, fission, motility, inter-organellar signalling
Context-dependent phenotypes	**Physiological states driven by social and environmental demands** Homeostasis, allostasis and allostatic load	**Cellular characteristics relevant only at the organ level** Hyperplasia, inflammation, elasticity	**Mitochondrial characteristics relevant in the context of the host cell** Mitochondrial content, mtDNAcn, cellular O_2 consumption



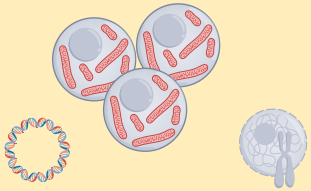
Cell-dependent phenotypes



Mito content/mass

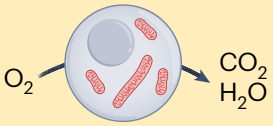
Volume density

mtDNAcn



500,000 — Copies — 1,000
mtDNAcn: 500

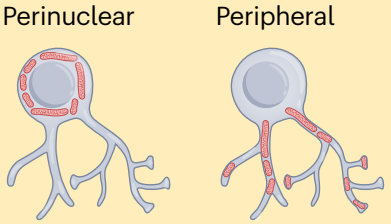
Cellular oxygen consumption



(pmol O_2/cell/min)

Energy expenditure

Cellular topology



Features

mtDNA sequence and integrity

SNPs, mutations, deletions

Molecular composition

Protein dynamics

Assembled protein complexes

Ultrastructure

Cristae structure

Morphology

Size, length, complexity

Activities

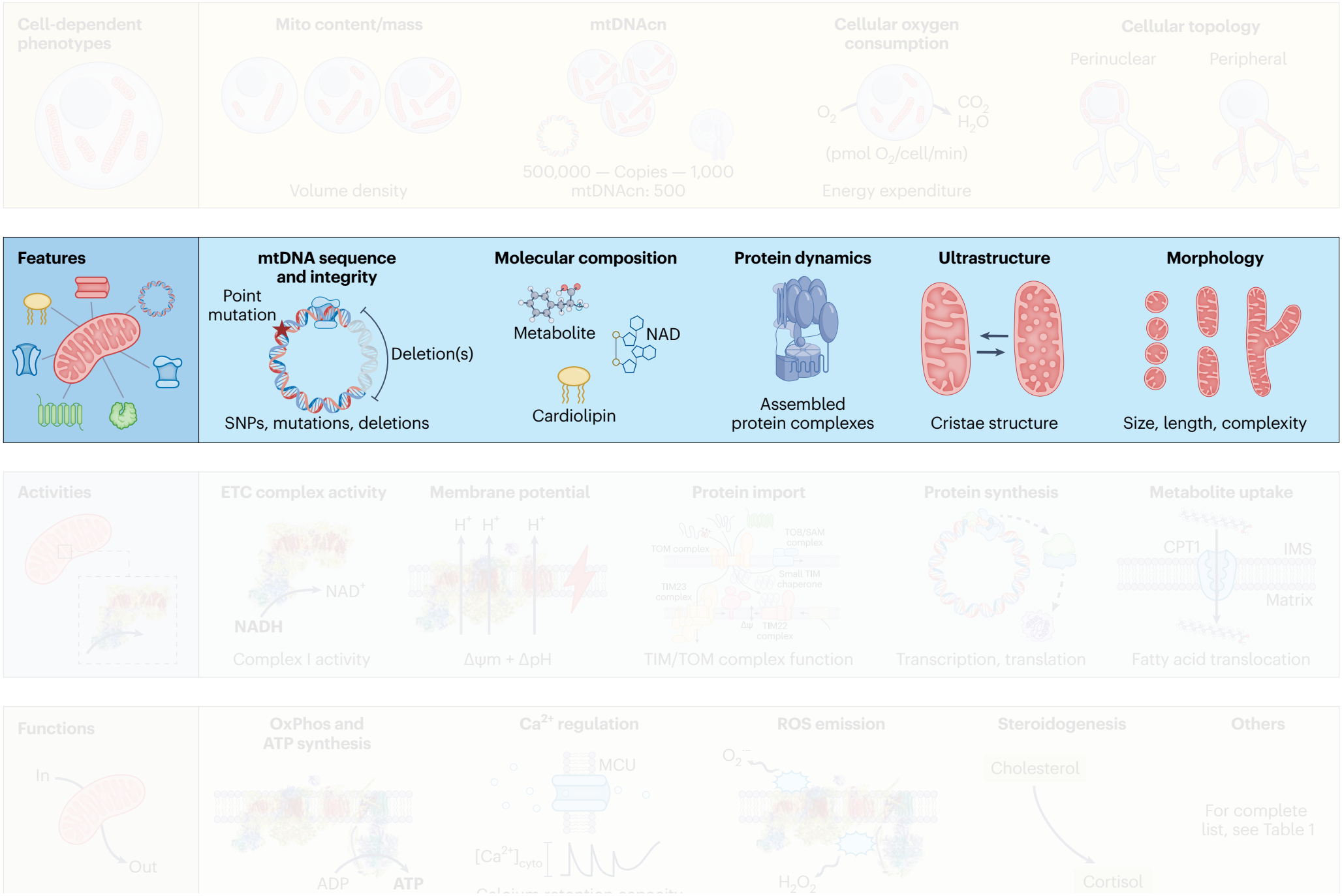
ETC complex activity

Membrane potential

Protein import

Protein synthesis

Metabolite uptake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Mito content/mass
Volume density
mtDNAcn
500,000 — Copies — 1,000
mtDNAcn: 500
Cellular oxygen consumption
O2
CO2
H2O
(pmol O2/cell/min)
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Cholesterol
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Others
For complete list, see Table 1

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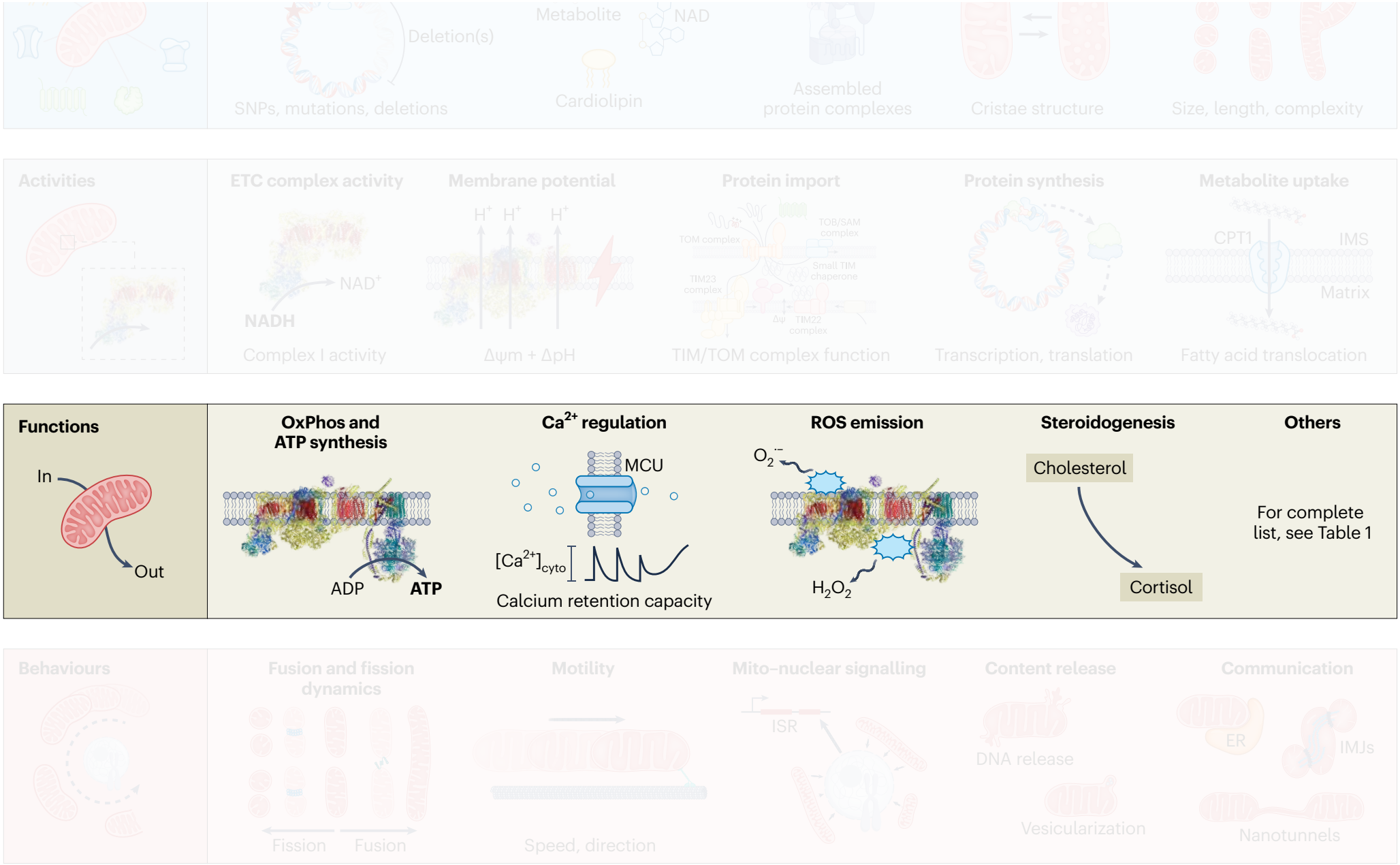
Fusion

Motility
Speed, direction

Mito–nuclear signalling
ISR

Content release
DNA release
Vesicularization

Communication
ER
IMJs
Nanotunnels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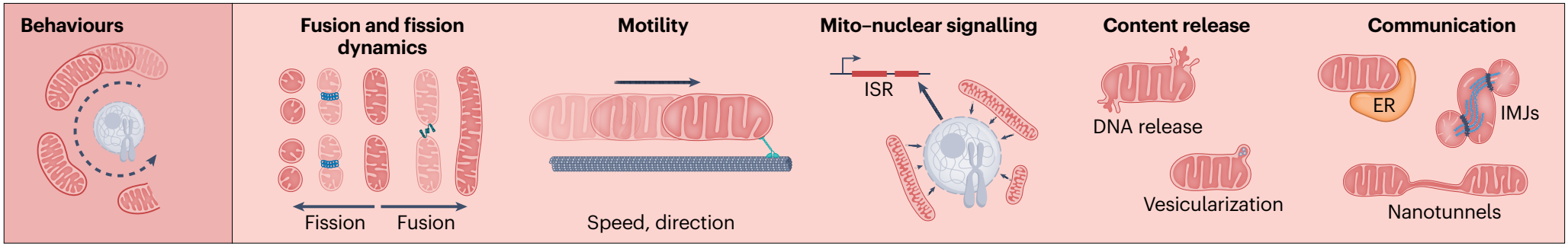
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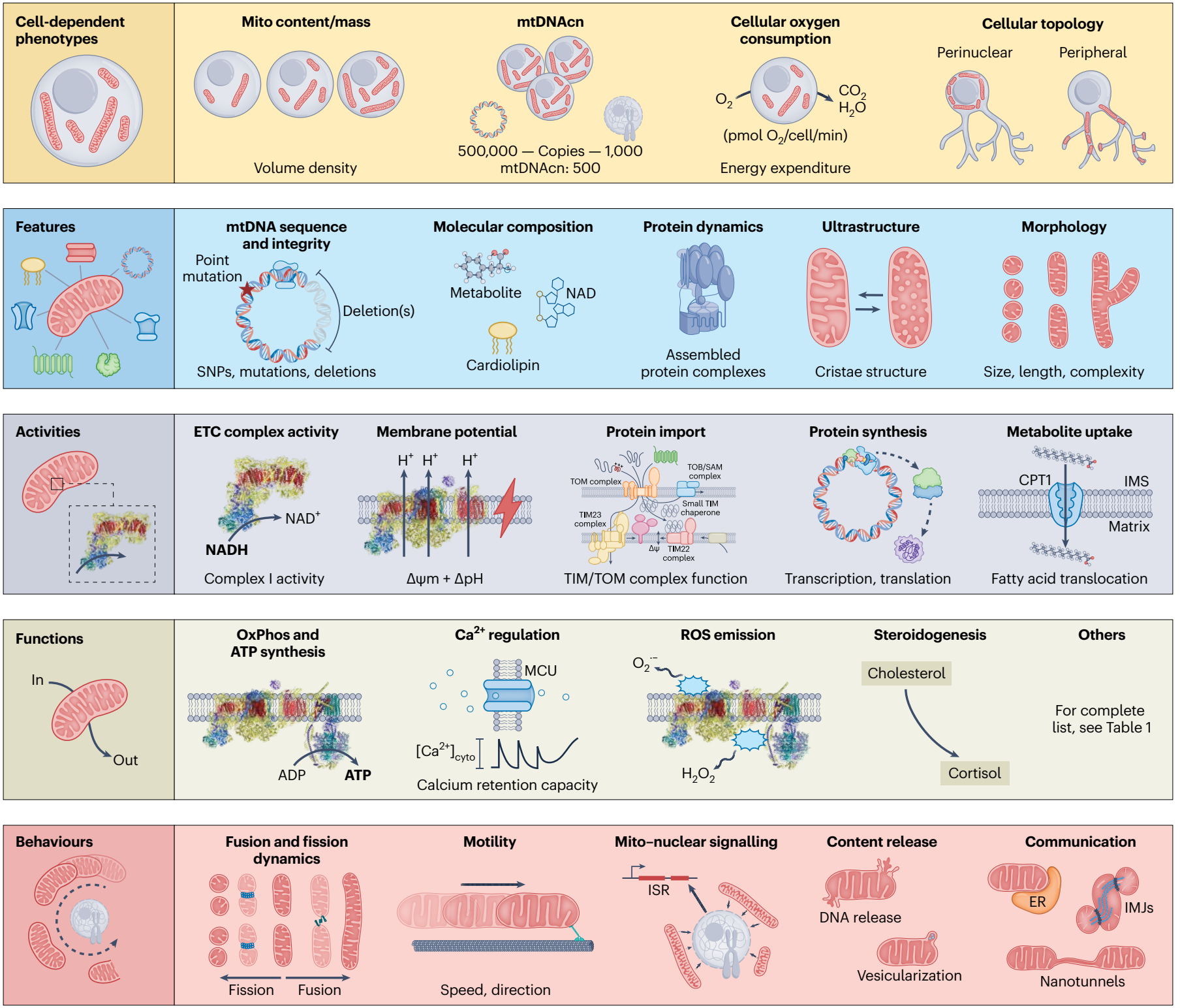
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https://doi.org/10.1038/s42255-023-00783-1

