



Photo by Kathy West, courtesy of the California National Primate Research Center.

NONHUMAN PRIMATE 1 2024 RESOURCES

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ORIP advances the NIH mission by providing the foundation for scientific innovation via research resources, including animal models for human diseases, cutting-edge scientific instrumentation, construction and modernization of research facilities, and research training opportunities for veterinary scientists. Through continued engagement with NIH institutes, centers, and offices and the biomedical research community, ORIP empowers and expands existing programs and develops new initiatives to support NIH research at the forefront of scientific progress.





National Institutes of Health Office of Research Infrastructure Programs

OVERVIEW

The Division of Comparative Medicine (DCM) within the Office of Research Infrastructure Programs (ORIP), Division of Program Coordination, Planning, and Strategic Initiatives (DPCPSI), Office of the Director, National Institutes of Health (NIH), advances biomedical research by supporting research resources, such as those that provide animal models for human disease.

Because of their genetic, anatomical, physiological, and behavioral similarities to humans, nonhuman primates (NHPs) are one of the best models for human disease research when studies in humans are not feasible. Among other important medical advances, NHPs have played key roles in the understanding and treatment of a variety of infectious diseases, such as AIDS, tuberculosis, Zika virus disease and congenital Zika syndrome, Ebola, and, recently, COVID-19. Additionally, NHPs have been critical in advancing therapeutics for type 2 diabetes and other metabolic disorders, treatment of glioblastoma (brain cancer), deep brain stimulation to treat Parkinson's disease, neuroprosthetics (including decoding of brain waves for brain–machine interfaces), pain management interventions, and organ transplantation. NHPs help determine the safety and efficacy of vaccines, devices, and therapies before they are used in humans. The COVID-19 pandemic demonstrated the pivotal role of NHPs in developing medical countermeasures for SARS-CoV-2, the virus that causes COVID-19; this included understanding infection progression and pathophysiology, preclinical development of vaccine candidates and therapeutics, and development of a variety of SARS-CoV-2specific diagnostic assays.

Costs related to stringent breeding and housing requirements limit access to NHPs throughout the biomedical research community. To mitigate these issues, ORIP's DCM supports multiple NHP facilities and research-related resources that are available to the community of NIH-funded researchers. These NHP resources support biomedical research spanning scientific disciplines, with studies supported across almost all NIH institutes, centers, and offices.

NONHUMAN PRIMATE RESOURCES



National Primate Research Centers

The <u>National Primate Research</u> <u>Centers (NPRCs)</u> are a national network of seven Centers that

increase access to and promote sharing of valuable NHPrelated resources among biomedical researchers. Additionally, the NPRCs advance the missions of NIH institutes, centers, and offices by providing the animals, facilities, expertise, and resources required by investigators in disease areas. ORIP's DCM funds NPRCs located in California, Georgia, Louisiana, Oregon, Texas, Washington, and Wisconsin. Collectively, the NPRCs maintain breeding colonies for rhesus, pigtail, and Japanese macaques; common marmosets; olive baboons; and titi monkeys. Each Center provides expertise on the use of various NHP species as models for human disease in specific research projects. Each provides a variety of services both individually and through inter-NPRC collaborations. The NPRCs provide services for research funded by NIH, other federal agencies, nonprofit foundations, and the private sector. Additionally, the program offers a Pilot Research Program that supports new investigators snd exploratory research, as well as a Visiting Scientist Program that offers advanced research training. The NPRCs have scientific programs addressing major research fields, such as infectious diseases, aging, cardiovascular disease, diabetes and metabolic disorders, neuroscience, pediatrics, regenerative medicine, reproductive health, and women's health. For detailed information on NPRC capabilities and programs, visit NPRCresearch.org. Recent advances by the NPRCs can be viewed at nprc.org.





NHP COVID-19 research being conducted in an Animal Biosafety Level 3 laboratory. Photo courtesy of the Southwest National Primate Research Center.

Specific-Pathogen-Free Macaque Colonies

Macaque monkeys are premier research models for HIV/AIDS. For example, macaques infected with the simian immunodeficiency virus (SIV)—the NHP analogue of HIV are used to address basic research questions about viral infection routes, acute phases of infection, and latent viral reservoirs because these cannot be studied directly in humans. Likewise, SIV-infected macaques serve as models for developing HIV vaccines, infection prevention devices, new therapeutics, microbicides, and cure strategies prior to first-in-human trials.

