



# 科技信息检索与利用

## 第五讲2

宋秀芳

中国科学院文献情报中心

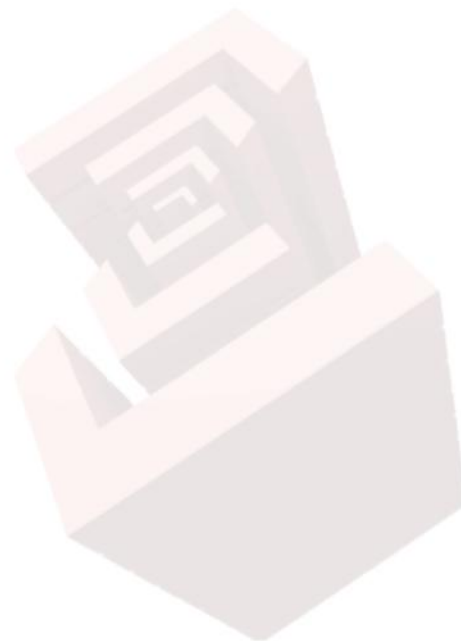


# 主要内容

图片表格

实验方法

基金信息





# 一、图表检索

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评级—RAT  
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语言学主题—LSU  
资源地点—RL  
起始页—PAGE  
输入中心编号, ASFA—TR  
过程—PRC  
通用名称—GN  
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遗传信息—GEN  
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- 命令行帮助
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# 命令行检索

高级检索

近期检索

运算符:

检索字段:

选择运算符

选择字段

CAP(coronavir\*)

所有日期

最近 7 天

最近 30 天

最近 3 月

最近 12 月

最近 3 年

在此日期...

在此日期之后...

在此日期之前...

特定日期范围...

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CAP(coronavir\*)

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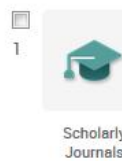
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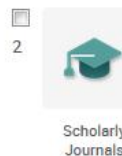
Bioinformatic prediction of immunodominant regions in spike protein for early diagnosis of the severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2)

Zhuang, Siqi; Tang, Lingli; Dai, Yufeng; Feng, Xiaojing; Fang, Yiyuan; 等. **PeerJ**; **San Diego** (Apr 8, 2021).

摘要/索引 全文文献 全文 - PDF 格式 (6 MB)

全文文献

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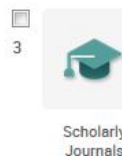
The Other Side of SARS-CoV-2 Infection: Neurological Sequelae in Patients

Alonso-Bellido, Isabel M; Bachiller, Sara; Vázquez, Guillermo; Cruz-Hernández, Luis; Martínez, Emilio; 等. **Frontiers in Aging Neuroscience; Lausanne** (Apr 6, 2021).

摘要/索引 全文文献 全文 - PDF 格式 (1 MB)

全文文献

显示摘要



The impact of coronavirus disease 2019 on surveillance colonoscopies in South Australia

Wassie, Molla M; Agaciak, Madelyn; Cock, Charles; Bampton, Peter; Young, Graeme P; 等. **JGH Open; [Richmond]** Vol. 5, Iss. 4, (Apr 2021): 486-492.

摘要/索引 全文文献 全文 - PDF 格式 (347 KB)

全文文献

显示摘要



## 二、实验方法查找

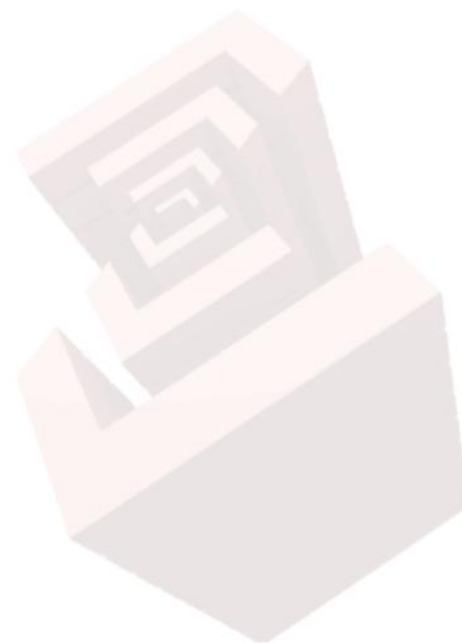
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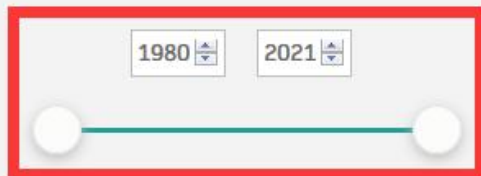
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## Publication Year