Table 1 | Mitochondrial functions and behaviours

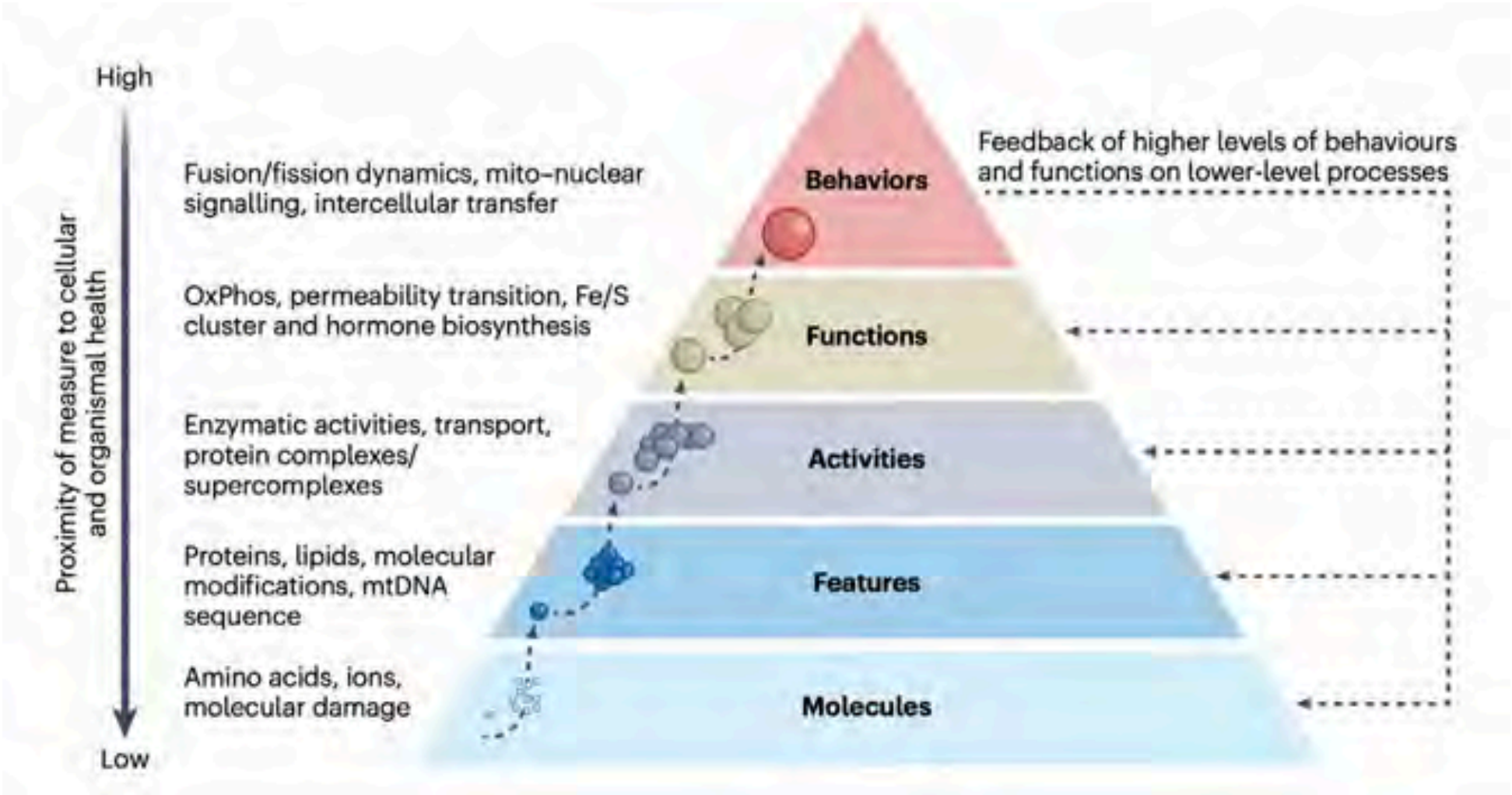
	Description	Reviewed in ref(s).	Methods described in ref(s).
Functions			
[a]**Membrane potential generation**	Formation of the electrochemical gradient (ΔΨm+ΔpH) across the IMM, usually by the electron pumping capacity of the respiratory complexes I, III and IV, but also by other processes including through ATP hydrolysis by the F_oF_1 ATP synthase (complex V).	104	105,106
Amino acid metabolism	Lysine metabolism (lysine-α-ketoglutarate reductase, encoded by *AASS*). Electrogenic malate–aspartate shuttle system, which is important for balancing pyridine dinucleotide redox states across subcellular compartments. Branched-chain keto and amino acids. Choline and derivatives as structural precursors for lipoproteins, membrane lipids and the neurotransmitter acetylcholine. Betaine as osmoregulator and an intermediate in the cytosolic transulfuration pathway.	107–111	112–119
Ascorbate metabolism	L-ascorbate (vitamin C) biosynthesis in many plants and animals, but not in primates, which serves as osmoregulator and antioxidant. Mitochondria may recycle oxidized (dehydro)ascorbic acid.	120	121,122
Bicarbonate metabolism	Production of bicarbonate (HCO_3^-) by mitochondrial carbonic anhydrase V (encoded by *CA5A*), used as a cofactor for anaplerotic reactions (for example, ureagenesis and gluconeogenesis) and acid–base balance. The TCA cycle is an important contributor to cellular/extracellular acidification due to CO_2 production.	123	–
Calcium uptake and extrusion	Uptake of cytoplasmic Ca^{2+} via the mitochondrial calcium uniporter in a ΔΨm-dependent manner; extrusion by the sodium/calcium exchanger NCLX (encoded by *SLC8B1*).	124–126	127,128
Hydrogen sulfide detoxification	Mitochondrial sulfide quinone oxidoreductase (encoded by *SQOR*) oxidizes hydrogen sulfide to glutathione persulfide by reducing CoQ.	129–132	133
Heat production	Heat generation is stimulated by uncoupling ΔΨm+ΔpH from ATP synthesis (thereby increasing electron flux and respiration) by UCP1 (encoded by *UCP1*), the ADP/ATP carrier (*AAC*, also *ANT1*), or by creatine-dependent substrate cycling and other futile cycles.	134–137	138
Intermediate metabolism	Enzymatic interconversion of metabolic intermediates to enable the synthesis of specific macromolecules, including five major anaplerotic ones. This includes the conversion of pyruvate into oxaloacetate by pyruvate carboxylase (encoded by *PC*), a critical step for de novo glucose synthesis (gluconeogenesis); citrate export to the cytoplasm where it is used for lipid synthesis or converted to acetyl-CoA for acetylation reactions; synthesis of itaconate, a derivative of *cis*-aconitate; succinate, α-ketoglutarate and others that participate in a variety of signalling	25,139,140	141,142

Lipid oxidation	Beta-oxidation of long-chain, medium-chain and short-chain fatty acids into acetyl-CoA.	145	146
Lipid synthesis	Synthesis of cardiolipin and phosphatidylethanolamine from ER precursors in the IMM.	147–150	–
mtDNA maintenance and expression	mtDNA replication, transcription, protein synthesis and assembly of the OxPhos system.	151,152	153,154
Na^+import/export	Sodium (Na^+) uptake and release against cytoplasmic Ca^{2+} by the sodium/calcium exchanger protein NCLX (encoded by *SLC8B1*) or by Na^+/H^+ antiporter (molecular identity pending).	124,155	156
Neurotransmitter synthesis and degradation	Synthesis of the cofactor BH4 (tetrahydrobiopterin), used by hydrolase enzymes to synthesize catecholamines and neurotransmitters (serotonin, melatonin, norepinephrine and epinephrine) and nitric oxide. Mitochondria with OMM-anchored monoamine oxidases (encoded by *MAOA* and *MAOB*, donate electrons and contribute to electron flow in the ETC) also degrade catecholamines. Mitochondria also participate in GABA metabolism.	9,157	158,159
One-carbon metabolism and pyrimidine synthesis	The one-carbon metabolism connects the synthesis of nucleotides (purine and pyrimidine), amino acids (methionine, serine and glycine), *S*-adenosyl-methionine and folate. Ubiquinone-mediated oxidation of dihydroorotate to orotate by dihydroorotate dehydrogenase (encoded by *DHODH*) is a key step in pyrimidine synthesis.	160–163	164
OxPhos	Transduction of $\Delta\Psi m + \Delta pH$ generated by the electron transport chain (ETC, also 'respiratory chain') into ATP synthesis by the F_oF_1 ATP synthase (complex V), abbreviated as OxPhos.	165	166
Oxygen sensing	The electron transport and free-radical generation by ETC complexes I and III is modulated by the partial pressure of oxygen, which can limit respiration at very low partial pressures of O_2.	167–170	–
Permeability transition	Opening of the high-conductance permeability transition pore (PTP), which dissipates membrane potential and promotes the release of intracristae and matrix-located components into the cytoplasm.	171,172	173–175
Protein import	Import, processing and folding of nuclear-encoded polypeptides from the cytoplasm by the translocator of the inner membrane (TIM) and outer membrane (TOM) complexes and associated proteins.	176	–
Redox homeostasis	Re-oxidation of enzymes and/or their redox cofactors (involved in anabolic and catabolic reactions) by the electron acceptors CoQ and cytochrome c (encoded by *CYTC*) within the mitochondrial respiratory chain, and production of NADPH by *NNT*.	177,178	–
Respiration	Electrons stored in reducing equivalents NADH and $FADH_2$, or derived from diverse redox reactions are sequentially delivered to respiratory complex I and CoQ, or cytochrome c, respectively, to promote the reduction of molecular oxygen at cytochrome c oxidase (complex IV).	179,180	181
ROS production	Production and release of ROS (H_2O_2, $O_2^{\bullet-}$, others) mainly at respiratory chain complexes I and III.	182,183	184
Steroidogenesis	Production of pregnanolone from cholesterol imported via IMM steroidogenic	33,34,185,186	187

	cytochrome c oxidase (complex IV).		
ROS production	Production and release of ROS (H_2O_2, $O_2^{\bullet-}$, others) mainly at respiratory chain complexes I and III.	182,183	184
Steroidogenesis	Production of pregnanolone from cholesterol imported via IMM steroidogenic acute regulatory protein (encoded by *STAR*) followed by enzymatic transformation by P450ssc (encoded by *CYP11A1*) in the matrix. Intermediate or terminal steps for some steroids occur in the ER. Cytochrome P450 family members participate also in xenobiotic metabolism as well as bile acid and vitamin D biosynthesis.	33,34,185,186	187
Behaviours			
Antiviral signalling	Assembly of the mitochondrial antiviral signal (encoded by *MAVS*) adaptor protein on the OMM to potentiate downstream signalling, and activation of nuclear interferon pathways in the nucleus by mtDNA release.	39,188	–
Apoptotic signalling	Release of cytochrome c (encoded by *CYCS*), apoptosis-inducing factor (encoded by *AIF*), and other proteins that trigger different forms of cell death by acting on cytoplasmic and nuclear effectors.	189,190	–
Cristae remodelling	Dynamic remodelling of IMM cristae junctions, cristae shape and distribution via the combined action of optic atrophy 1 (encoded by *OPA1*) and mitochondrial contact site and cristae organizing system (MICOS) proteins.	103,191	95
DNA signalling	mtDNA extrusion in the cytoplasm, particularly in the form of oxidized mtDNA fragments via proteinaceous pores forming across the IMM and OMM, which trigger inflammasome activation.	189,190,192,193	175
Epigenetic remodelling	Transduction of mitochondrial states into changes in epigenome via several functions including metabolic intermediates, DNA release, ROS production and others.	30,194	–
Inter-organelle communication	Exchange of information between mitochondria and other organelles, particular the ER, where mitofusin 2 (encoded by *MFN2*) plays a key role in tethering organelles.	195,196	197,198
Mitochondrial dynamics	Mitochondrial fusion and fission through OMM-anchored and IMM-anchored GTPase proteins capable of merging or constricting mitochondrial membranes to enact fragmentation of larger organelles into smaller ones.	191,199–201	202
Mito–mito communication	Exchange of information between mitochondria by soluble signals (for example, ROS-induced ROS release, RIRR), by complete membrane fusion, or by physical extensions of thin protein-carrying OMM and IMM membrane protrusions (that is, nanotunnels) and trans-mitochondrial cristae alignment between energized mitochondria.	203–206	207–209
Motility	Movement of energized mitochondria across the cytoplasm via the combined action of motor and adaptor proteins interacting with cytoskeletal elements.	6,210	211
Vesicle formation	Release of MDVs destined to different cellular fates by the action of motor and accessory proteins acting on the OMM and IMM.	212	213,214

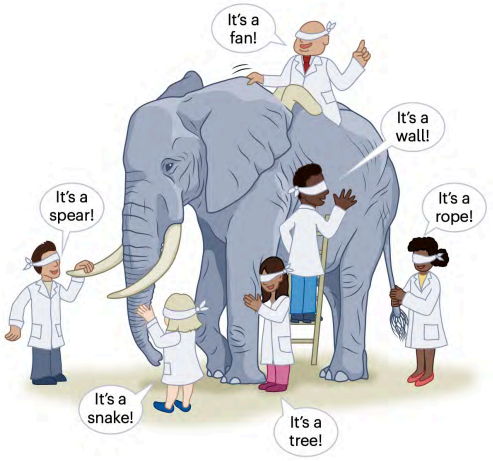
[a]Generation of mitochondrial membrane potential is the 'mother' of many other functions and behaviours, providing the driving force for the movement of ions, solutes and proteins across the IMM, the driving force for key enzymes and processes, including the phosphorylation of ADP into ATP (OxPhos). Mitochondrial features (that is, molecular components) and activities (individual enzyme and non-enzymatic activities) are too numerous to be comprehensively listed, so only functions and behaviours are included. CoQ, coenzyme Q.

