

# Psychedelics Drug Development Startup Seeking Experienced Clinical Trials Coordinator

DELOS is dedicated to rigorously researching and developing psychedelic medicines with the aim of producing commercially available treatments that are safe, effective, and affordable.

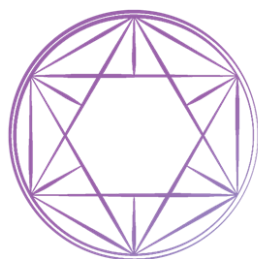
## Job Description

### Position Summary:

Delos is hiring for a clinical trials coordinator position which will involve work with clinical trials research on scheduled substances like LSD and 5-MeO-DMT. The clinical trials to be conducted are related to LSD microdosing for treatment of depression and obesity along with full-dose 5-MeO-DMT for treatment of depression. The endpoints for these studies will be opening a new drug application with the FDA to create marketable psychedelics products that could benefit millions. Prior experience with conducting clinical trials along with FDA and DEA requirements will be essential, while interest in therapeutic use of psychedelics is preferred.

### Key Responsibilities:

Responsibilities will include protocol development; achieving IRB, state, and federal approval; running phase 1-3 clinical trials up to the NDA level; assuring adherence of study operating procedures and protocols when trials are being conducted; participation in preparing study reports, journal publications, and presentations; maintaining confidential, accurate, and detailed records consistent with high-quality research; assisting with recruitment, consent, and randomization of participants into trials, plan daily activities with the team, monitor activities to ensure completion of tasks within the specified period, oversee data entry, and participate in regular meetings with the other investigators.



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[www.delos-research.com](http://www.delos-research.com)

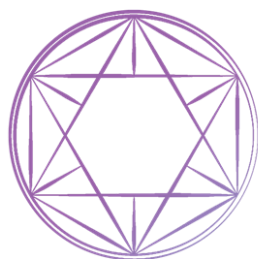
### **Essential:**

- Formal education in medicine or a medically-related profession.
- Experience with procedures and FDA, state, and DEA requirements for running at least phase 1 clinical trials.
- 3-5 years working with clinical trials studies.
- Experience in drug development clinical study design, cost estimates, and execution.
- Experience in collaboration with FDA project managers and regulatory officers.

### **Desirable:**

- Education in medicine or a medically-related profession from a top school.
- 10+ years working in clinical trials research up to phase 3 and NDA status.
- Interest in therapeutic use of psychedelic substances.
- MD or Ph.D. degree in a medical or medically-related profession.
- Has worked with scheduled substances (especially schedule 1 substances) in clinical trials previously.
- San Francisco Bay Area resident.

**Qualified applicants, please contact Richard Knowles, PhD at**  
[Richard.Knowles@delospyche.org](mailto:Richard.Knowles@delospyche.org)



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