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- ☐ PCR 11324

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Protocol

Series: Methods In Molecular Biology

## Effects of Malignant Melanoma Initiating Cells on T-Cell Activation

Tobias Schatton, Ute Schütte, Markus H. Frank ✉

Although human malignant melanoma is a highly immunogenic cancer, both the endogenous antitumor immune response and melanoma immunotherapy often fail to control neoplastic progression. Accordingly, characterizing melanoma cell subsets capable of ...[more](#)

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- ☐ Transfection 5280
- ☐ Gel Electrophoresis 4825
- ☐ HPLC 4457
- ☐ Flow Cytometry 4090

## Antibody <sup>BETA</sup> ⓘ

- ☐ Anti-phospho-Akt antibody
- ☐ Anti-Caspase 3 antibody
- ☐ Anti-AKT antibody

## Organism

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- ☐ Homo sapiens 10278
- ☐ Mus musculus 7164
- ☐ Saccharomyces cerevisiae 6187
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- ☐ Rattus (rat) 3058
- ☐ Bos taurus 3042
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- ☐ Capra hircus 1099
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- ☐ Arabidopsis thaliana 738



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2021

# Selective Regional Isolation of Brain Microvessels

**Authors:**

Fernanda Medina-Flores <sup>1, 2</sup>, Gabriela Hurtado-Alvarado <sup>2</sup>, Beatriz Gómez-González  <sup>2</sup>

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Full text

## Abstract

The study of the regionalized function of the blood-brain barrier at the level of brain endothelial cells and pericytes is essential to understand the biological properties and molecular mechanisms regulating this biological barrier. The isolation of blood vessels from specific brain regions will allow to understand regional differences in susceptibility to pathological phenomena such as ischemia, traumatic brain injury, and neurodegenerative diseases, such as Alzheimer disease. Here, we propose an efficient and fast method to isolate brain endothelial cells and pericytes from a specific cerebral region. The isolated brain endothelial cells and pericytes are viable to perform conventional molecular and histological techniques such as Western blots, immunocytofluorescence, and scanning electron microscopy. [less](#)





## Selective Regional Isolation of Brain Microvessels

Fernanda Medina-Flores, Gabriela Hurtado-Alvarado,  
and Beatriz Gómez-González

### Abstract

The study of the regionalized function of the blood-brain barrier at the level of brain endothelial cells and pericytes is essential to understand the biological properties and molecular mechanisms regulating this biological barrier. The isolation of blood vessels from specific brain regions will allow to understand regional differences in susceptibility to pathological phenomena such as ischemia, traumatic brain injury, and neurodegenerative diseases, such as Alzheimer disease. Here, we propose an efficient and fast method to isolate blood vessels from specific brain regions.

## 2 Materials

Prepare fresh solutions with ultrapure water at room temperature. Filter the solutions, and then store them at the corresponding temperature indicated below.

### 2.1 Solutions for Brain Blood Vessel Isolation

#### 溶液配制

1.  $5\times$  PBS solution: Weigh 40.03 g NaCl, 1.01 g KCl, 7.2 g  $\text{Na}_2\text{HPO}_4$ , and 1.2 g  $\text{KH}_2\text{PO}_4$ , and transfer them to a 1 L beaker. Add 800 mL ultrapure water to the cylinder, mix with a magnetic stir bar, and adjust pH to 7.4 with a solution of 0.3 M HCl (pH 0.13) or 0.5 M NaOH (pH 13.6). Make up to 1 L with ultrapure water and mix. Store at  $4^\circ\text{C}$ .
2.  $1\times$  PBS solution: Add 200 mL of  $5\times$  PBS solution to the 1 L beaker. Add 600 mL of ultrapure water, mix, and adjust pH to 7.4 with a solution of 0.3 M HCl or 0.5 M NaOH. Make up to 1 L with water; mix with a magnetic stir bar. Filter the solution.