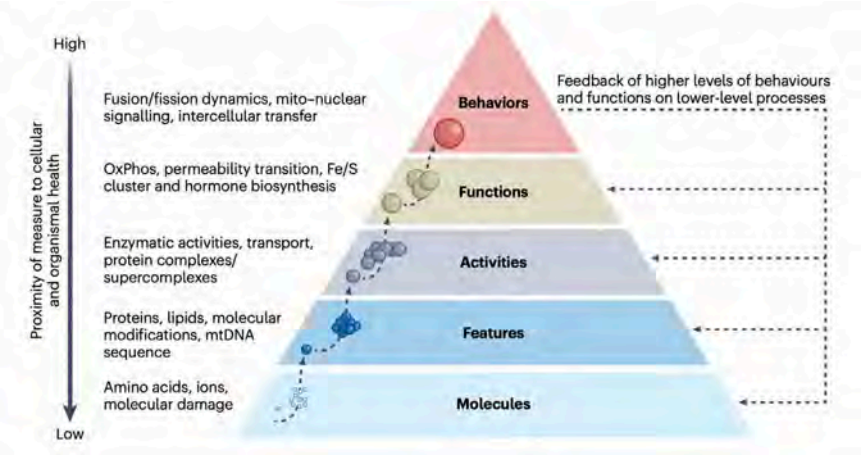
Hierarchy of mitochondrial needs

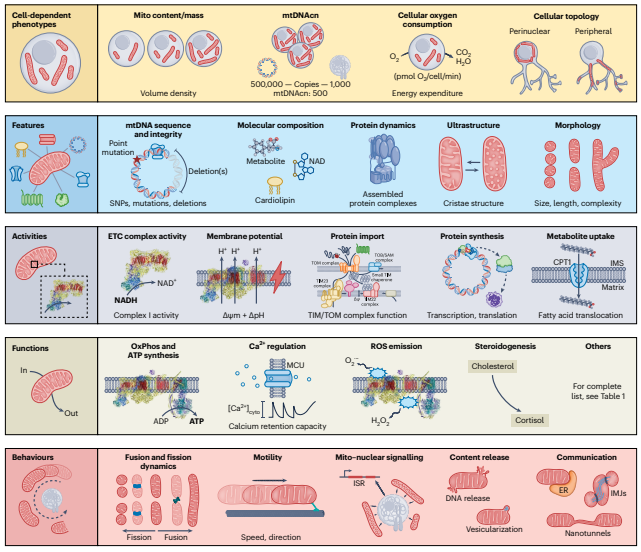


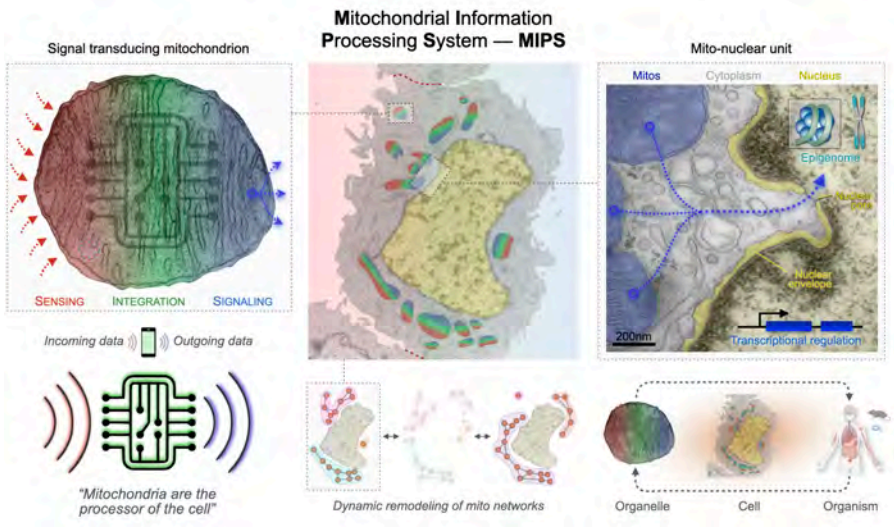
Monzel et al. *Nat Metab* 2023

Mitochondria are diverse, multifunctional organelles that transduce information



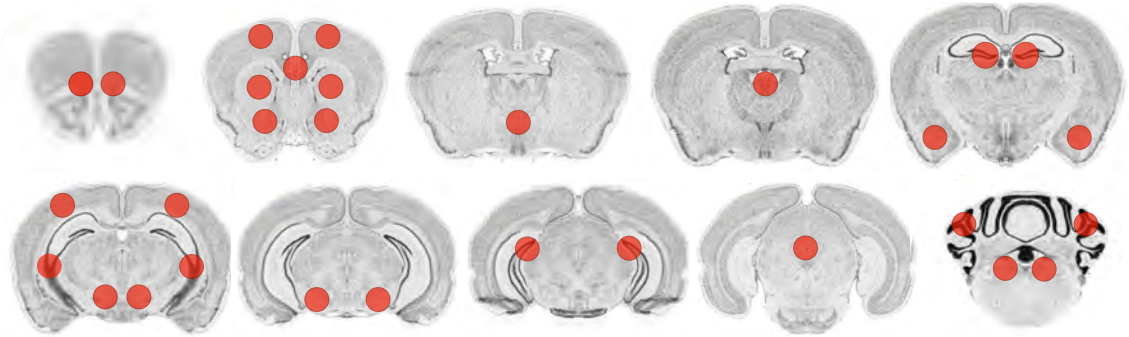


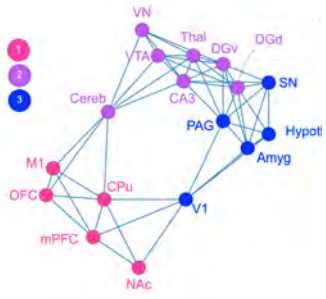


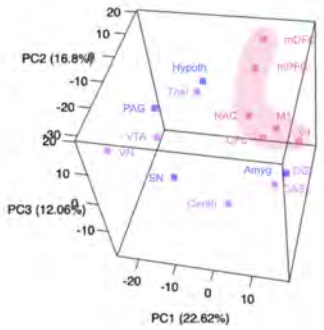


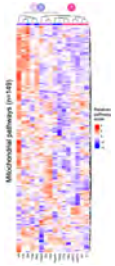
Article https://doi.org/10.1038/s41467-023-39941-0

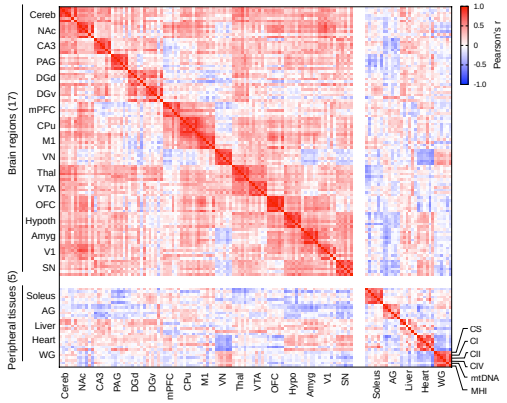
Brain mitochondrial diversity and network organization predict anxiety-like behavior in male mice

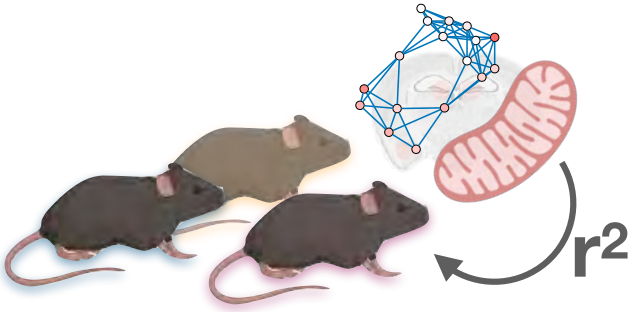






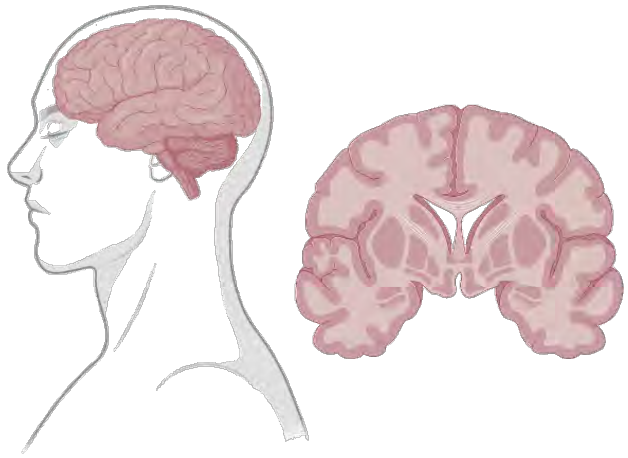




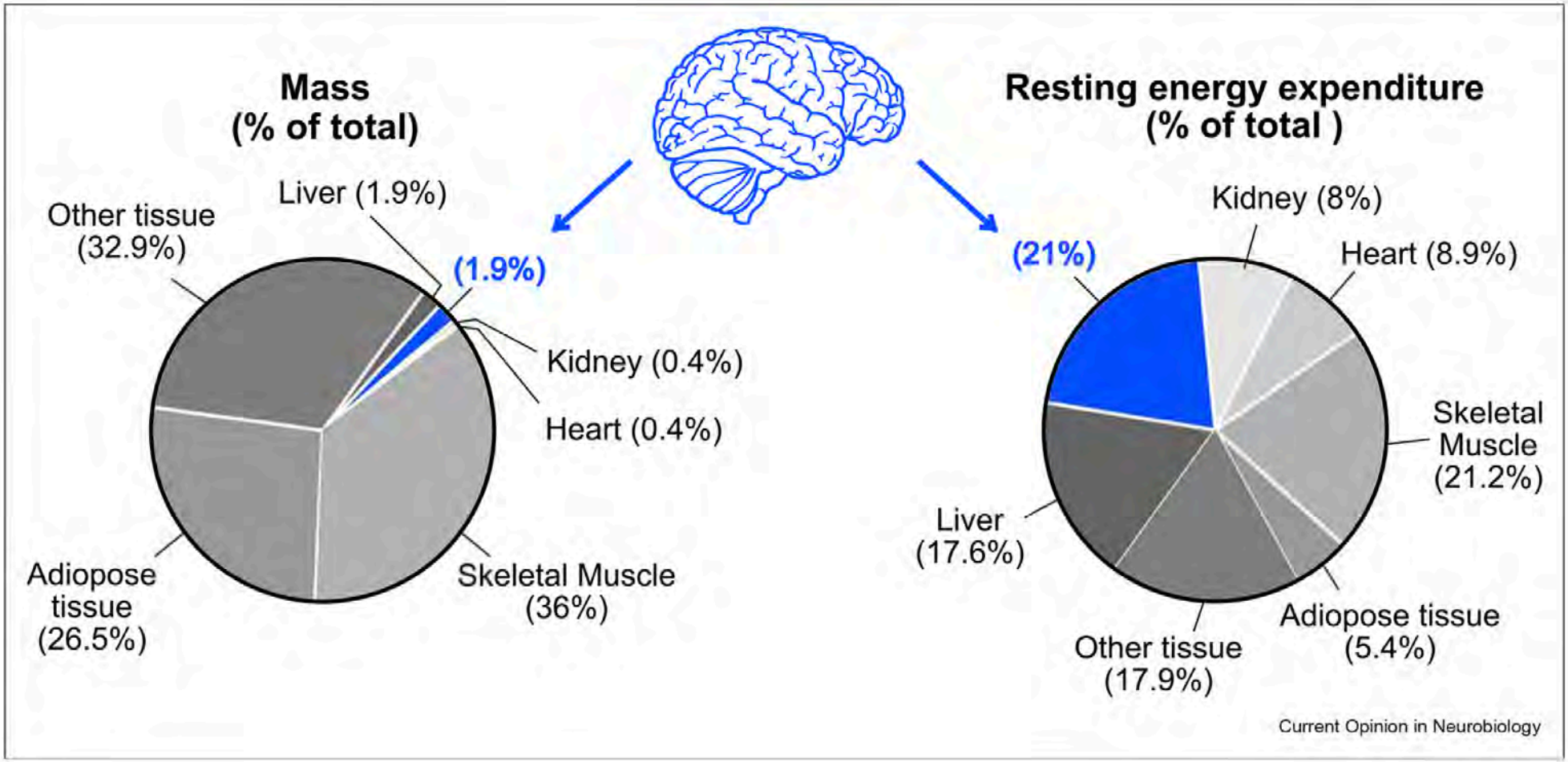


Rosenberg et al. *Nat Commun* 2023

How are mitochondria distributed, and do they specialize across the *human* brain?



The brain's enormous, constant energy demand

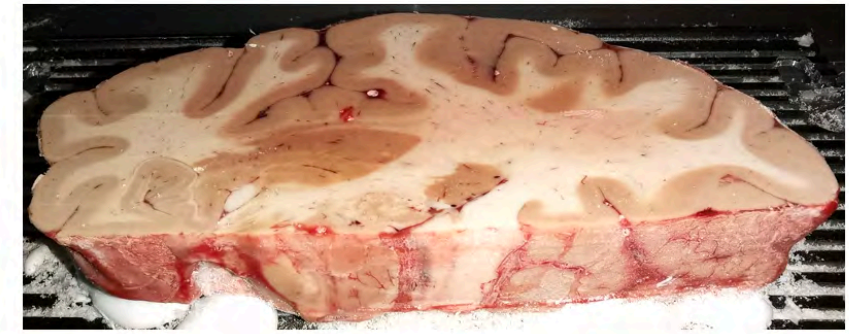


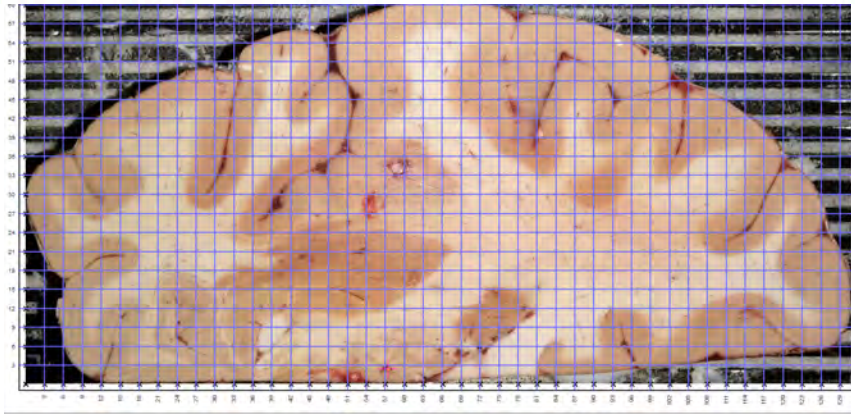
Variations from rest to activity is ~5%

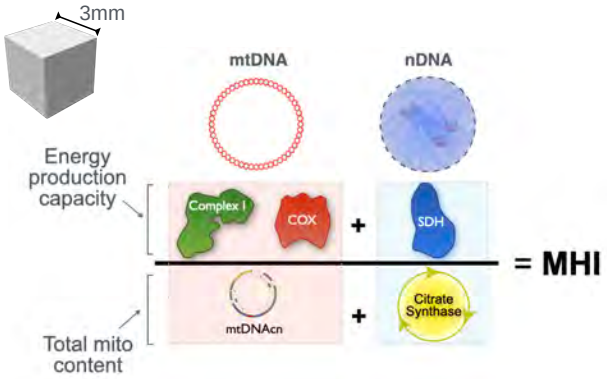
Padamsey and Rochefort. *Curt Opin Neurobiol* 2023

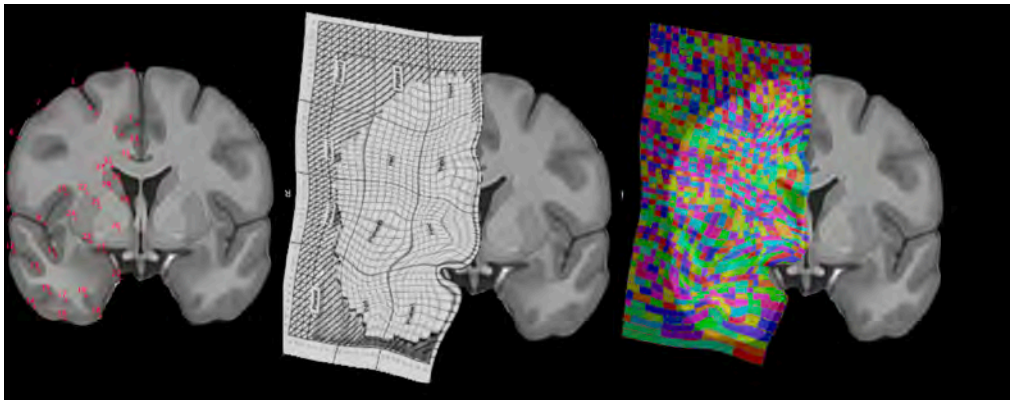
MitoBrainMap v1.0

A multi-function mitochondrial atlas of a single human coronal brain section at fMRI resolution



