

## Warning Notice

OMB Control Number: 0693-XXXX Expiration Date: XX/XX/XXXX

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Privacy Act Statement Authority: The Bayh-Dole Act (35 USC §§ 200-212) and its implementing regulations (37 CFR 401). Purpose: The Bayh-Dole Act (35 USC §§ 200-212) and its implementing regulations (37 CFR 401) require Contractors to report information related to inventions and patents to the federal agency which awarded the relevant funding. The purpose of this legislation included promoting utilization of inventions resulting from federal R&D funding, encourage collaboration between non-profit and for-profit entities, increase commercialization and public availability of inventions made in the United States, and to ensure that the Government obtains sufficient rights in federally funded inventions to meet the needs of the Government and protect the public. To facilitate these and other objectives as well as to track both compliance with and effectiveness of this legislation, collection of data is required.

Routine Uses: NIST will utilize the information collected to facilitate the objectives of the Bayh-Dole Act and to track both compliance with and effectiveness of this legislation. DEPT-23: Information Collected Electronically in Connection with Department of Commerce Activities, Events, and Programs. Disclosure: Furnishing this information is mandatory to obtain benefits. When you submit the form, you are indicating your voluntary consent for NIST to use of the information you submit for the purpose stated.



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Invention/Patent/Utilization Tree

Invention Details

Invention Report Number \*

Institution

System only accepts a valid invention



Please search and choose a valid invention

Invention Title

Primary Agency

Invention Report Date

Title Elect Date

Utilization Details

Reporting Year (YYYY) \*

Indicate the latest stage of development of any product arising from this invention, according to the following categories: \*

If any product arising from this invention has reached the market, what was the calendar year of the first commercial sale? (YYYY) \*

In the designated reporting period, what was the total income received as a result of license or option agreements? Do not include specific patent costs reimbursement.

\$0.00

Did the grantee organization/contractor or any of the exclusive licensees request a waiver of the U.S. manufacturing requirements in the designated reporting period?

Yes  No

If yes, how many such waivers were obtained?

0

In the designated reporting period, how many exclusive licenses and/or options are active?

0

Number of domestic manufacturing licenses

0

In the designated reporting period, how many non-exclusive licenses and/or options are active?

0

How many licenses and/or options of any type to small businesses (<500 employees) are active in the designated reporting period?

0

Total Gross Sales

\$0.00

FDA Approved Commercial Product Details

Please provide the commercial name of any FDA-approved products, utilizing this invention, that have first reached the market during the designated reporting period. Please note: Commercial names should be limited to FDA-approved products that first reached the market during the designated reporting period. Please remove the "Public" checkmark from any FDA-approved product that you do not want to appear on a publicly available list of products arising from your funding agreement.

Commercial Products

Product Name	FDA Approval Number	FDA Approval Type	Public (Y/N)	Government Review Status	Report Date
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No records found.

+ Add FDA Approved Commercial Product

active in the designated reporting period?

Total Gross Sales

### FDA Approved Commercial Product Details

Please provide the commercial name of any FDA-approved products, utilizing this invention, that have first reached the market during the reporting period. Please note: Commercial names should be limited to FDA-approved products that first reached the market during the reporting period from any FDA-approved product that you do not want to appear on a publicly available list of products arising from this invention.

**Commercial Products**

Product Name	FDA Approval Number	FDA Approval Type	Public (Y/N)
No records found.			

[+ Add FDA Approved Commercial Product](#)

### Submitted Utilization Reports

### Add FDA Approved Commercial Product

Commercial Product Name \*

FDA Approved Number

Commercial Product Type

Government Review Status

Public Announced

Yes  No

[Cancel](#) [Reset](#) [Add Commercial Product](#)

### Impact on U.S Economy

Approximate number of new US-based jobs created because of commercialization efforts during the reporting period.

Number of new US-based companies created from the commercialization efforts during the reporting period.

Describe how the development and commercialization of the subject invention comply with any required U.S. manufacturing requirements (e.g., U.S. Preference, a U.S. Competitive Clause, U.S. Manufacturing Plan, etc.)

Licensed to a US company, and made in the US

*955 characters remaining.*

### Unique Commercial Products Made Through the Use of or Embodying the Subject Inventions

Please note: For each product, also list the associated NAICS code (if applicable)

[+ Add Manufacturing Commercial Product](#)

### Impact on U.S Economy

Approximate number of new US-based jobs created because of commercialization efforts during the reporting period.

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### Unique Commercial Products Made Through the Use of or Embodying the Subject Inventions

Please note: For each product, also list the associated NAICS code (if applicable)

[+ Add Manufacturing Commercial Product](#)

### Submitted Utilization Reports

### Add Manufacturing Commercial Product

Unique Commercial Product Name \*

NAICS Code

Manufacturing Country	Manufacturing State	First Date of Manufacturing	Manufacturing Date Type (actual/expected)	Manufacturing Product Quantity
No records found.				

[+ Add Manufacturing Location](#)

[Cancel](#) [Reset](#) [Add Commercial Product](#)