## 3 Methods

### 3.1 Blood Vessel Isolation from Selective Brain Regions

#### 操作步骤

1. Dissection instruments must be cleaned and placed in a disinfectant solution diluted in water 1:1000 to sterilize for 1 h at room temperature.
2. Add 1.5 mL 1% BSA solution to each 1.5 mL conical microtube, and store over night at  $4^\circ\text{C}$ . This avoids the attachment of microvessels to the walls of the microtubes.
3. Discard the 1% BSA solution from microtubes by decantation. Add 1 mL of SB to the microtube, and maintain it on ice during the experiment.
4. Obtain the head of one euthanized rat, and immediately place it on ice (see Note 4).
5. Use large scissors to remove the skin that is on top of the skull. Next, use dental extracting forceps to remove the skull, begin by carefully introducing the forceps tip on the foramen mag-

## 4 Notes

### 注意事项

1. The volume of the solution depends on the number of samples you will work with; 100 mL of 1% BSA solution and sucrose buffer are sufficient to process two brain regions from six rat brains.
2. The avidin-biotin complex solution must be prepared and stored at  $4^\circ\text{C}$  30 min before using it.
3. Osmium tetroxide is extremely toxic; use coat, gloves, and goggles to prepare it in a lab extraction hood.
4. The euthanasia method does not modify the viability of brain microvessels. Animals may be euthanized by anesthesia overdose (e.g., with sodium pentobarbital) or by any other accepted method for rats. Consider obtaining the tissue quickly as possible, and process it immediately.
5. To dissect the cerebral cortex, use iris scissors, and carefully cut horizontally to the brain surface a piece of 1.5 cm long and 3 mm depth for each hemisphere. To dissect the hippocampus, remove the debris from the cerebral cortex with medium scr-

## References

### 参考文献

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2. Keane J, Campbell M (2015) The dynamic blood-brain barrier. *FEBS J* 282(21):4067–4079. <https://doi.org/10.1111/febs.13412>
3. Armulik A, Abramsson A, Betsholtz C (2005) Endothelial/pericyte interactions. *Circ Res* 97(6):512–523. <https://doi.org/10.1161/01.RES.0000182903.16652.d7>
4. Bell RD, Winkler EA, Sagare AP et al (2010) Pericytes control key neurovascular functions and neuronal phenotype in the adult brain and during brain aging. *Neuron* 68(3):409–427. <https://doi.org/10.1016/j.neuron.2010.09.043>
5. Rustenhoven J, Jansson D, Smyth LC et al (2017) Brain Pericytes as mediators of Neuroinflammation. *Trends Pharmacol Sci* 38(3):291–304. <https://doi.org/10.1016/j.tips.2016.12.001>
6. Hamilton NB (2010) Pericyte-mediated regulation of capillary diameter: a component of neurovascular coupling in health and disease. *Front Neuroener* 2:2. <https://doi.org/10.3389/fnenc.2010.00005>
7. Sweeney MD, Sagare AP, Zlokovic BV (2018) Blood-brain barrier breakdown in Alzheimer disease and other neurodegenerative disorders. *Nat Rev Neurosci* 14(3):133–150. <https://doi.org/10.1038/nrn.2017.188>
8. Winkler EA, Bell RD, Zlokovic BV (2011) Central nervous system pericytes in health and disease. *Nat Neurosci* 14(11):1398–1405. <https://doi.org/10.1038/nn.2946>

## Featured

### Protocol

25 Sept 2020

### Quantification of SARS-CoV-2 neutralizing antibody by a pseudotyped virus-based assay

The production and titration of the SARS-CoV-2 S pseudotyped virus using a VSV-based pseudovirus production system in this protocol enable its use under biosafety level 2 conditions as well as in a neutralization assay to assess the level of neutralizing antibodies or molecular inhibitors in a sample.

Jianhui Nie, Qianqian Li ... Youchun Wang

### Protocol Extension

09 Apr 2021

### Assessing biofilm inhibition and immunomodulatory activity of small amounts of synthetic host defense peptides synthesized using SPOT-array technology

Here, we present a protocol for high-throughput screening of SPOT-peptide arrays to assess the antibiofilm, antimicrobial and immunomodulatory activities of synthetic peptides.

Hashem Etayash, Evan F. Haney & Robert E. W. Hancock

### Protocol

12 Apr 2021

### Fabrication, characterization and application

This protocol describes how to fabricate graphene electronic tattoos (GETs). GETs can be used for a variety of applications, including wearables, personal health monitoring, and biomedical research.