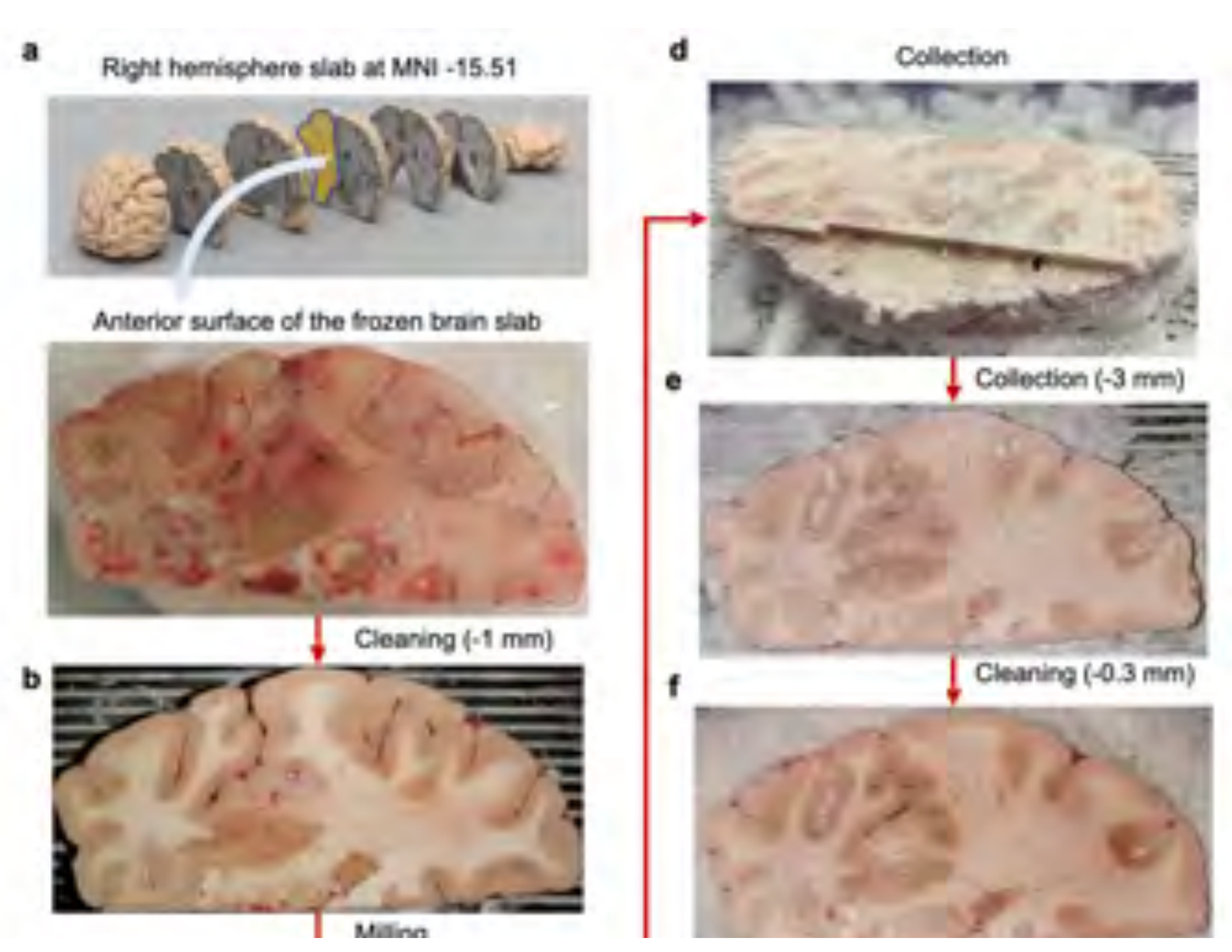


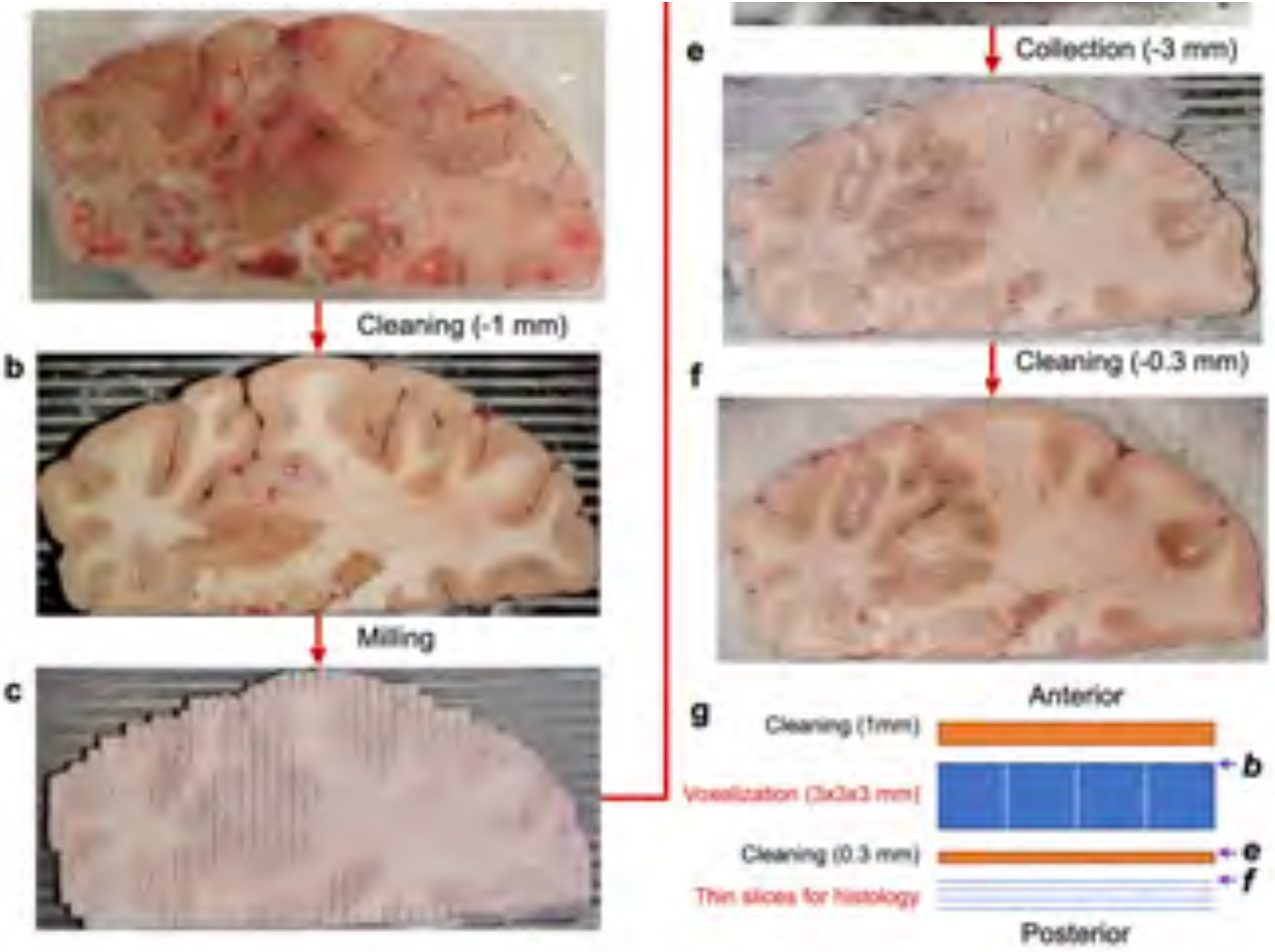


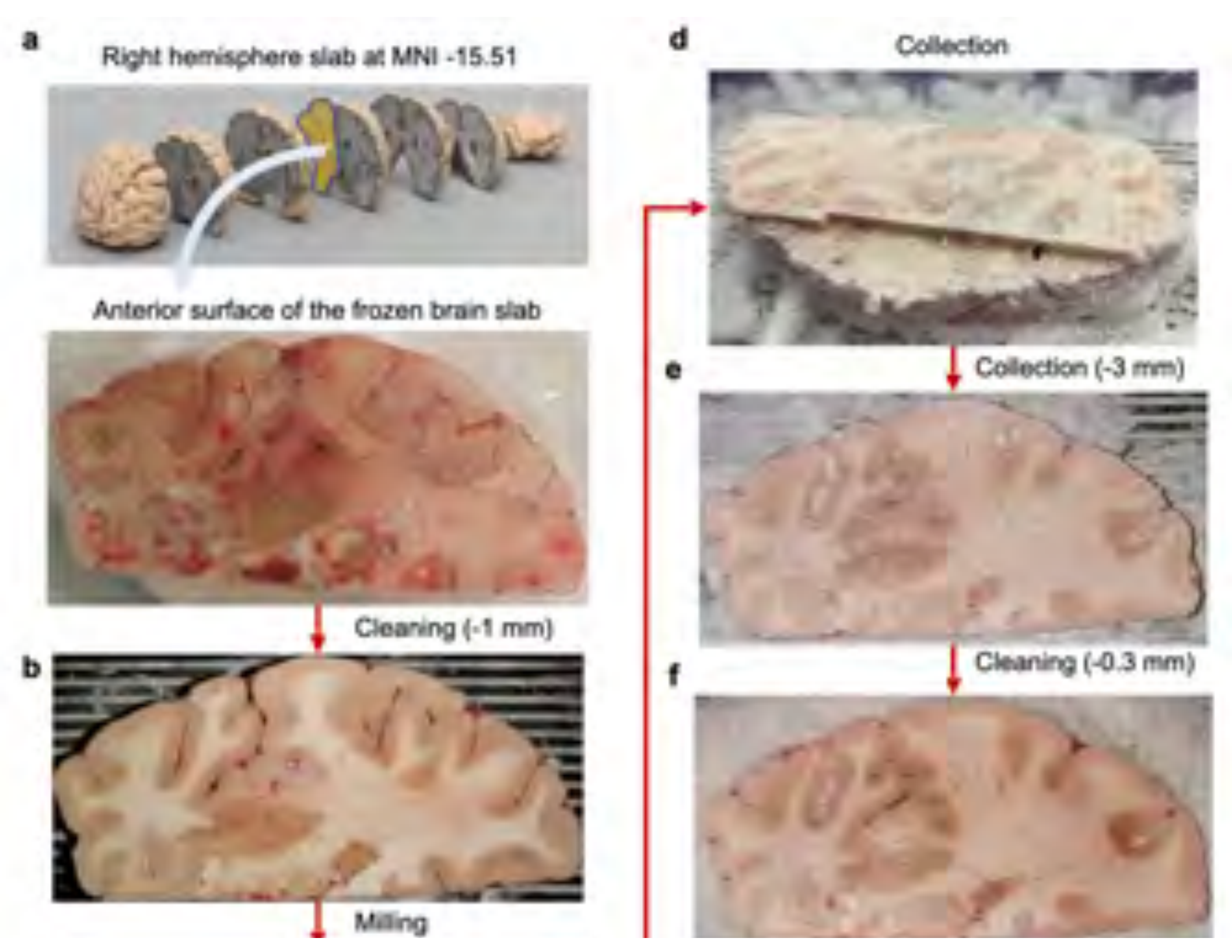


Closing the gap between organellar bioenergetic profiling and whole-brain neuroimaging modalities (fMRI, PET, CBV, DWI, etc)

Eugene Mosharov


a
Right hemisphere slab at MNI -15.51
Anterior surface of the frozen brain slab
Cleaning (-1 mm)
b
d
Collection
Collection (-3 mm)
e
Cleaning (-0.3 mm)
f


Cleaning (-1 mm)
b
Milling
c
e
Collection (-3 mm)
Cleaning (-0.3 mm)
f
Anterior
g
Cleaning (1mm)
b
Voxelization (3x3x3 mm)
Cleaning (0.3 mm)
e
f
Thin slices for histology
Posterior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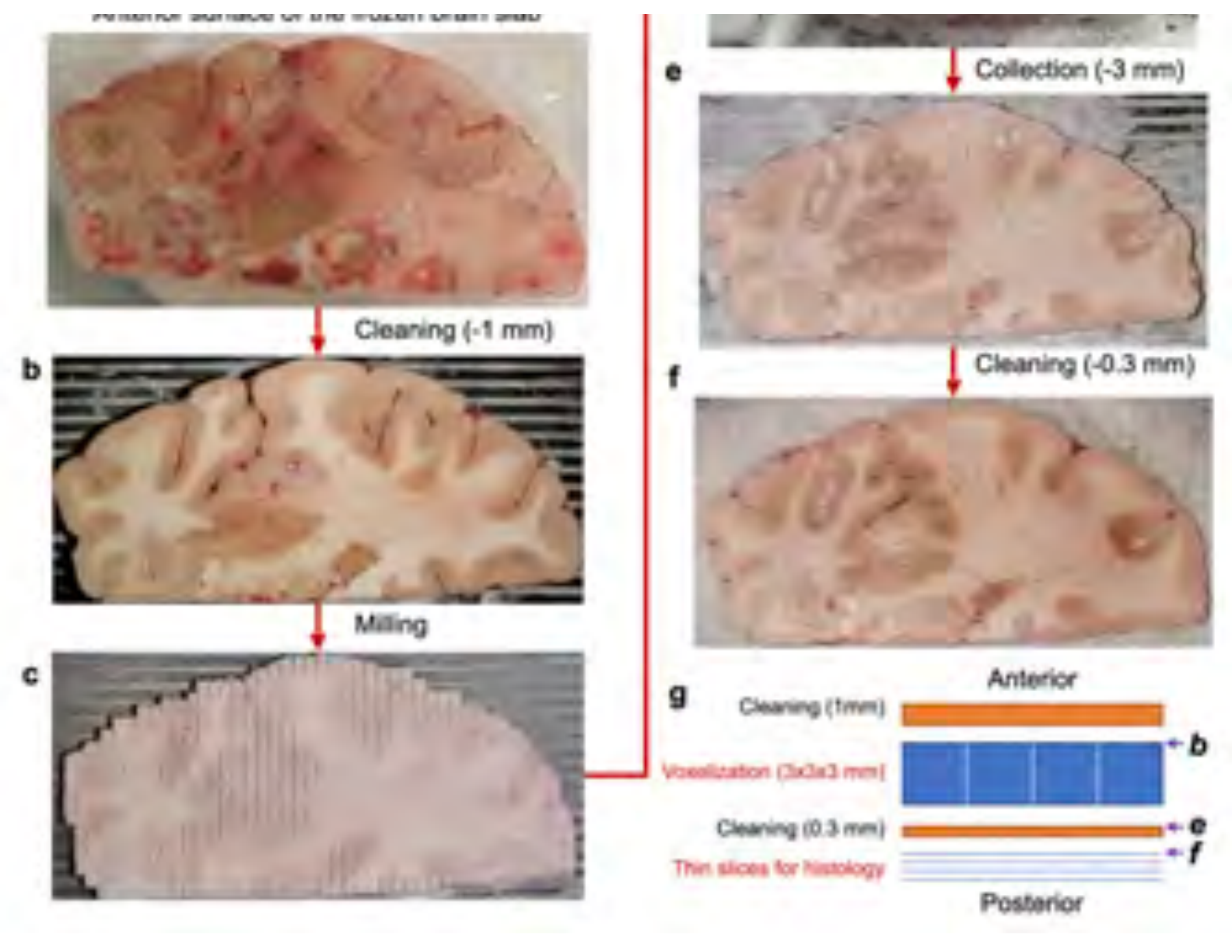
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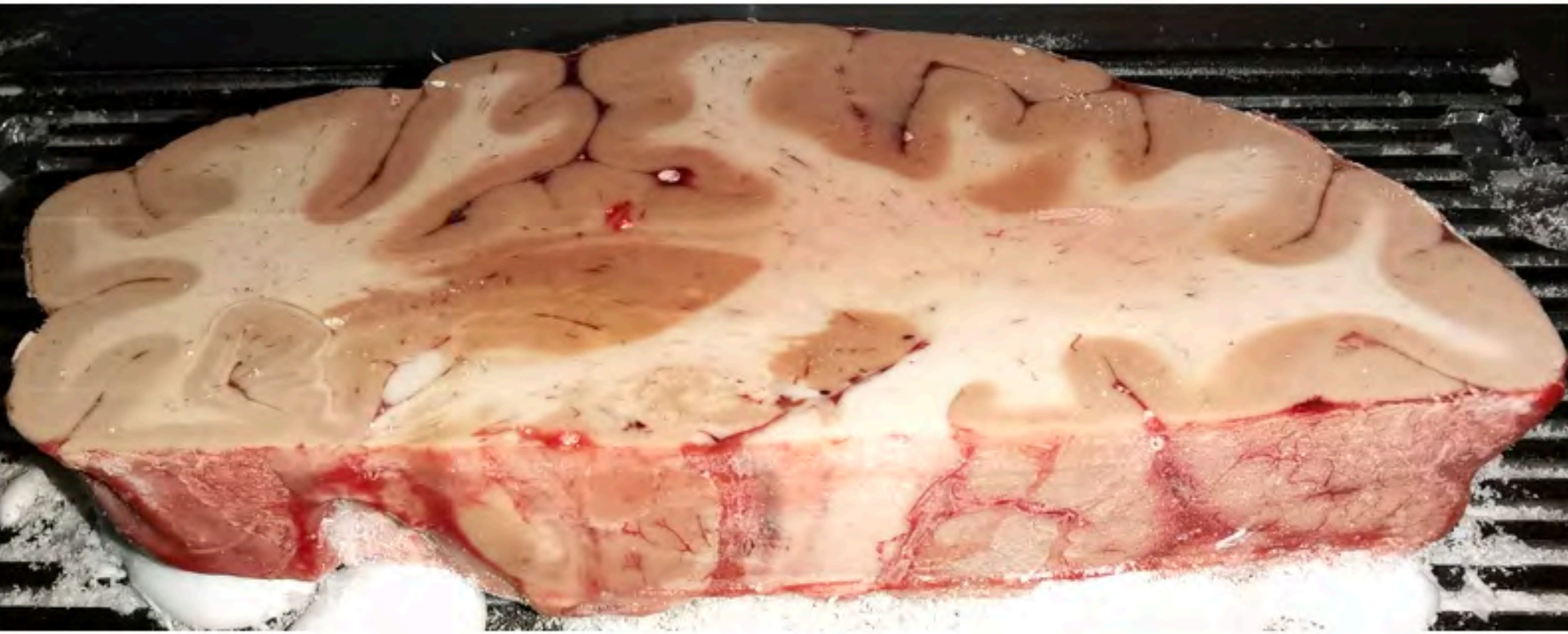
g
Anterior
Cleaning (1mm)
b
Voxelization (3x3x3 mm)
Cleaning (0.3 mm)
e
f
Thin slices for histology
Posterior

