Jazz Pharmaceuticals.	FORM	
Document Number	Name	Version
JV-QDOC-203141	Global Job Description Template	1.0

Job Description

JOB TITLE:	TMF Operations Intern	COUNTRY:	United	States
LOCATION:	Remote GCDO Business and Technology Capabilities	ALT LOCATION:	N/A	
DEPT NAME:		DEPT NO:	21600	
REPORTS TO:	Chloe Bovis	FLSA STATUS: WORK LOCATION TYPE:	Exemp	t
JOB TYPE:	Temporary			Home Based
IS POSITION SU	PERVISORY? Choose Drop Down	BUDGET NUMBER	:	Enter Budget Number
% OF TRAVEL: <10% JOB LEVEL: Grade 5 Lead Coordinator Executive Assistant Associate Analyst		TRAVEL TYPE:		US Only
		DATE WRITTEN/RE	EVISED	17/01/2024

Brief Description:

The TMF Operations Intern (TOI) supports non-complex clinical trial and TMF activities in support of the GCDO Business & Technology Department. The TOI works closely with GCDO Business & Technology personnel to ensure department activities are conducted according to required activities and tasks delegated. As appropriate, they will identify issues and escalate to GCDO Business & Technology personnel. The core duties and responsibilities of the TOI are delineated below.

Essential Functions/Responsibilities

- General administrative support to the GCDO Business & Technology Department
- Manages eTMF content
- Perform Quality Control (QC) activities for TMF submissions in Jazz Clinical Vault (JV eTMF), in accordance with documented process
- Run eTMF metrics in Jazz Clinical Vault (JV eTMF), in accordance with documented process
- Assist in generating and managing trial dashboards for eTMF metrics
- General administrate support of GCDO Business & Technology department storage platforms (e.g., SharePoint, etc.)
- Assist in tracking of trial activities which may include but are not limited to study lists, QC review, FAQs, etc.
- Routinely participates in department and/or clinical trial team meetings and participate in collaborative efforts (e.g., departmental initiatives, etc.)

Required Knowledge, Skills, and Abilities

- Matriculated in a Bachelor's degree program
- No industry experience required
- Exposure to in Good Clinical Practices (GCP) regulations and Standard Operating Procedures preferred
- Understanding of study phases and general knowledge of how they apply to clinical development is preferred
- Knowledge of Word, Excel, and PowerPoint and shared platforms (e.g., SharePoint, Box, Smartsheets)
- Strong verbal and written communication skills required

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Signed: _____

Position Holder, Position Title

Date: _____

Signed: ______ Line Manager, Line Manager Title

Date: _____