Bo Liu, Keren, Chidab, Kishor, Anand ... Delli, Akhmad

## Current issue >

### Tutorial: assessing metagenomics software with the CAMI benchmarking toolkit

Fernando Meyer, Till-Robin Lesker ... Alice C. McHardy

**Review Article** | 01 Mar 2021

### The in vitro multilineage differentiation and maturation of lung and airway cells from human pluripotent stem cell-derived lung progenitors in 3D

Ana Luisa Rodrigues Toste de Carvalho, Hsiao-Yun Liu ... Hans-Willem Snoeck

**Protocol Extension** | 01 Mar 2021

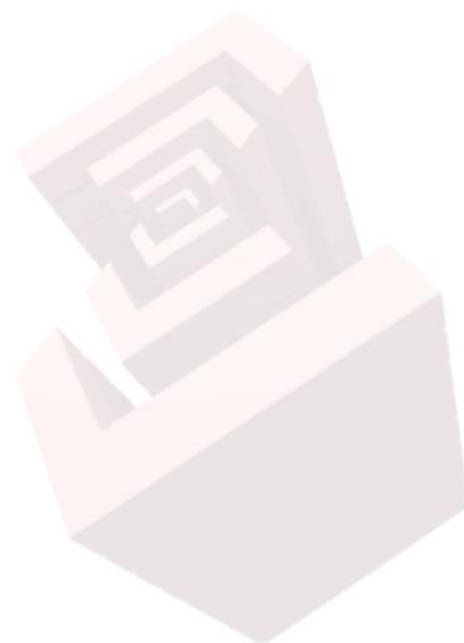
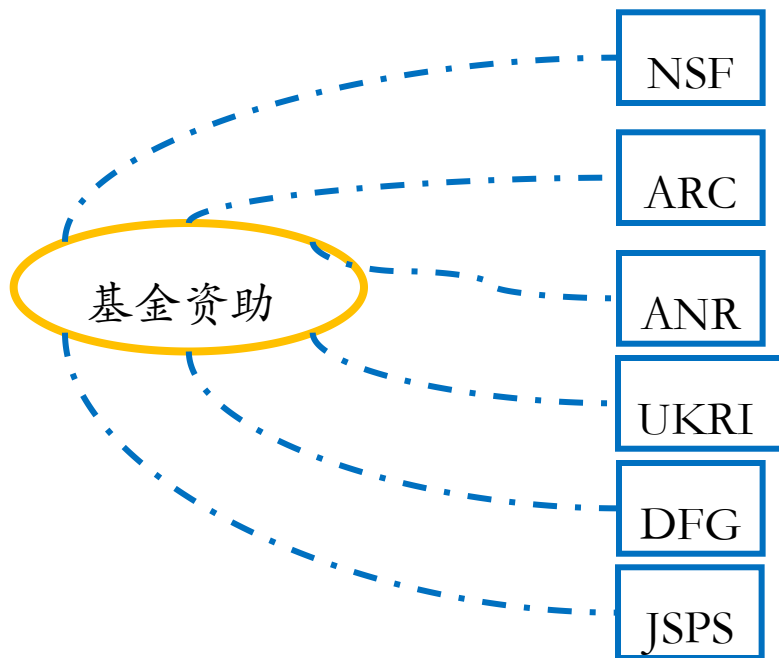
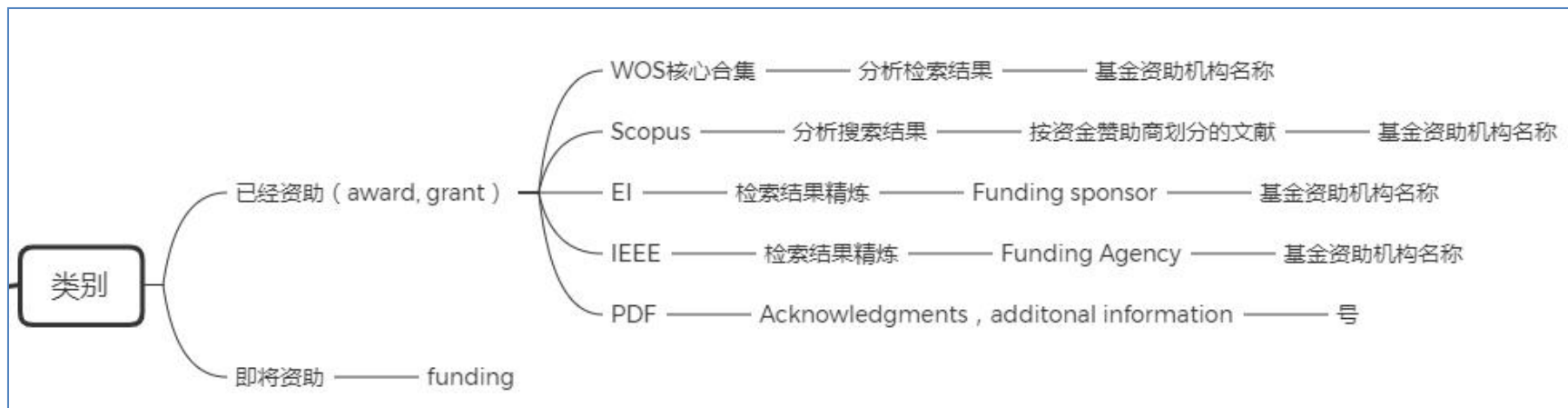
### Infection of zebrafish larvae with human norovirus and evaluation of the in vivo efficacy of small-molecule inhibitors