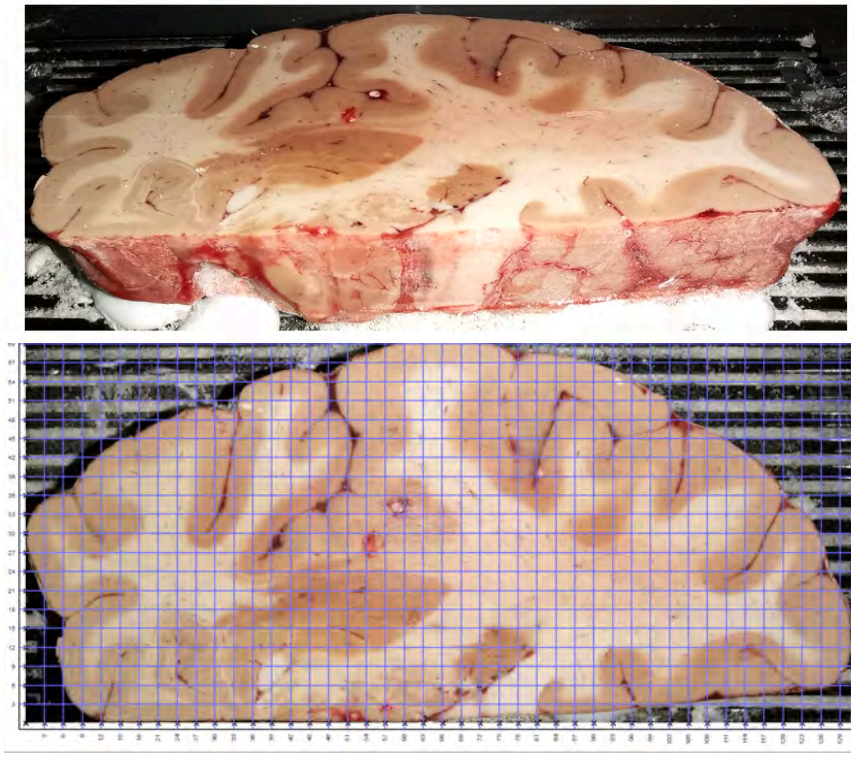
MitoBrainMap v1.0

A multi-function mitochondrial atlas of a single human coronal brain section at fMRI resolution

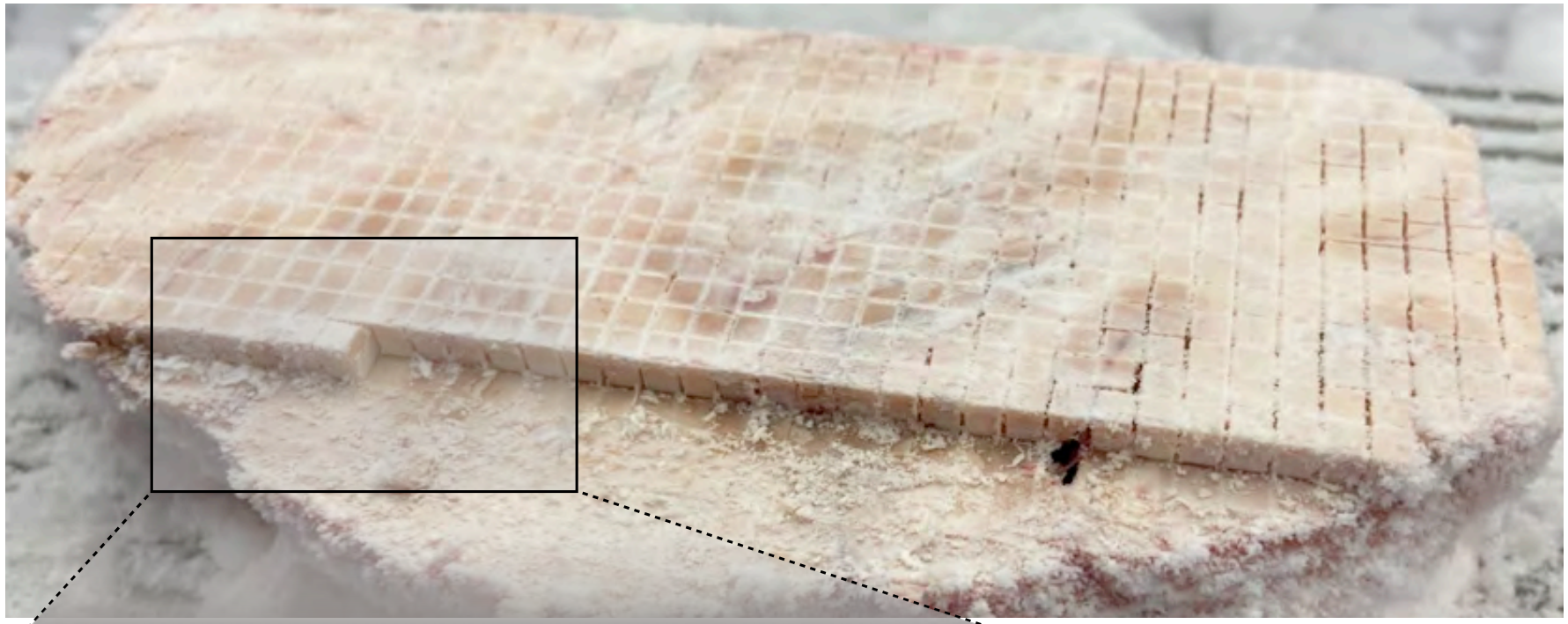


MitoBrainMap v1.0

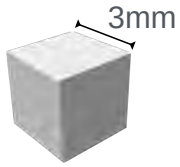
A multi-function mitochondrial atlas of a single human coronal brain section at fMRI resolution



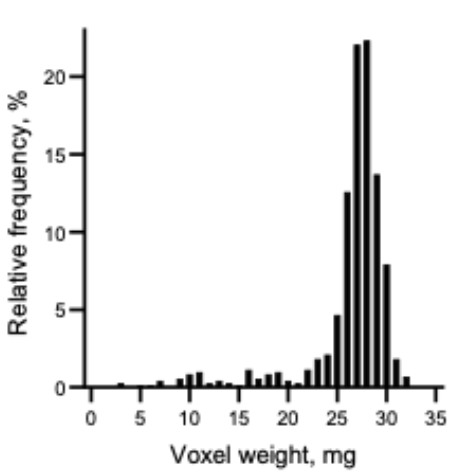
Eugene Mosharov

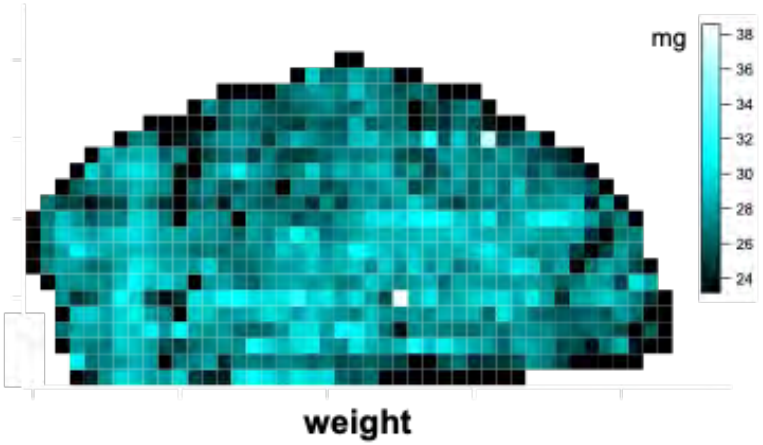


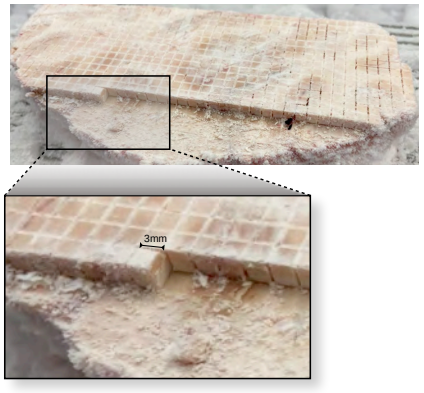
Physical *voxelization*
of the human brain
at fMRI resolution

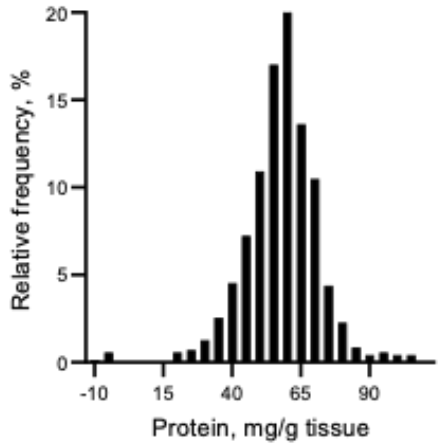


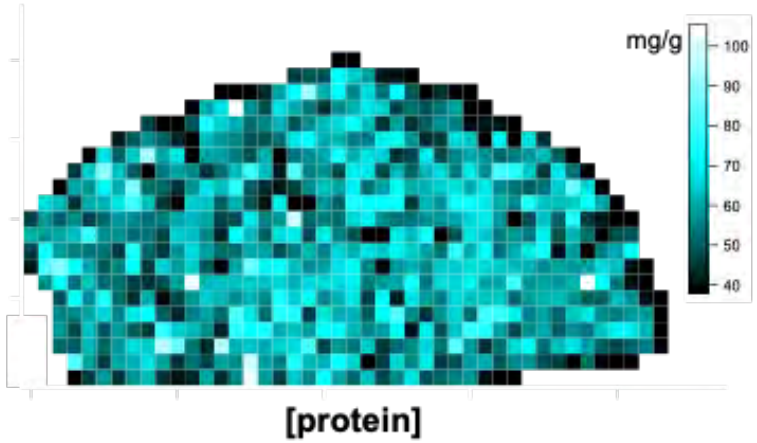
Quality control on 702 human brain voxels

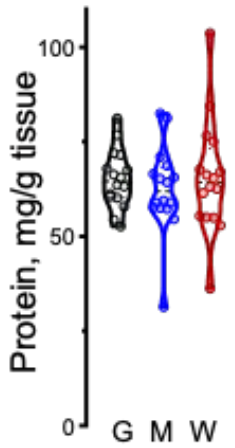






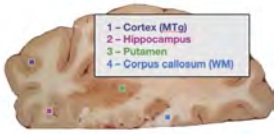






Eugene Mosharov


1 – Cortex (MTg)
2 – Hippocampus
3 – Putamen
4 – Corpus callosum (WM)


1 – Cortex (MTg)
2 – Hippocampus
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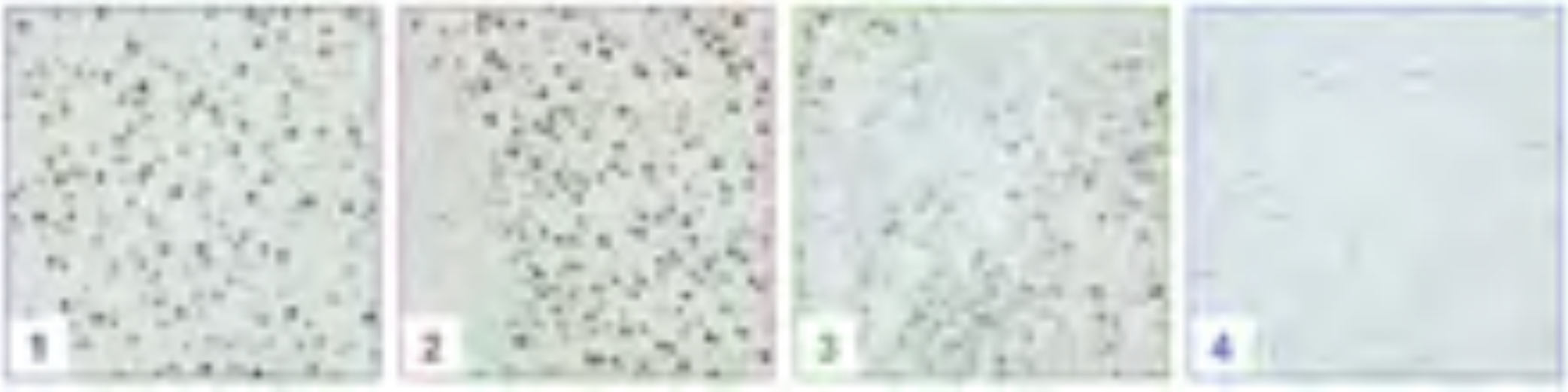
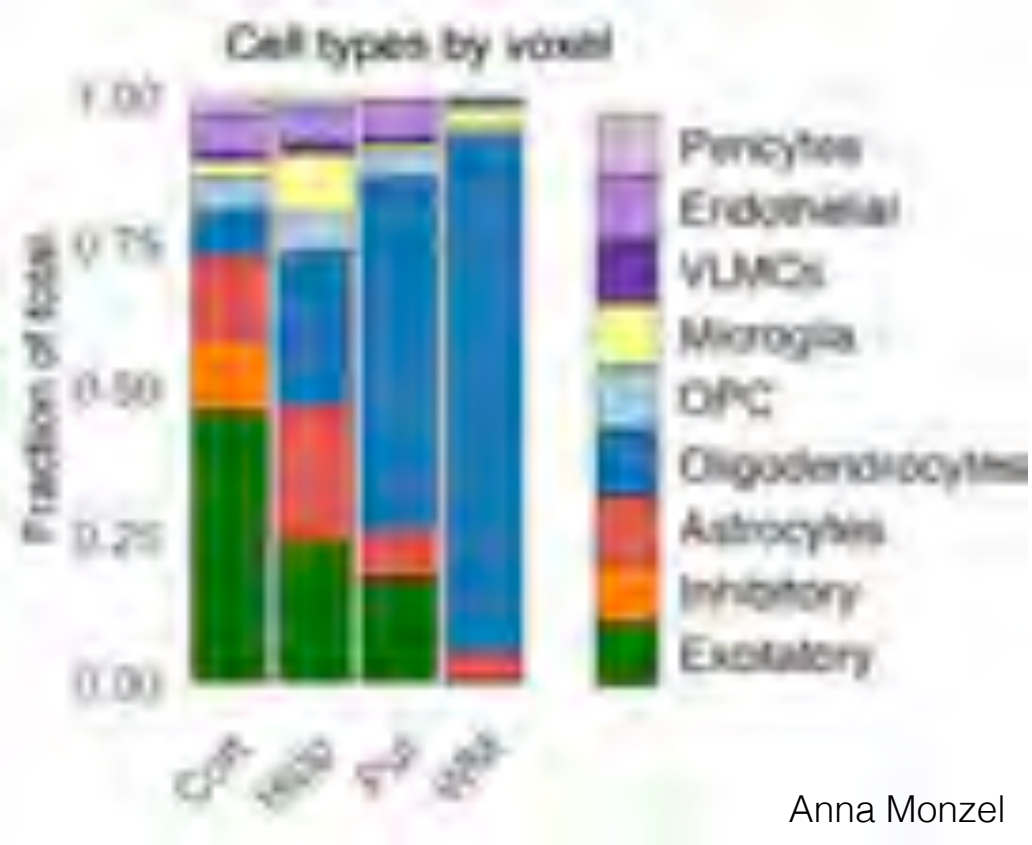
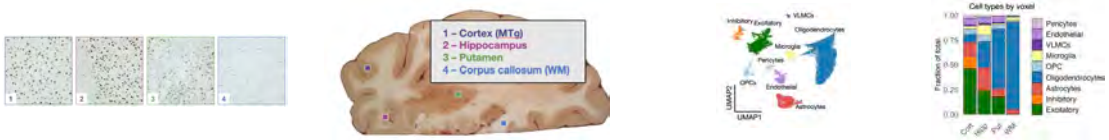
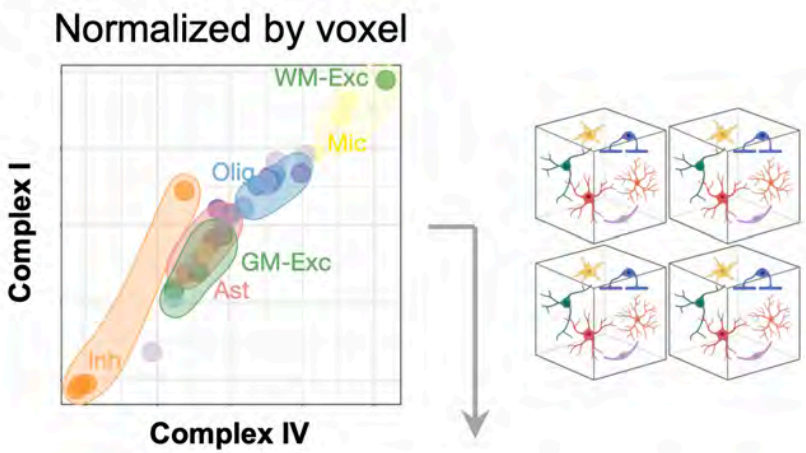

