



JOB INFORMATION

<i>Job Code:</i>	188017
<i>Job Title:</i>	cGMP Core Operator
<i>FLSA Status:</i>	Exempt
<i>Supervisory:</i>	May lead one or more employees performing similar work.
<i>Job Family:</i>	Clinical Research
<i>Job Family Group:</i>	USC Job Families
<i>Management Level:</i>	7 Individual Contributor

JOB SUMMARY

Supports daily manufacturing operations in current Good Manufacturing Practice (cGMP) facilities. Maintains materials, gowning, cleaning supplies, and consumable supplies, manages manufacturing waste streams, cleans equipment, oversees documentation, and supports in other manufacturing duties as required.

JOB QUALIFICATIONS:

Education

<i>Req</i>	<i>Pref</i>	<i>Degree</i>	<i>Field of Study</i>	
X		Bachelor's degree		In
X			Pharmacy	Or
X			Biology	Or
X			in related field(s)	
	X	Master's degree		In
	X		Biotechnology	Or
	X		in related field(s)	

Additional Education

Check here if experience may substitute for some of the above education.

Combined experience/education as substitute for minimum education

Work Experience

<i>Req</i>	<i>Pref</i>	<i>Work Experience</i>	<i>Experience Level</i>	
X		3 years	in cellular or biological manufacturing with laboratory responsibilities.	

Additional Work Experience

Check here if education may substitute for some of the above work experience.

Combined experience/education as substitute for minimum work experience

Knowledge, Skills and Abilities

Req	Pref	Functional Skills
X		Demonstrated knowledge base with Good Manufacturing Practices (e.g., cGMPs, GLPs, GDPs).
X		Extensive experience with standard operating procedures in a cGMP laboratory setting.
X		Demonstrated ability to work as an individual contributor and in a dynamic team environment.
X		Excellent written and oral communication skills.
	X	Demonstrated knowledge of all aspects of biotechnology and cell therapy.
	X	Demonstrated passion for solving complex scientific issues.
	X	Experience with Food and Drug Administration regulations and clinical trials.
	X	Extensive leadership experience.

Other Job Factors

JOB ACCOUNTABILITIES

	% Time	Essential	Marginal	N/A
Performs all product manufacturing duties in cGMP facilities, adhering to all applicable gowning and safety requirements. Prepares and properly stores all incoming, intermediate and final products. Performs quality control for incoming, in-process and final products. Generates batch records, chain of custody forms, certificates of analysis and other records. Maintains complete documentation throughout the manufacturing process.				
Orders cGMP facility equipment and supplies. Develops an inventory system for ongoing supplies and orders. Supports senior facility staff in maintaining cGMP budgets to ensure efficient use of equipment and materials. Establishes preventative maintenance schedule routines, and designs forms for maintaining documentation of equipment maintenance.				
Acts as technical consultant to clinicians and researchers, providing technical assistance within the facility as necessary. Develops, implements and documents standard operating procedures, making updates as necessary. Instructs and oversees cGMP technicians and instructs graduate students and post-doctoral fellows in proper procedures within cGMP facilities. Serves as a resource to cGMP facility management in identifying and assessing the appropriate complement of resources and support needed to successfully implement and execute projects.				
Applies appropriate labels to incoming and outgoing products and monitors for proper labeling and handling during manufacturing. Ensures biohazard waste is handled appropriately and that medical waste stream rules and regulations are followed. Works with management to ensure facilities' compliance with all applicable regulations.				
Plans and organizes workflows to meet established turnaround times, dispense dates and deadlines. Attends routine meetings with management team for progress reports on projects, facility needs, and discussion of any other required items. Recommends process improvements as appropriate.				
Promotes an environment that fosters inclusive relationships and creates unbiased opportunities for contributions through ideas, words, and actions that uphold principles of the USC Code of Ethics.				

Other Requirements

Essential:	Emergency Response/Recovery	Essential:	Mandated Reporter
	In the event of an emergency, the employee holding this position is required to "report to duty" in accordance with the university's Emergency Operations Plan and/or the employee's department's emergency response and/or recovery plans. Familiarity with those plans and regular training to implement those plans is required. During or immediately following an emergency, the employee will be notified to assist in the emergency response		A mandated reporter who in his or her professional capacity has knowledge of, or reasonably suspects a person who is under the age of 18 years, elderly, or a dependent adult has been the victim of abuse or neglect must report the suspected incident. The reporter must contact a designated agency immediately or as soon as practically possible by telephone or in writing within 36 hours. By virtue of the associated job duties, this position qualifies as a mandated reporter as required by state law

Other Requirements

<i>Essential:</i>	<i>Emergency Response/Recovery</i>	<i>Essential:</i>	<i>Mandated Reporter</i>
	efforts, and mobilize other staff members if needed.		and USC's policy at: https://policy.usc.edu/mandated-reporters/
<i>Campus Security Authority (CSA)</i>			<i>Essential:</i>
By virtue of the associated job duties, this position qualifies as a Campus Security Authority as required by law and USC's policy at: https://dps.usc.edu/alerts/clery/			

ACKNOWLEDGMENTS

The above statements reflect the essential and non-essential functions as necessary to describe the principle contents of the job. They are not intended to be a complete statement of all work requirements or duties that may be required of the position. I understand that I may be asked to perform other duties as assigned. USC reserves the right to add or change duties at any time.

The University of Southern California is an Equal Opportunity Employer. USC prohibits discrimination on any basis protected under federal, state, or local law, regulation, or ordinance or university policies. All employment decisions are based on individual qualifications and business need.

I acknowledge receipt of this job description and its associated physical requirements. I have read and understand the job description and job requirements and agree to abide by their contents. I realize that duties may be requested of me that are not specifically stated herein. I understand that I will be expected to adjust to potential fluctuations in work volume. I understand that, if I have any questions about the essential functions or expectations of my position, my supervisor and/or HR partner are available to discuss them with me.

Print Employee Name

Signature

Date

Print Manager Name

Signature

Date

This job description describes the general nature and level of work required by the position. It is not intended to be an all-inclusive list of qualifications, skills, duties, responsibilities or working conditions of the job. The job description is subject to change with or without notice, and Management reserves the right to add, modify or remove any qualification or duty. Nothing in this job description changes the existing at-will employment relationship between the university and the employee occupying the position.