Jana Van Dycke, Arno Cuvry ... Joana Rocha-Pereira

**Protocol Extension** | 09 Apr 2021



### 三、基金信息检索







# (一) 美国国家科学基金

## 1、NSF 主页

The screenshot shows the NSF homepage with a blue header. The NSF logo is on the left, followed by the text "National Science Foundation" and "WHERE DISCOVERIES BEGIN". On the right, there is a search bar and links for "Operating Status" (with an "ALERT" badge) and "Contact | Help". Below the header is a navigation bar with links: "Research Areas", "Funding" (highlighted with an orange box), "Awards" (highlighted with an orange box), "Document Library", "News", and "About NSF". Below the navigation bar is a green banner with the text "ALERT" and "资助动态 资助项目" (Funding Dynamics Funding Projects). Below the banner is a white box with the text "Read the latest information from NSF on coronavirus (COVID-19)" and "Read statement on deadline extensions and flexibilities." Below the white box is a large image of a desert landscape with a wooden sign that reads "ENTERING PINE RIDGE INDIAN PRESERVATION". To the right of the image is a black box with the text "SCIENCE MATTERS" and "Tribal colleges partner to fill COVID-19 data gaps".

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# (一) 美国国家科学基金

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<input type="checkbox"/> Include Co-Principal Investigator in name search	<input type="text" value="Zip Code"/>
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受资助人

Program Information	
<input type="text" value="NSF Organization"/> Select one	<p>HINT: The "Program" box searches both program element and program reference names and codes.</p> <input type="text" value="Program"/>
<input type="text" value="Element Code"/> Any All	<input type="text" value="Program Officer"/>
<input type="text" value="Reference Code"/> Any All	

项目信息

Additional Information	
<input type="text" value="Keyword"/> HINT: The Keyword field searches on the title and abstract only. <input type="checkbox"/> Search Award Title Only	<p>HINT: Data prior to 1976 may be less complete.</p> <input checked="" type="checkbox"/> Active Awards <input type="checkbox"/> Expired Awards
<input type="text" value="Award Number"/> From To	<input type="text" value="Original Award Date"/> From To
<input type="text" value="Award Amount"/> Select one	<input type="text" value="Start Date"/> From To
<input type="text" value="Award Instrument"/> Select one	<input type="text" value="End Date"/> From To