1
2
3
4

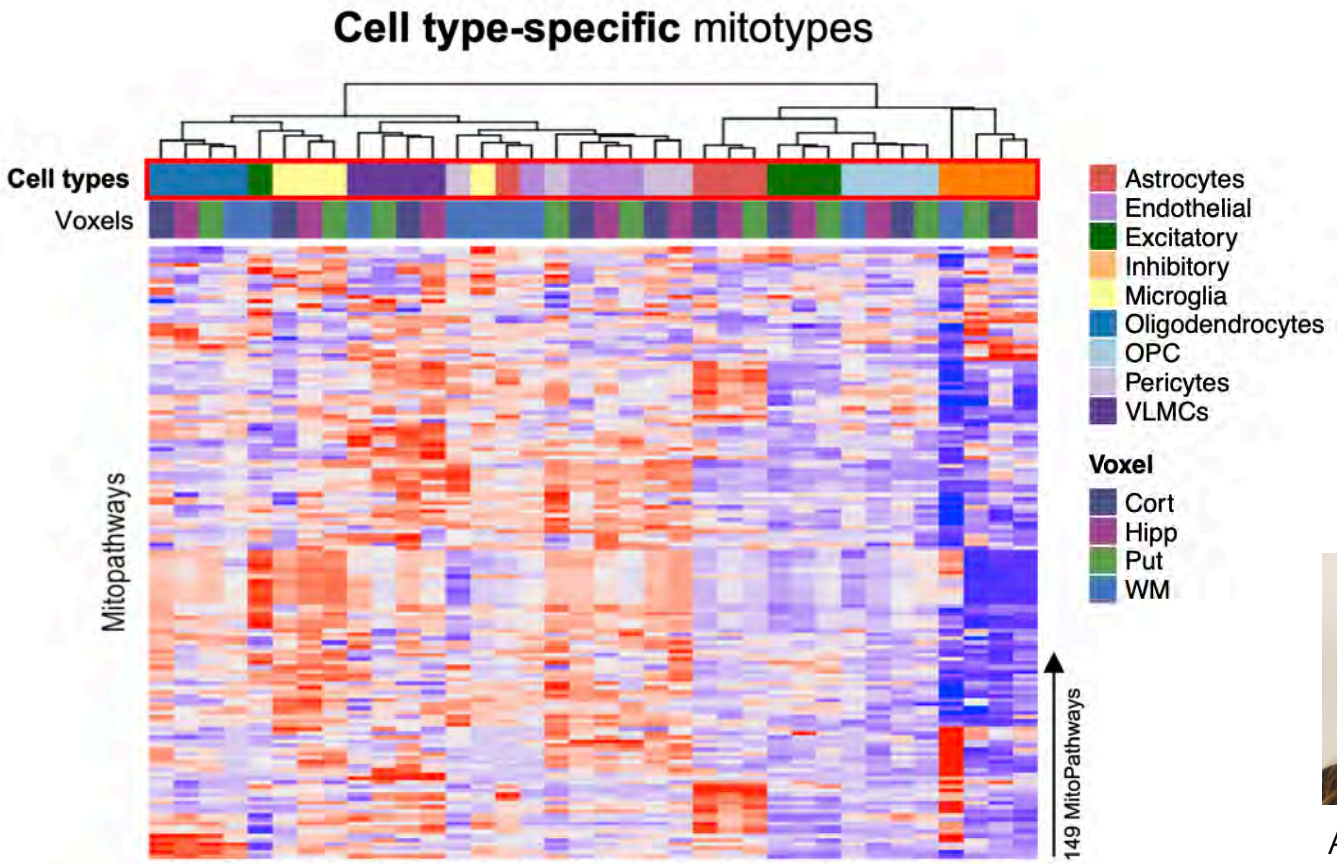

VLMCs
Inhibitory
Excitatory
Oligodendrocytes
Microglia
Pericytes
OPCs
Endothelial
Astrocytes
UMAP2
UMAP1


Cell types by voxel
Fraction of total
1.00
0.75
0.50
0.25
0.00
Pericytes
Endothelial
VLMCs
Microglia
OPC
Oligodendrocytes
Astrocytes
Inhibitory
Excitatory

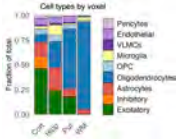
Anna Monzel

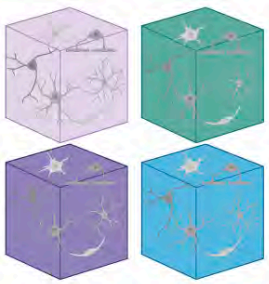

1 – Cortex (MTg)
2 – Hippocampus
3 – Putamen
4 – Corpus callosum (WM)
UMAP1
UMAP2
Cell types by voxel
Fraction of total
Cort
Hipp
Put
WM
Pericytes
Endothelial
VLMCs
Microglia
OPC
Oligodendrocytes
Astrocytes
Inhibitory
Excitatory


Normalized by voxel
WM-Exc
Mic
Olig
GM-Exc
Ast
Inh
Complex I
Complex IV

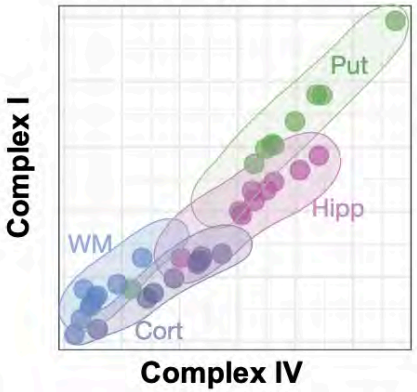

Cell type-specific mitotypes
Cell types
Voxels
Mitopathways
149 MitoPathways
Astrocytes
Endothelial
Excitatory
Inhibitory
Microglia
Oligodendrocytes
OPC
Pericytes
VLMCs
Voxel
Cort
Hipp
Put
WM

Anna Monzel

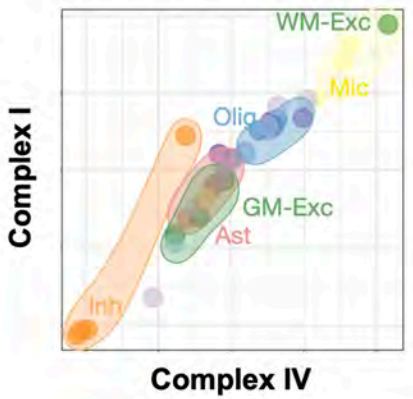


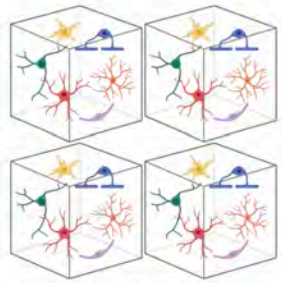


Raw expression

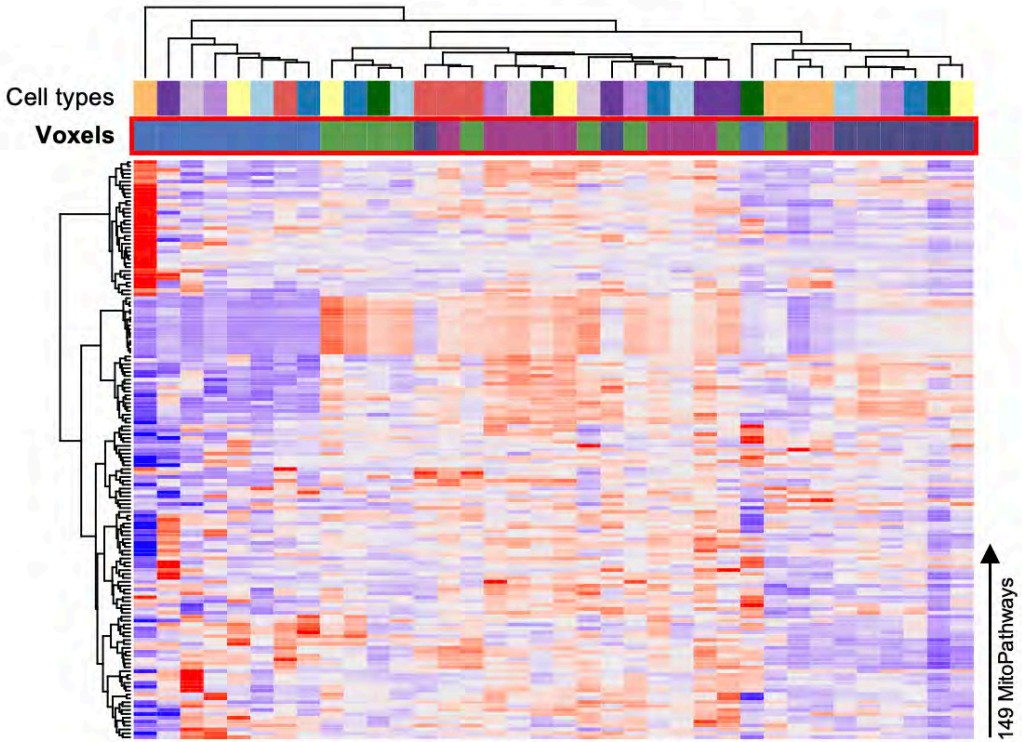


Normalized by voxel

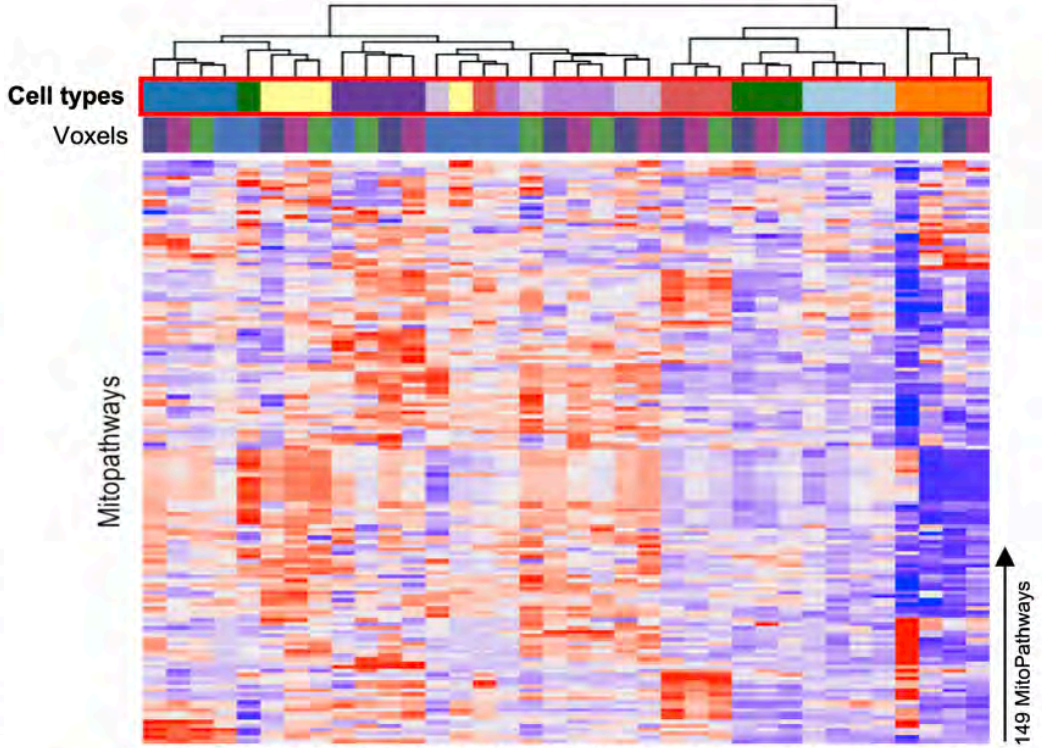




Voxels-specific mitotypes



Cell type-specific mitotypes

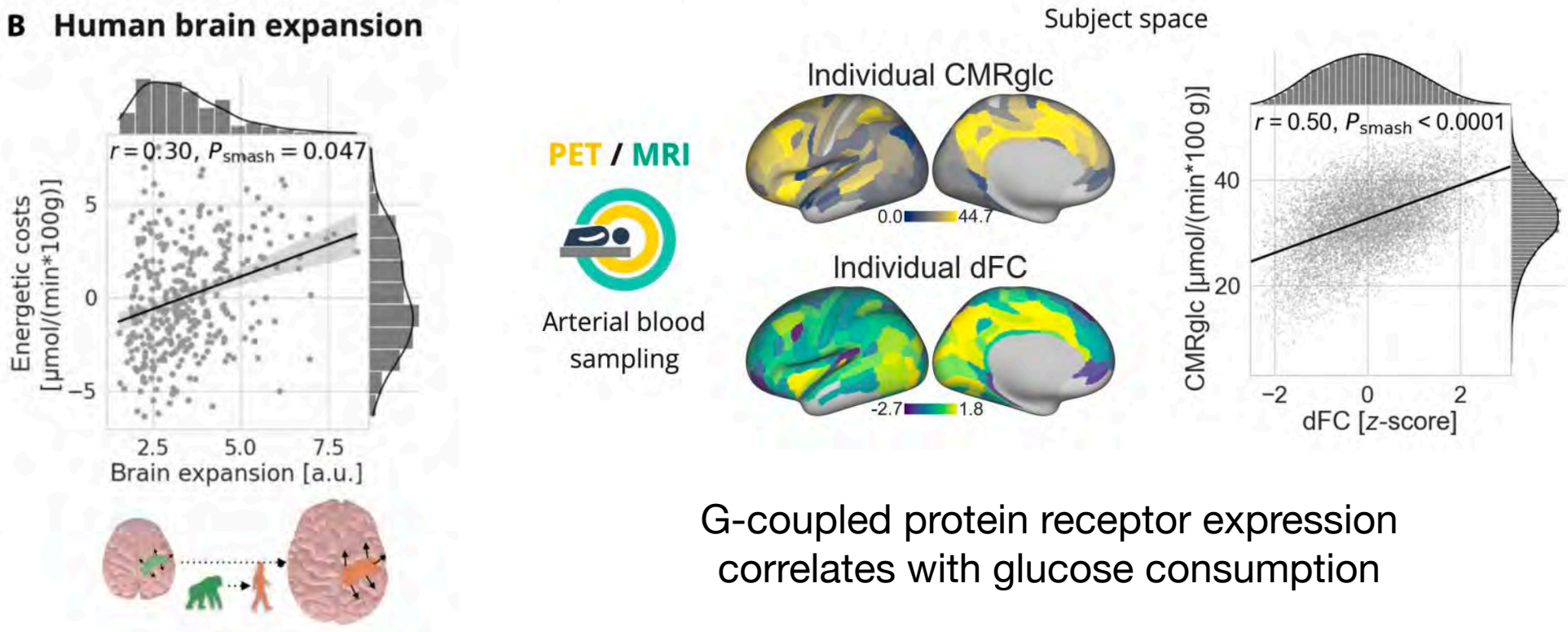


SCIENCE ADVANCES | RESEARCH ARTICLE

NEUROSCIENCE

An energy costly architecture of neuromodulators for human brain evolution and cognition