The presence of specific viral pathogens in experimental animals can confound the results of HIV/AIDS-related investigations or pose a health risk to staff. Therefore, a consortium of colonies was developed to provide specificpathogen-free (SPF) macaques for AIDS research that are negative for SIV, type D simian retrovirus, simian T-cell lymphotropic virus, and herpes B virus. Additionally, SPF macaques are characterized for major histocompatibility complex class I alleles, which are known to be associated with SIV viral load and rate of disease progression. ORIP supports SPF rhesus macaque colonies at the California, Emory, Oregon, Southwest, and Tulane NPRCs, as well as the Caribbean Primate Research Center in Puerto Rico. ORIP also supports SPF pigtail macaque colonies at the Washington NPRC and the Johns Hopkins University School of Medicine. Visit the <u>ORIP website</u> for more details on these critical SPF macaque resources.

Other Nonhuman Primate Research Resources

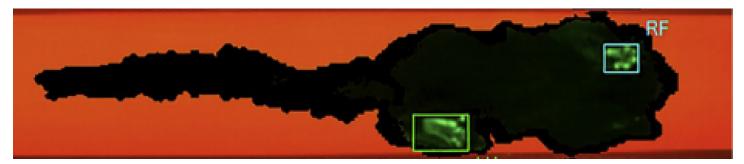
Baboon Research Resources: Relative to most other NHP models, baboons share the greatest similarity to humans in terms of their larger size, year-round breeding, and placental biology. Because of similarities between the baboon and human immune systems, baboons are critical for vaccine development, xenotransplantation, and studies of infectious disease and bacterial sepsis. The baboon colony at MD Anderson Cancer Center's Michale E. Keeling Center for Comparative Medicine and Research (KCCMR) is maintained free of infection from an extensive list of at least 18 viruses, bacteria, and parasites and is a valuable research resource for studies requiring the unique similarities of baboons to humans without the complicating influences of coinfections. A baboon colony also exists at the Southwest NPRC.

Squirrel Monkey Breeding and Research Resource:

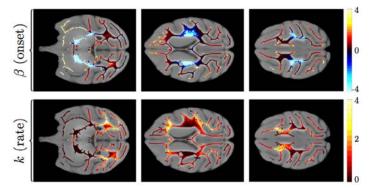
Housed at KCCMR, this is the only national <u>squirrel</u> <u>monkey</u> breeding and research resource available for biomedical research and one of the few NIH-supported national research resources that specialize in a New World (neotropical) primate species. Squirrel monkeys are valuable for neuroscience research because of their small size and similarities to humans in brain structure, which makes them superior neuroscience models compared to small nonprimate mammals, such as rodents. Squirrel monkeys are used in neuroscience, vision, and hearing research; in studies of infectious diseases (malaria vaccine, polyoma virus disease, Zika virus); as a model for sporadic cerebral amyloid angiopathy; and in research on drug addiction and its behavioral and physiological consequences.

Caribbean Primate Research Center Program:

The <u>Caribbean Primate Research Center (CPRC) Program</u> maintains conventional and SPF macaque colonies. Additionally, the CPRC maintains a free-ranging colony of rhesus macaques of purely Indian origin that was established over 80 years ago from a substantial founder population. This colony exhibits the lowest levels of genetic admixture with non-Indian-origin rhesus macaques among all rhesus monkey colonies surveyed in the United States, providing a unique resource for research in a naturalistic



Data on gait parameters as a common marmoset in the Wisconsin National Primate Research Center's Preclinical Parkinson's Research Program walks through a Noldus CatWalk XT10.6 (apparatus not shown).



Postnatal brain structural maturation in infant rhesus macaques during the first 18 months of age. Top: White matter onset intensity at birth (β). Bottom: Median rate (k) of normalized white matter intensity change per day. Image courtesy of the Emory National Primate Research Center and collaborators, Drs. M. Styner and M. Niethammer.

setting. The CPRC provides resources to researchers at other U.S. institutions, as well as collaborations onsite, and it has active programs in virology (especially SIV and West Nile, dengue, and Zika viruses), genetics, diabetes, parasitology, behavior, cognition, and anatomy.