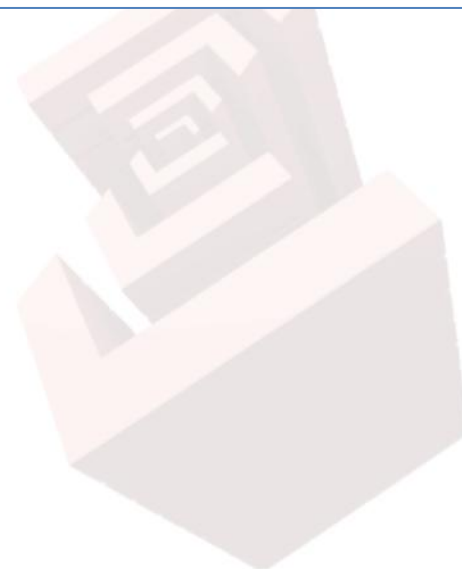
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<div>You Searched For:</div> <div>3D print</div> <div>Active Awards true</div> <div>Refined by</div> <div>Refine Search</div> <div>State</div> <div>Alaska(5)</div> <div>Alabama(37)</div> <div>Arkansas(12)</div> <div>Arizona(59)</div> <div>California(250)</div> <div>Show More ...</div> <div>NSF Organization</div> <div>Office Of The Director(30)</div> <div>Office Of Information &amp; Resource Mgmt(5)</div>	<div><div>Structured Filaments for High Performance 3D Printed Plastic Objects</div><div>Award Number:2011289; Principal Investigator:Bryan Vogt; Co-Principal Investigator;; Organization:Pennsylvania State Univ University Park;NSF Organization:CMMI Start Date:08/14/2019; Award Amount:\$331,309.00; Relevance:47.63;</div></div> <div><div>SBIR Phase II: Increasing Maker Manufacturing through 3D Printing with Reclaimed Plastic &amp; Direct Drive Pellet Extrusion</div><div>Award Number:1853153; Principal Investigator:Samantha Snabes; Co-Principal Investigator;; Organization:RE3D Inc.;NSF Organization:IIP Start Date:04/15/2019; Award Amount:\$1,442,973.00; Relevance:47.63;</div></div> <div><div>EAGER/GOALI: 3D Printing of Nanostructured Battery Electrodes</div><div>Award Number:1938787; Principal Investigator:Vibha Kalra; Co-Principal Investigator:Thomas Greszler; Organization:Drexel University;NSF Organization:CMMI Start Date:10/01/2019; Award Amount:\$124,436.00; Relevance:47.63;</div></div> <div><div>I-Corps: Swarm Three Dimensional Printing and Assembly Platform</div><div>Award Number:1928756; Principal Investigator:Wenchao Zhou; Co-Principal Investigator;; Organization:University of Arkansas;NSF Organization:IIP Start Date:06/15/2019; Award Amount:\$50,000.00; Relevance:47.63;</div></div> <div><div>CAREER: Adaptive Tactile Picture Books for Blind Children during Emergent Literacy</div><div>Award Number:1453771; Principal Investigator:Tom Yeh; Co-Principal Investigator;; Organization:University of Colorado at Boulder;NSF Organization:IIS Start Date:02/01/2015; Award Amount:\$581,999.00; Relevance:47.63;</div></div>





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**Award Amount**

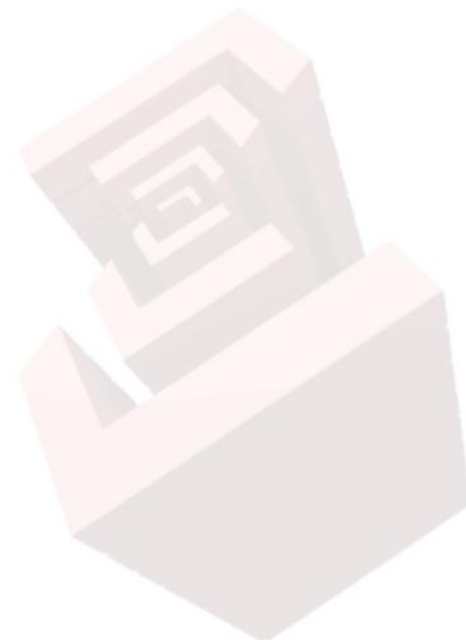
less than or equal \$50,000(118)  
between \$50,001 - \$100,000(119)  
between \$100,001 - \$500,000(1541)  
between \$500,001 - \$1,000,000(426)  
more than \$1,000,000(182)

额度  
范围

**Award Instrument**

Standard Grant(1819)  
Continuing Grant(501)  
Cooperative Agreement(36)  
Fellowship(25)  
Contract Interagency Agreement(5)

