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**Regulatory affairs specialist -Design History file-Medical devices**

PharmEng has immediate opening for a Regulatory affairs associate with experience in medical devices

**Essential Job Requirements:**

EDUCATION Bachelor's degree preferably in Science or technology

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EXPERIENCE Need to have: previous experience in **submitting medical device regulatory filing**  
Excellent technical writing skills for writing design history files with minimum of 5-7 years of experience  
Must know FDA regulations and must have experience working with **medical devices**

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REQUIRED SKILLS Advanced thinking and analytical skills  
Ability to select the best course of action depending on the organization needs  
Ability to make decisions, act upon them with some direction and accept accountability for outcome of those decisions  
Ability to maintain confidential information at appropriate levels within the organization

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PREFERRED SKILLS RAC-Regulatory affairs Certification Completed

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OTHER REQUIREMENTS Canadian or USA citizen  
Valid Driving license and Passport  
Ability to Travel

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QUALIFIED CANDIDATES  
SUBMIT RESUME TO [humanresources@pharmeng.com](mailto:humanresources@pharmeng.com)

Only those candidates selected will be contacted for an interview.