Vervet Research Colony: Vervets, or African green monkeys (AGMs), are critical research models owing to their similarities to humans in reproductive biology, development of cardiovascular disease and type 2 diabetes on a Western diet, and growth of amyloid plaques with age. The <u>Vervet</u> <u>Research Colony (VRC)</u> at Wake Forest School of Medicine maintains a multi-generational, pathogen-free, genotyped colony of Caribbean-origin AGMs. The colony consists of individuals ranging in age from newborns to geriatric animals over 27 years old. VRC animals, biospecimens, and data have contributed to research on diabetes, obesity, cardiovascular diseases, Alzheimer's disease, microbiome influences, metabolomics, and neuroscience. Additionally, VRC animals have supported vaccine research for SIV, neonatal influenza, respiratory syncytial virus, dengue virus, and SARS-CoV-2.

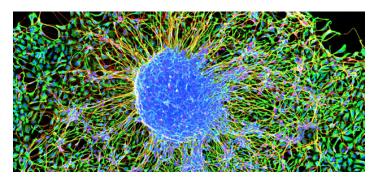
NHP Antibody Resources: The Nonhuman Primate Reagent Resource (NHPRR) and the Neotropical Primate Reagent Resource are located at Mass Biologics, a business unit of the University of Massachusetts Medical School. With ORIP support, the NHPRR develops, manufactures, and distributes immune cell–depleting antibody reagents to optimize



Pyramidal cell from the prefrontal cortex of a rhesus macaque. The NPRCs offer a wide variety of resources for research with nonhuman primates, including advanced microscopy. Photo courtesy of John Morrison of the University of California, Davis.

the usefulness of Old World NHPs (e.g., rhesus and pigtail macaques) in biomedical research. These reagents support research on HIV and other infectious diseases, transplantation, cancer, and gene therapy. The Neotropical Primate Reagent Resource characterizes the immunoglobulin repertoire and antibody responses of New World NHPs (e.g., marmosets and squirrel monkeys) and uses this information to engineer species-matched lymphocyte-depleting and diagnostic antibodies for New World monkeys. Investigators may make inquiries of, and request reagents from, both resources via the <u>NHPRR website</u>.

The New World Monkey Immunoreagent Resource, located at Trinity University in San Antonio, Texas, develops mouse monoclonal antibodies to be used in novel immunoassays specific for biomarkers of inflammation and metabolic hormones in marmosets, squirrel monkeys, and owl monkeys. Immunoreagents for such biomarkers are useful in studies of aging, infectious diseases, neurodegenerative diseases, diabetes, metabolic syndrome, and obesity, all of which are being modeled in New World monkeys.



Common marmoset-derived embryonic stem cells differentiating into neurons, courtesy of Marina Emborg's laboratory at the Wisconsin National Primate Research Center. A neurosphere was stained to visualize nuclei (blue) and immature neural progenitors (green) transitioning to neurons (red). Image by Scott Vermilyea, Ph.D.

NHP Centers of the Somatic Cell Genome Editing

Program: The NIH Common Fund's <u>Somatic Cell Genome</u> <u>Editing (SCGE) Program</u> includes approaches for development and testing of gene editing in NHPs to improve their efficacy and specificity with the ultimate aim of reducing the burden of common and rare genetic diseases in humans. Regulatory authorities currently require in-animal studies of safety, efficacy, and gene target specificity for nearly all genomeediting therapeutics under development for clinical use. Using NHP models, several ORIP-supported programs are currently developing and testing novel regeneration and gene therapy tools that span a wide range of organ systems, including kidney, brain, musculoskeletal, lung, liver, heart, and hematopoietic.

In addition, the NPRCs have identified at least a dozen rare genetic disorders within existing NHP groups, which serve as valuable disease models for treatment development and testing. Visit the <u>ORIP website</u> for more details on other supported NHP research resources.

Biospecimens Query System of the National Primate Research Centers Consortium: The Biospecimens Query System (BQS) is an informatics tool that helps investigators obtain such biological resources as tissues, serum, blood, DNA, and selected cells from the NPRCs and other participating NHP facilities. The BQS allows investigators to make prospective requests or to search for existing NHP biospecimens according to NHP species, sex, age, tissue type, and preservation method. Use of the BQS <u>requires registration</u> for approved access. **Primate Pathology Image Database:** The NPRCs' Consortium Primate Pathology Image Database (PPID) is an informatics tool that allows investigators to search for images of NHP organs and tissues based on disease name, etiology, organ, NHP genus and species, and image type. Some images are annotated, allowing the user to go directly to the anatomical region of interest. The PPID is also widely used as a learning tool by trainees in NHP clinical medicine and pathology. Use of the PPID requires <u>registration for approved</u> <u>access</u>.