资助类型







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Award Number	Title	NSF Organization	Programs	Start Date	Principal Investigator	Co-Principal Investigator	State	Organization	Awarded Amount To Date	Program Officer
2011289	Structured Filaments for High Performance 3D Printed Plastic Objects	CMMI	Special Initiatives , Materials Eng. & Processing	08/14/2019	Vogt, Bryan		PA	Pennsylvania State Univ University Park	\$331,309.00	Andrew Wel
1853153	SBIR Phase II: Increasing Maker Manufacturing through 3D Printing with Reclaimed Plastic & Direct Drive Pellet Extrusion	IIP	SBIR Phase II	04/15/2019	Snabes, Samantha		PR	RE3D Inc.	\$1,442,973.00	Rajesh Meh
1938787	EAGER/GOALI: 3D Printing of Nanostructured Battery Electrodes	CMMI	AM-Advanced Manufacturing , GOALI-Grnt Opp Acad Lia wIndus	10/01/2019	Kalra, Vibha	Greszler, Thomas;	PA	Drexel University	\$124,436.00	Khershed Cooper
1928756	I-Corps: Swarm Three Dimensional Printing and Assembly Platform	IIP	I-Corps	06/15/2019	Zhou, Wenchao		AR	University of Arkansas	\$50,000.00	Ruth Shums
1453771	CAREER: Adaptive Tactile Picture Books for Blind Children during Emergent Literacy	IIS	HCC-Human-Centered Computing	02/01/2015	Yeh, Tom		CO	University of Colorado at Boulder	\$581,999.00	Ephraim Glinert
1829664	SBIR Phase II: Simulation for structural integrity of as manufactured 3D printed parts	IIP	SBIR Phase II	10/01/2018	Adams, Brady		WY	Teton Composites	\$1,418,740.00	Peter Atherton
1815585	CHS: Small: Collaborative Research: 3D Printing for High Fidelity Image Reproduction	IIS	HCC-Human-Centered Computing	09/01/2018	Matusik, Wojciech		MA	Massachusetts Institute of Technology	\$250,000.00	Ephraim Glinert



## 4、资助详细信息

### Award Abstract # 2011289 Structured Filaments for High Performance 3D Printed Plastic Objects

<b>NSF Org:</b>	<a href="#">CMMI Div Of Civil, Mechanical, &amp; Manufact Inn</a>
<b>Awardee:</b>	PENNSYLVANIA STATE UNIVERSITY, THE
<b>Initial Amendment Date:</b>	December 30, 2019
<b>Latest Amendment Date:</b>	June 26, 2020
<b>Award Number:</b>	2011289
<b>Award Instrument:</b>	Standard Grant
<b>Program Manager:</b>	Andrew Wells awells@nsf.gov (703)292-7225 CMMI Div Of Civil, Mechanical, & Manufact Inn ENG Directorate For Engineering
<b>Start Date:</b>	August 14, 2019
<b>End Date:</b>	July 31, 2022 (Estimated)
<b>Total Intended Award Amount:</b>	¥276,309.00
<b>Total Awarded Amount to Date:</b>	\$331,309.00
<b>Funds Obligated to Date:</b>	FY 2018 = ¥276,309.00 FY 2020 = ¥55,000.00
<b>History of Investigator:</b>	Bryan Vogt (Principal Investigator) bdv5051@psu.edu (814)863-5459

<b>Awardee Sponsored Research Office:</b>	Pennsylvania State Univ University Park 201 Old Main CENTRE, University Park, PA US 16802-1503 (814)865-1372
<b>Sponsor Congressional District:</b>	05
<b>Primary Place of Performance:</b>	Pennsylvania State Univ University Park 201 Old Main CENTRE, University Park, PA US 16802-1503
<b>Primary Place of Performance Congressional District:</b>	05
<b>DUNS ID:</b>	003403953
<b>Parent DUNS ID:</b>	003403953
<b>NSF Program(s):</b>	Special Initiatives, Materials Eng. & Processing
<b>Primary Program Source:</b>	040100 NSF RESEARCH & RELATED ACTIVIT 040100 NSF RESEARCH & RELATED ACTIVIT
<b>Program Reference Code(s):</b>	091Z, 1444, 1467, 1773, 8021, 8025
<b>Program Element Code(s):</b>	088y, 1642, 8092
<b>Award Agency Code:</b>	4900
<b>Fund Agency Code:</b>	4900
<b>CFDA Number(s):</b>	47.041





