

Downstate Biotechnology Incubator Application

Business Name: _____

SECTION I: CONTACT INFORMATION

Contact Person				
Dr. <input type="checkbox"/>	Mr. <input type="checkbox"/>	Ms. <input type="checkbox"/>	First Name	Last Name
Phone #:		Mobile Phone #:	Email:	

SECTION II: BUSINESS INFORMATION

Current Business Address:		

<i>Number and Street</i>		

<i>City, State and Zip Code</i>		
_____	_____	_____
<i>Phone #</i>	<i>Fax #</i>	<i>Email</i>

Principal #1:

Dr. <input type="checkbox"/>			
Mr. <input type="checkbox"/>			
Ms. <input type="checkbox"/>			
	_____	_____	_____
	<i>First Name</i>	<i>Last Name</i>	<i>Title</i>
	_____	_____	_____
	<i>Ownership (%)</i>	<i>Phone #</i>	<i>Email</i>

Principal #2:

Dr. <input type="checkbox"/>			
Mr. <input type="checkbox"/>			
Ms. <input type="checkbox"/>			
	_____	_____	_____
	<i>First Name</i>	<i>Last Name</i>	<i>Title</i>
	_____	_____	_____
	<i>Ownership (%)</i>	<i>Phone #</i>	<i>Email</i>

Principal #3:

Dr. <input type="checkbox"/>			
Mr. <input type="checkbox"/>			
Ms. <input type="checkbox"/>			
	_____	_____	_____
	<i>First Name</i>	<i>Last Name</i>	<i>Title</i>
	_____	_____	_____
	<i>Ownership (%)</i>	<i>Phone #</i>	<i>Email</i>

Please attach separate sheet(s) for any additional Principals.

Other Person(s) Authorized to Negotiate/Contract on behalf of Business:

Dr. <input type="checkbox"/>			
Mr. <input type="checkbox"/>			
Ms. <input type="checkbox"/>			
	<u>First Name</u>	<u>Last Name</u>	<u>Title</u>
<u>Phone #</u>	<u>Email</u>		

Dr. <input type="checkbox"/>			
Mr. <input type="checkbox"/>			
Ms. <input type="checkbox"/>			
	<u>First Name</u>	<u>Last Name</u>	<u>Title</u>
<u>Phone #</u>	<u>Email</u>		

Please attach separate sheet(s) for any additional authorized persons.

Describe exactly what the Company will do in the space:

Business Operations began/will begin in (month/year): _____ Incorporated in: State _____
Year _____

Capitalization \$ _____ FY 20 _____ Operating Budget: \$ _____ FY 20 _____ Sales Revenues: \$ _____
FY 20 _____

Number of Employees: _____ Full Time _____ Part Time _____ Research/Science _____ Tech/Other

Describe how the Company will create a symbiotic relationship with SUNY Downstate Medical Center (e.g., internships, hiring graduates, collaborations, teaching, etc.).

SECTION III: START-UP NY

Are you interested in applying for START-UP NY? ____ Yes ____ No ____ Already a member of START-UP NY

If the Company is not incorporated in New York State, has it registered with the New York Secretary of State as a "Foreign Company Doing Business in New York State"? ____ Yes ____ No

SECTION IV: REQUIREMENTS

Approximate Space Required: Wet Lab _____ SF Office/Other _____ SF Total _____ SF

Required Start Date of Occupancy (approx): Month _____ Year _____ Length of Occupancy (approx): _____ yrs

	<u>Required</u>	<u>Number (if applicable)</u>
Vacuum	<input type="checkbox"/>	_____
Fume Hoods (1 per lab)	<input type="checkbox"/>	_____
Biosafety Hoods	<input type="checkbox"/>	_____
Gas	<input type="checkbox"/>	_____
Benches	<input type="checkbox"/>	_____
Animal facility	<input type="checkbox"/>	_____
Other (specify):		_____

NO RADIOACTIVITY CAN BE USED IN THE INCUBATOR

SECTION V: COMPANY OPERATIONS

Please provide descriptions for each of the following as they pertain to your space usage and requirements. Attach additional sheets if necessary.

☐ **HUMAN SAMPLES** ☐ **Not Applicable**

- Specimen type:**
- ☐ Blood
 - ☐ Body fluid
Type: _____
 - ☐ Cell/Organ/Tissue
(Both primary and commercially procured)
Name: _____
 - ☐ Cell line/culture
Name: _____

Known hazards and infectious agents and required Biosafety level for proper handling (e.g., HIV-1, HBV, HCV):

Describe measures to protect personnel: _____

☐ **INFECTIOUS AGENTS** (attach additional forms for each infectious agent) ☐ **Not Applicable**

Is this agent infectious to animals? ☐ No ☐ Yes

Is this agent infectious to humans? ☐ No ☐ Yes

Does this agent elaborate a toxin? ☐ No ☐ Yes

Is there a vaccine available for use in humans against this agent or its components?

☐ No ☐ Yes _____

Identify any precautionary medical practices that will be implemented, if any _____

Identify all personnel who will work on this project, providing documentation indicating their level of training and experience in working with infectious agents. List all certifications required by FDNY, including C-14 Certificate of Fitness for Non-Production Chemical Laboratories:

If a bacterial agent, provide an antibiogram: (attach additional sheets as needed)

How is the infectious agent propagated in the laboratory?