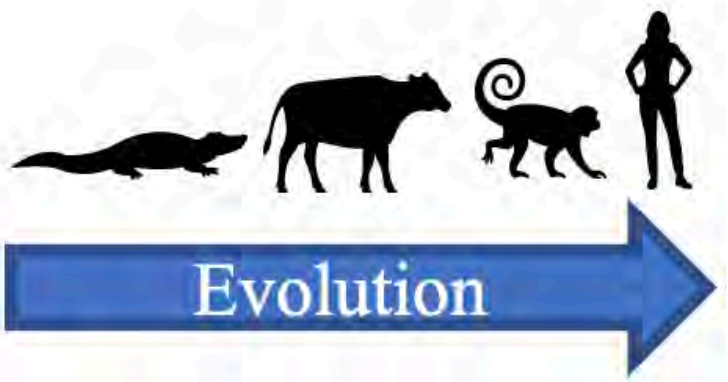
Gabriel Castrillon[1,2,3], Samira Epp[1,4], Antonia Bose[1,4], Laura Fraticelli[1,4], André Hechler[1,4], Roman Belenya[1,4], Andreas Ranft[5], Igor Yakushev[6], Lukas Utz[1], Lalith Sundar[7], Josef P Rauschecker[8,9], Christine Preibisch[1,10], Katarzyna Kurcyus[1], Valentin Riedl[1,3]*

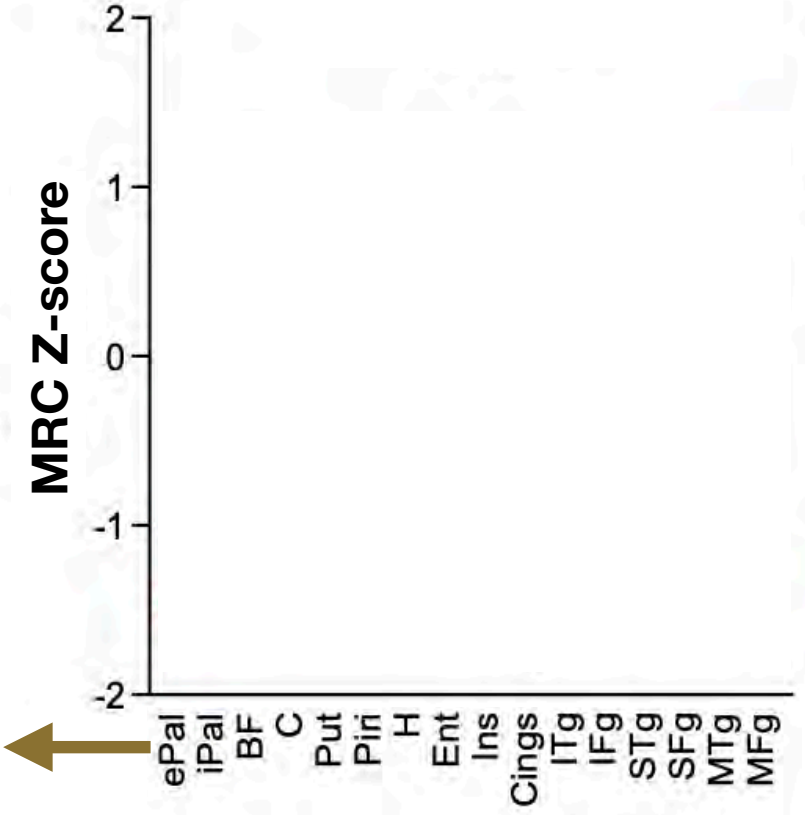


G-coupled protein receptor expression correlates with glucose consumption

Castrillon et al. *Sci Adv* 2023

Evolutionary correlate of mitochondrial OxPhos specialization

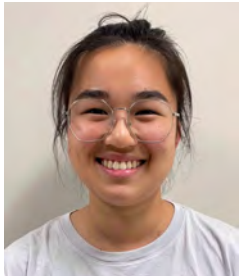




Most **"ancient"** ← → Most **"recent"**

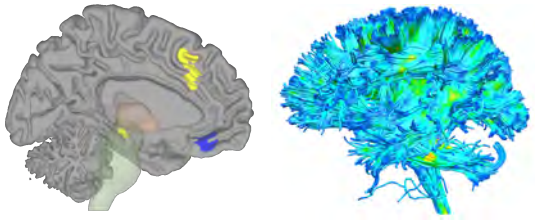
MITO BRAIN

ROSMAP



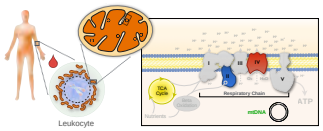
Cynthia Liu

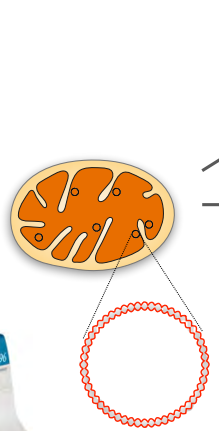
Mitochondrial **S**tress, **B**rain **I**maging, and **E**pigenetics — **MiSBIE**

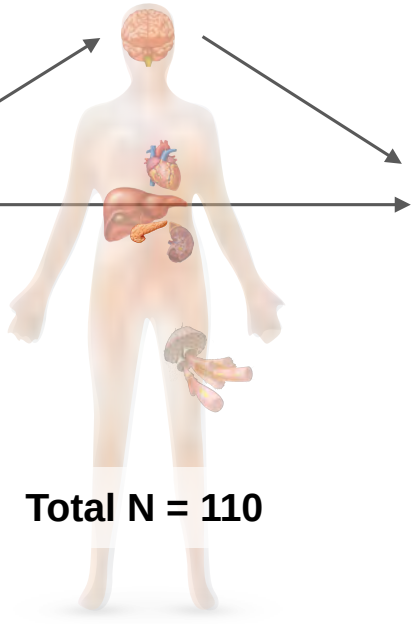


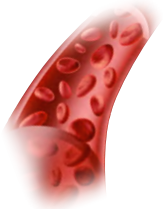
Brain structure and function
Neuropsychological function

mtDNA heteroplasmy
Mitochondrial OxPhos
Lymphocytes, Monocytes,
Neutrophils, Platelets





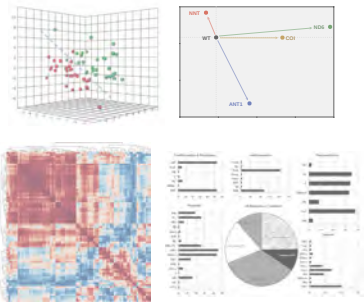




Total N = 110

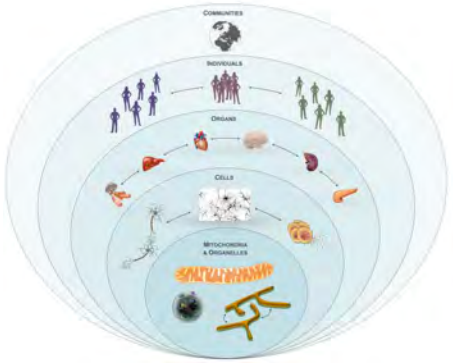
Disease biomarkers
Stress reactivity
Energy expenditure

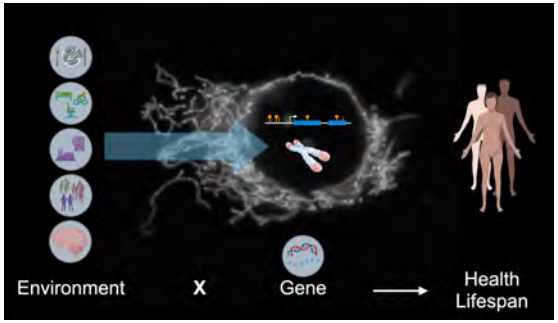
e.g., GDF15, Lactate,
Pyruvate, Alanine, etc.

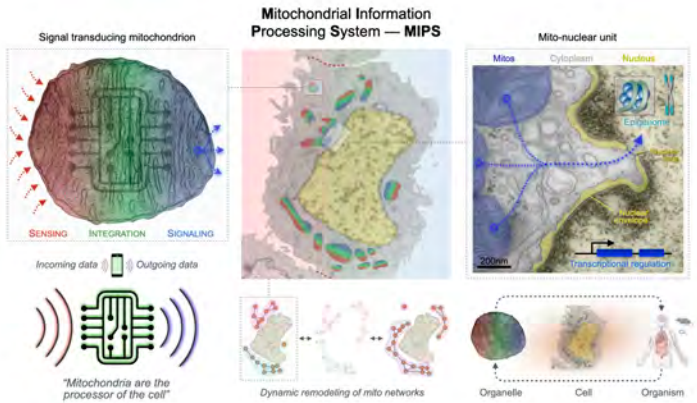


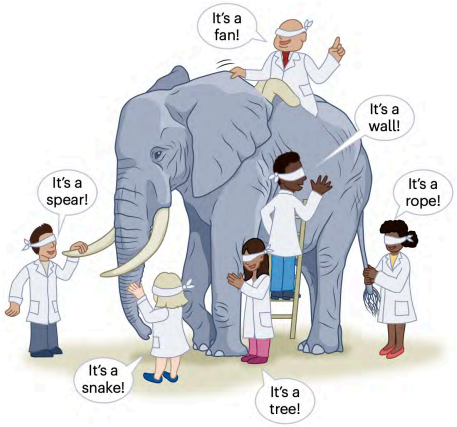
- **Healthy controls** (n = 70)
- **mtDNA defects**
  - **3243A>G (group A)** (n = 20)
  - **3243A>G (group B)** (n = 5)
  - **Single deletion** (n = 15)

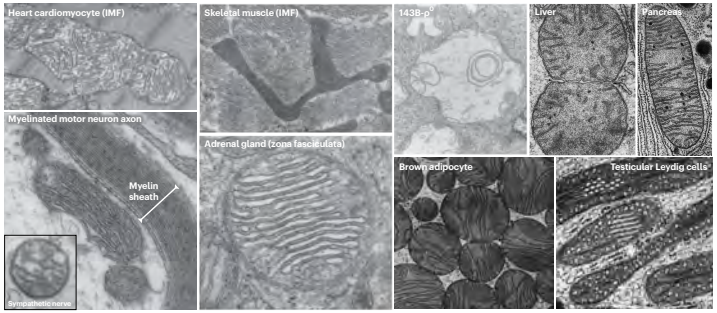
MITOCHONDRIA STRESS BRAIN EPIGENETICS MiSBIE

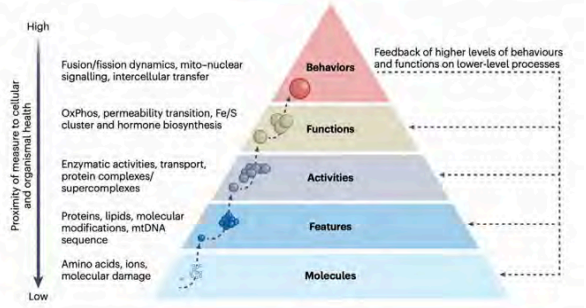


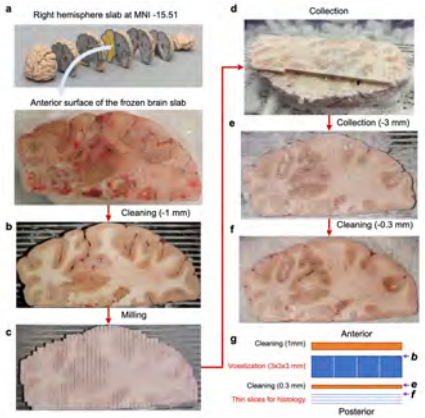


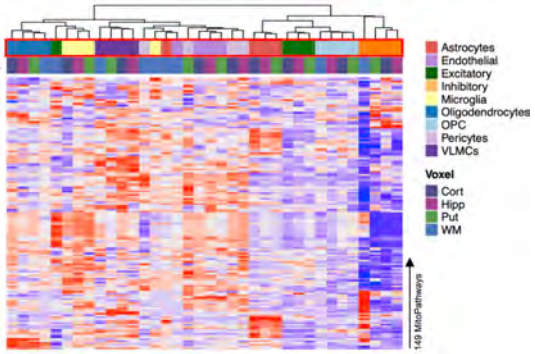


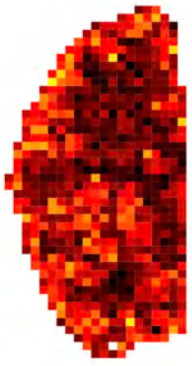


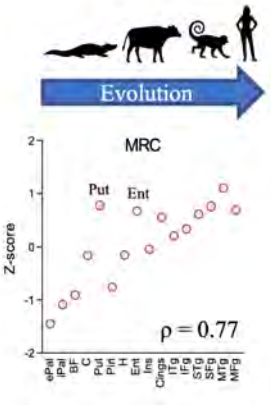




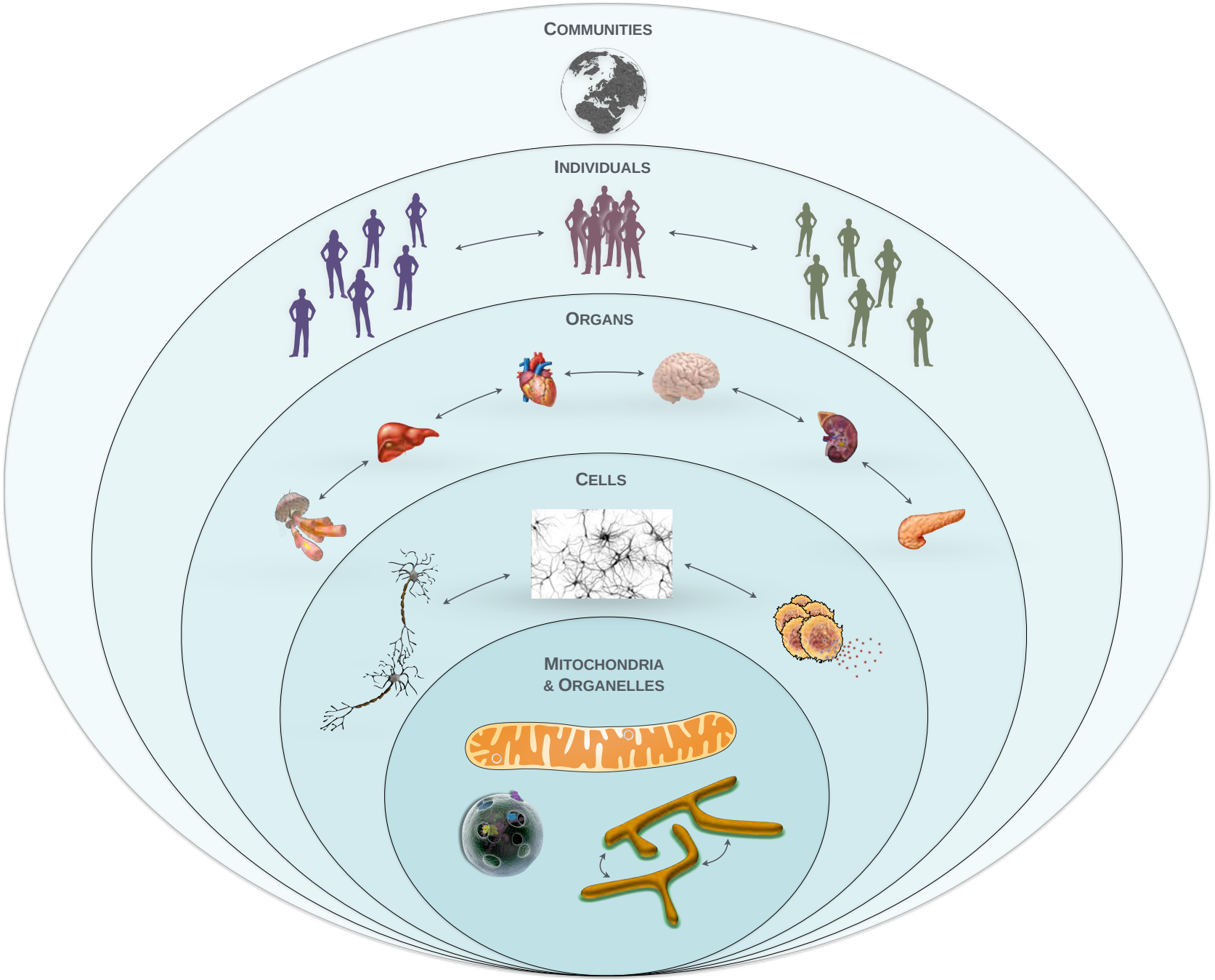








MITO BRAIN
ROSMAP



Picard and Sandi. *Neurosci Biobehav Rev* (2021)