## ABSTRACT

This grant will support research that will contribute new fundamental knowledge related to the design of the feedstock for one common method of 3D printing to provide guidance for improved plastic parts to enable the translation from rapid prototyping to manufacture. Additive manufacturing generates near net shape object of virtually any shape from a digital computer model and is critical to the development of new manufacturing approach essential for the national productivity. Additive manufacturing is commonly called 3D printing and offers revolutionary possibilities in terms of massive customization for the individual consumer, so the fit is ergonomically perfect. However, almost all additive manufacturing processes for plastic parts lead to inherent weaknesses in the parts that make them inferior to traditionally manufactured plastics. This grant supports fundamental research to provide needed knowledge for the development of improved performance of additive manufactured plastic parts through scalable changes in the feedstock for one type of 3D printing called fused filament fabrication. The new materials will be compatible with existing printers, including consumer printers that are available to the general public, but will enable significant improvements in obtaining parts that better match the desired dimensions and with improved toughness. Additive manufacture of plastic parts is growing with applications from healthcare to assist doctors with planning surgery and custom implants for craniofacial restoration to aerospace parts to decrease the weight of non-critical components. Through improving the performance of plastic parts, this research will benefit the U.S. economy and society by extending the potential applications for additively manufactured plastic parts. This research involves several disciplines including manufacturing, materials science, and mechanical engineering, which will provide a unique educational experience for the students involved in this research. Additionally, the materials produced will be compatible with many commercial 3D printers, including those found in some K-12 schools, so outreach to these schools will provide the students with an opportunity to learn about additive manufacturing and design of materials with a hands-on approach.

The design of structured filaments is hypothesized to overcome the intrinsic trade-off between mechanical properties and dimensional accuracy associated with extrusion-based polymer additive manufacturing. Generally, there is poor interlayer strength during the print as the interdiffusion of polymer chains is limited by the temperature and increasing the printing temperature leads to flow and deformation of the printed part. This research seeks to overcome this trade-off with a core-shell structure to the feedstock filament, where the core provides mechanical reinforcement to inhibit flow, while the shell solidifies at lower temperature to provide interlayer strength. However, the fundamental requirements associated with the materials selection and the print processing are poorly understood for the core-shell materials in additive manufacturing. This research will fill the knowledge gap on the relationships between solidification temperature, mechanical properties and miscibility of the core and shell polymers through systematic experimental investigation. The research team will establish relationships between process parameters, intrinsic material properties of the polymers, the dimensional accuracy of the printed part, and mechanical properties to provide insights into the limitations of extrusion-based additive manufacturing for plastic objects.

This award reflects NSF's statutory mission and has been deemed worthy of support through evaluation using the Foundation's intellectual merit and broader impacts review criteria.



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## 3D printing of hollow geometries using blocking liquid substitution stereolithography

[Aftab A. Bhanvadia](#), [Richard T. Farley](#), [Youngwook Noh](#) & [Toshikazu Nishida](#) [✉](#)

[Scientific Reports](#) **13**, Article number: 434 (2023) | [Cite this article](#)

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### 1. 3D printing of hollow geometries using blocking liquid substitution stereolithography

<https://doi.org/10.1038/s41598-022-26684-z> [DOI](#)

Bhanvadia, Aftab A. ; Farley, Richard T. ; Noh, Youngwook ; Nishida, Toshikazu ( December 2023 , Scientific Reports)

Abstract Micrometer scale arbitrary hollow geometries within a solid are needed for a variety of applications including microfluidics, thermal management and metamaterials. A major challenge to 3D **printing** hollow geometries using stereolithography is the ability to retain empty spaces in between the solidified regions. In order to prevent unwanted polymerization of the trapped resin in the hollow spaces—known as **print-through**—significant constraints are generally imposed on the primary process parameters such as resin formulation, exposure conditions and layer thickness. Here, we report on a stereolithography process which substitutes the trapped resin with a UV blocking liquid to mitigate **print-through**. We investigate the mechanism of the developed process and determine guidelines for the formulation of the blocking liquid. The reported method decouples the relationship between the primary process parameters and their effect on **print-through**. Without having to optimize the primary process parameters to reduce **print-through**, hollow heights that exceed the limits of conventional stereolithography can be realized. We demonstrate fabrication of a variety of complex hollow geometries with cross-sectional features ranging from tens of micrometer to hundreds of micrometers in size. With the framework presented, this method may be employed for 3D **printing** functional hollow geometries for a variety of applications, and [more »](#)

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