Example of Clinical Director Job Description



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Our innovative and growing company is looking for a clinical director. Thank you in advance for taking a look at the list of responsibilities and qualifications. We look forward to reviewing your resume.

Responsibilities for clinical director

- Ensure that all pipelines and weekly reports are consistently completed and submitted to the appropriate stakeholders, including updates/written reports as required by Sheridan leadership
- Work collaboratively with the Vice President of Clinical Recruitment on all sourcing and practice recruitment activities and challenges
- Proactively make recommendations for improvements where opportunities and challenges exist, including sourcing and selection
- Engage in on-site practice profiles and development of interview panels for effective and legal hiring practices
- Adhere to the department's financial budget with all candidate activity and sourcing initiatives
- Proactively pursue new candidate sourcing strategies for candidate identification
- Ensure where vacancies exist that accurate candidate requirements, practice profiles and strategy action plans are in place
- Participate in medical specialty trade shows and conferences as needed
- Manage and control the development work in the Quality Measures team
- Lead, build and deliver automated sets of clinical measures that provide insights into performance and clinical outcomes

Qualifications for clinical director

• Drives the development of new services, with an emphasis on niche areas of

- Responsible for imaging equipment maintenance, physics inspections and adherence to all regulatory guidelines to ensure American College of Radiology (ACR) and TDSHS compliance
- Works with the radiologists, physician leaders and executive team to ensure that the department tools and systems promote high levels of patient care and staff productivity
- Familiar with a variety of imaging concepts, practices and procedures
- 10+ years of experience in dialysis preferably in all areas (acute, chronic, PD)
- Experience with protocol writing, IRB submissions, planning and executing studies according to GCP and SOPs