Mitochondrial PsychoBiology Lab

Linking molecular processes within mitochondria with the human experience

OUR RESEARCH

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NIA Intramural

Dan Belsky
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CUIMC Mailman & Aging Center

BASZUCKI BRAIN RESEARCH FUND
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Downloadable presentation slides

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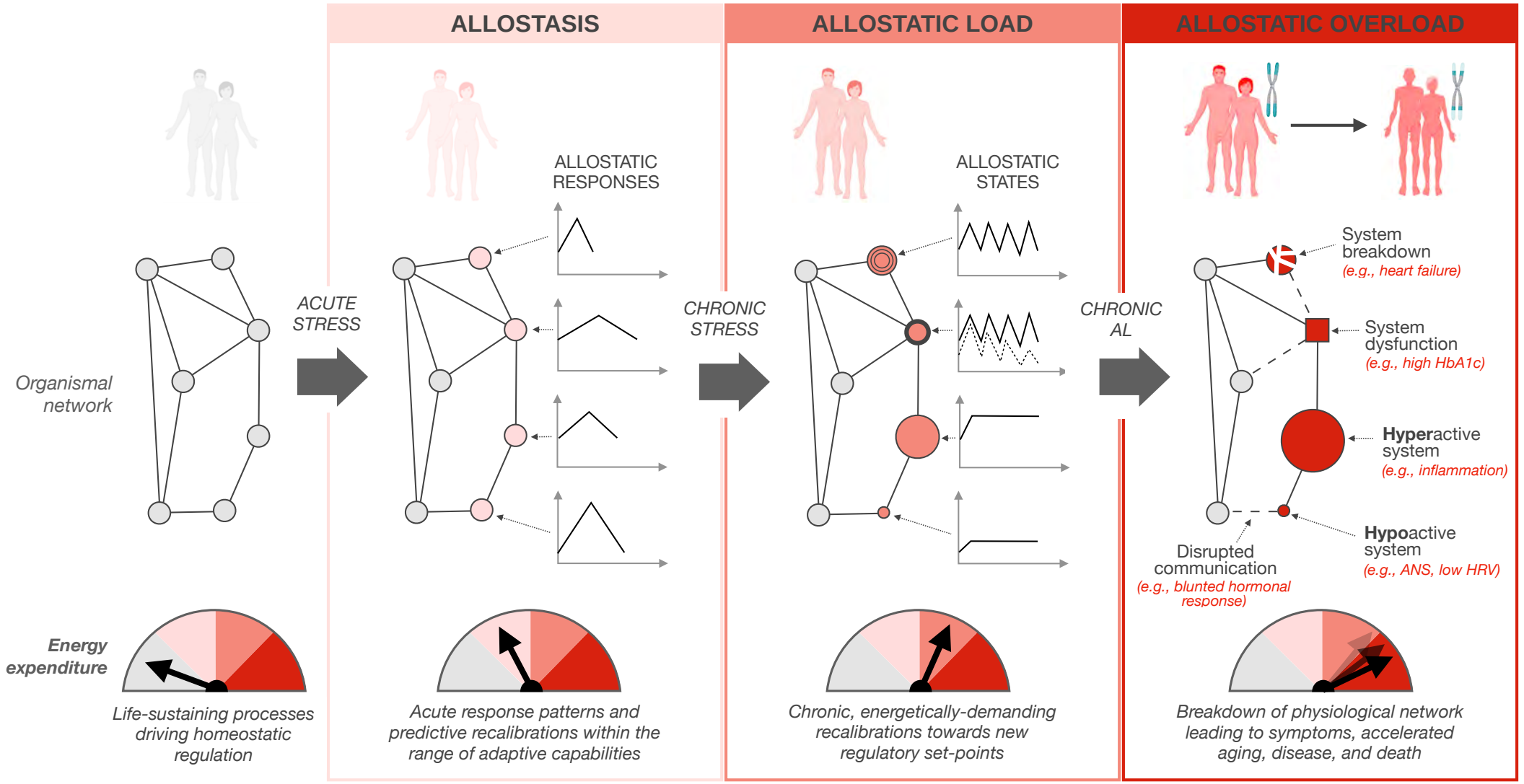
IRVING MEDICAL CENTER

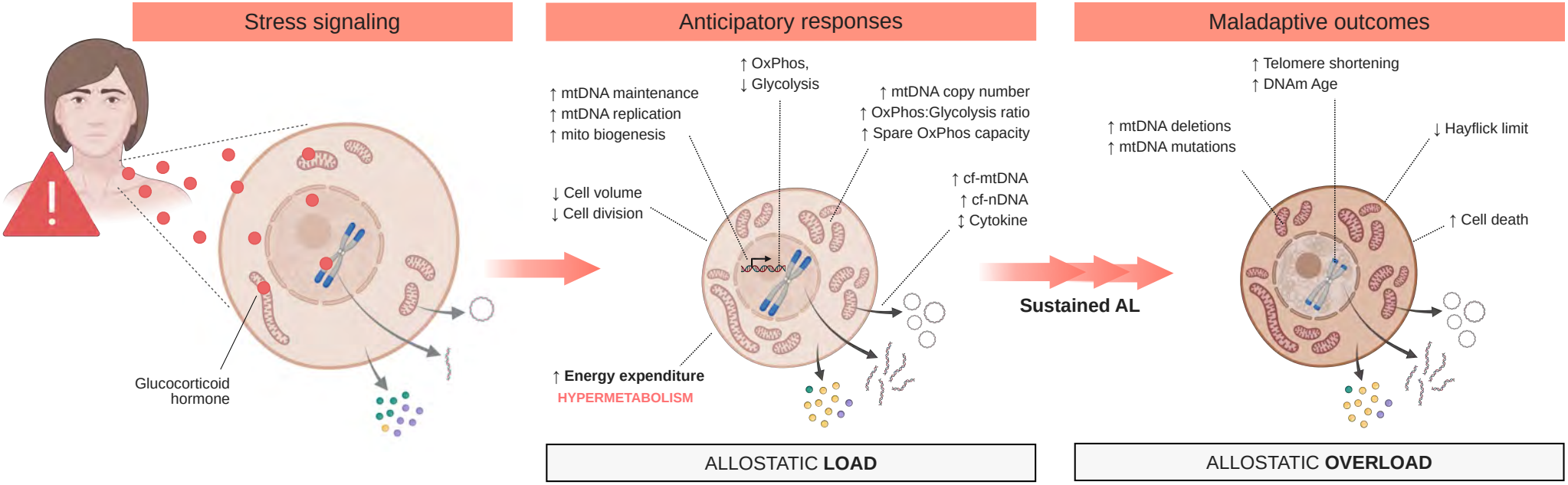
MITOCHONDRIAL PSYCHOBIOLOGY LAB

MITOCHONDRIA STRESS BRAIN EPIGENETICS iSBIE

Energetic Model of Allostatic Load (EMAL)



Bobba-Alves, Juster, Picard. *Psychoneuroendocrinol* 2022

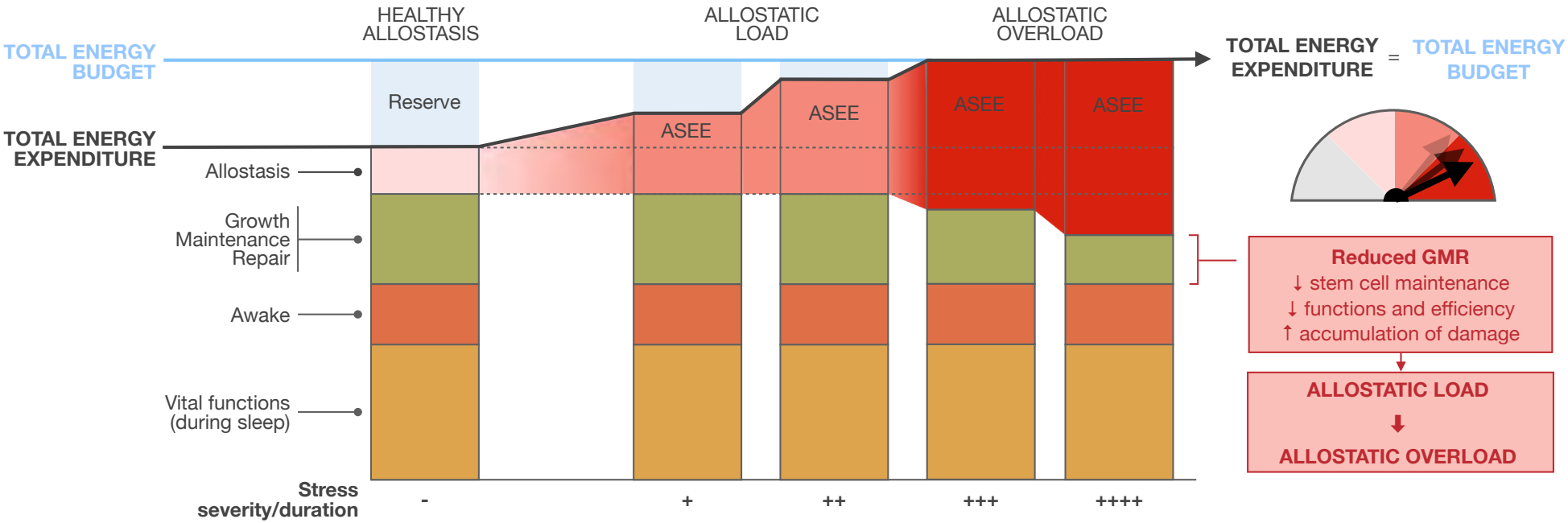


Glucocorticoid signaling increases energy expenditure by **60%**

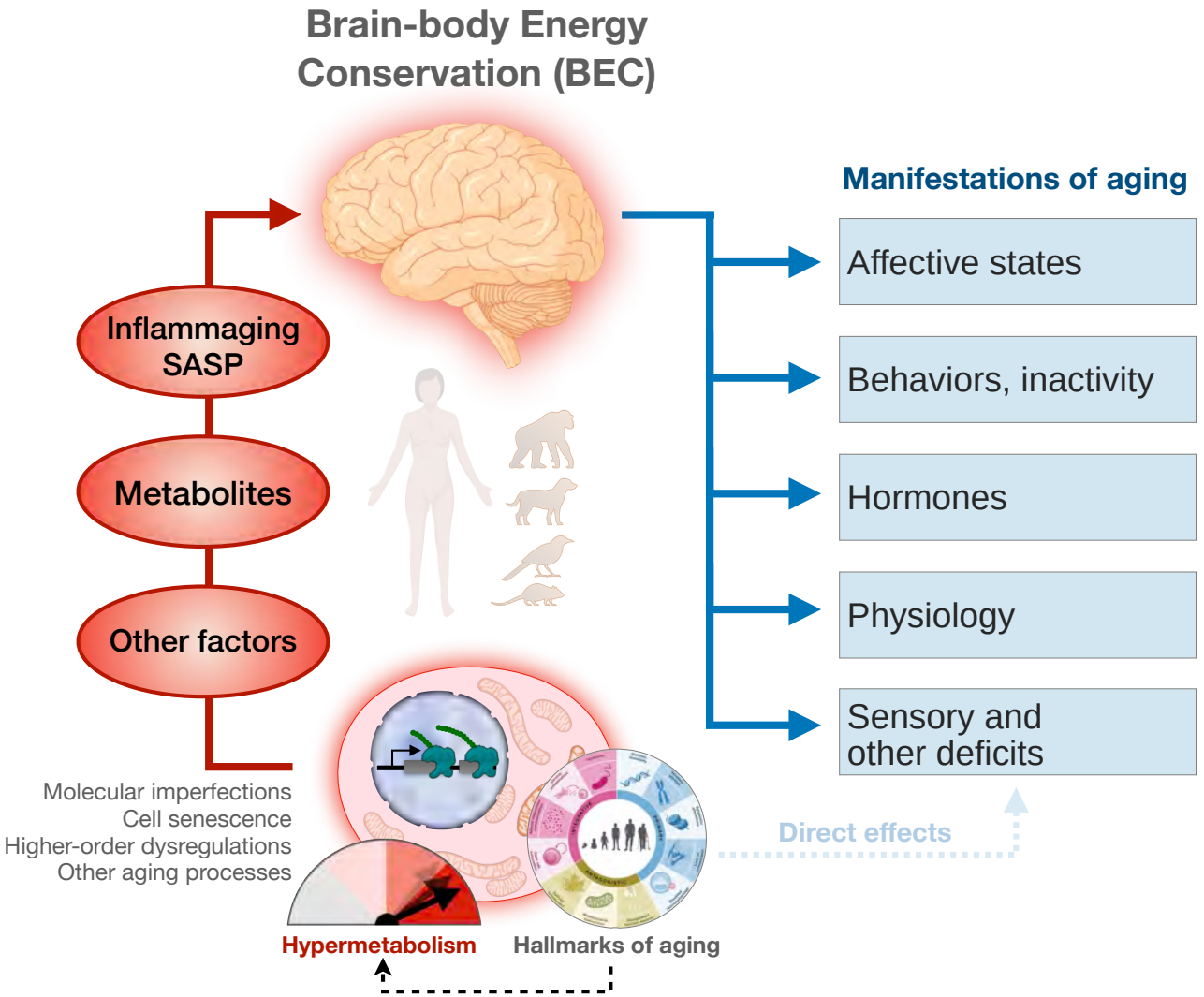
And accelerates cellular aging by **10-40%**

Bobba-Alves et al. *PNEC* 2023

Model: Aging and lifespan determined by the partitioning of limited energetic resources



Bobba-Alves et al. The energetic cost of allostasis and allostatic load. *Psychoneuroendocrinol* 2022



Shaulson, Cohen, Picard. *Preprint* (osf.io/zuey2)