Specify methods of inactivation/decontamination and disposal of the agent or contaminated materials:

How will the agent stored in your laboratory?

☐ **ANIMAL WORK**

☐ **Not Applicable**

Will you be working with animals? ___ Yes ___ No

If yes, where will this be done? _____

☐ **RECOMBINANT DNA**

☐ **Not Applicable**

Are recombinant DNA procedures used in your laboratory limited to PCR amplification of DNA fragments (i.e., no subsequent cloning of amplified DNA)?

☐ Yes (Only check this if your recombinant DNA studies are exempt from restrictions described in the *NIH Guidelines for Research Involving Recombinant DNA Molecules*).

☐ No (Please provide the following information using a separate table for each gene):

Biological source of DNA or gene (2):			
Name and function of the gene:			
Selectable marker			
Host:			
Cell/animal recipient:			
Assessment of levels of physical and biological containment (consult current <i>NIH Guidelines for Research Involving Recombinant DNA Molecules</i> at http://www.nih.gov/od/orda/toc.html)	<input type="checkbox"/> Risk group 1	<input type="checkbox"/> BSL - 1	<input type="checkbox"/> Animal BSL-1
	<input type="checkbox"/> Risk group 2	<input type="checkbox"/> BSL - 2	<input type="checkbox"/> Animal BSL-2
	<input type="checkbox"/> Risk group 3	<input type="checkbox"/> BSL - 3	<input type="checkbox"/> Animal BSL-3

☐ **TOXIC/HAZARDOUS SUBSTANCES**

☐ **Not Applicable**

Name of the toxic/hazardous substance (include carcinogenic, mutagenic, teratogenic substances):
Attach a Material Safety Data Sheet (MSDS) for each substance.

Each Company must maintain on-site an up-to-date file of MSDS documents for each reagent in their lab.

Is this substance to be given to animals? ☐ No ☐ Yes

Amount of the substance to be kept in the laboratory:

Storage location:

Use location:

Inventory control procedure:

Method of deactivation:

Risk of human exposure and containment procedure?
(describe measure to protect personnel)

☐ **FLOW CYTOMETRIC HAZARD ASSESSMENT** ☐ **Not Applicable**

1. Cells to be used:

- ☐ Fresh or frozen animal cell
☐ Fresh or frozen human cells
☐ Cell lines

2. If a cell line to be used, indicate name(s)/designation(s):

3. If the cells are from human donors, were the donors screened for bloodborne pathogens?

☐ Yes; proceed to # 4

☐ No; proceed to # 6

4. Any pathogens the sample may contain?

☐ None

- ☐ HIV ☐ HCV
☐ HBV ☐ Other

5. Has the infectious agent been inactivated?

☐ No

☐ Unknown

☐ Yes; describe method

6. Do the cells contain infectious agents such as viruses, bacteria, fungi, protozoa?

☐ No

☐ Yes; give name(s):

7. Were the cells genetically engineered?

- ☐ No
☐ Yes

Was a virus used?

- ☐ Adenovirus ☐ Retrovirus
☐ Lentivirus ☐ Herpes virus

☐ **CHEMICAL USAGE** (detail types, quantities, and method of storage) ☐ **Not Applicable**

The Company is responsible for the safe storage and handling of all chemicals, including appropriate disposal.

☐ **WASTE GENERATION** ☐ **Not Applicable**

[**Regulated Waste** means liquid or semi-liquid blood or other potentially infectious materials, chemical waste or hazardous substances; contaminated items that would release blood or other potentially infectious materials in a liquid or semi-liquid state if compressed; items that are caked with dried blood or other potentially infectious materials and are capable of releasing these materials during handling; contaminated sharps; and pathological and microbiological wastes containing blood or other potentially infectious materials.]

Does your work generate waste that would be considered “regulated waste”? ____ Yes ____ No
If “Yes”, detail types, quantities and disposal plan

- Will the waste be autoclaved before leaving the facility? ____ Yes ____ No
- Will the waste be “red bagged” before leaving the facility? ____ Yes ____ No
- Will you have sharps disposal containers appropriately placed in your laboratory? ____ Yes ____ No

The Company is responsible for the proper storage, handling and disposal of all regulated waste.

☐ **IS THERE SPECIALIZED EQUIPMENT YOU WILL BE USING THAT WE SHOULD BE AWARE OF?** ☐ **Not Applicable**

Are you or do you plan on being CLIA (CLEP)? ☐ Yes ☐ No

Do you have a Safety Plan? ☐ Yes ☐ No ☐ Under Development
(All companies occupying laboratory space are required to have a Safety Plan.)

Company's Safety Officer is responsible for implementing company's safety plan and monitoring ongoing compliance. Please note the Company is responsible for all applicable safety guidelines, approvals and training.

Safety Officer Name: _____ Phone #: _____ Email: _____

This application has been completed by:

Name

Signature

Date

Please submit this completed application along with your Business Plan, R&D Plan, audited financial statements, resumes of principals, and supporting documentation to:

**Blake Adair
Executive Director
Downstate Biotechnology Incubator
760 Parkside Avenue
Brooklyn, NY 11226**

Email: blake.adair@downstate.edu