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David Zavala

1 Main Street, New Cityland, CA 91010
Home: 000-000-0000 | Cell: (555) 322-7337
example-email@example.com

Professional Summary

Creative Clinical Research Associate with extensive project experience from concept to development. Talents include in-depth knowledge of ICH guidelines SAE reporting and GCP auditing. Integral team-player with excellent communication skills; fluent in French.

Core Qualifications

- Serious Adverse Event (SAE) reporting
- Project development
- Excellent interpersonal and communication skills
- Data management
- Recruitment and screening
- Accounts Payable / Receivable
- Project management
- Detailed documentation

Experience

Clinical Research Associate   1/1/2012 - Current
Green Care International   New Cityland, CA

- Monitor subject follow-up from 3-year clinical trial.
- Coordinate laboratory samples and compile data.
- Collaborate with medical professionals to determine changes in subjects' conditions.

Clinical Research Associate   2/1/2010 - 1/1/2012
Tanlam International   New Cityland, CA

- Managed Geo-nix project and oversaw team of 4 Research Associates.
- Maintained SOP's and Good Clinical Practice in compliance with ICH guidelines.
- Scheduled and led team study meetings; reported findings to Lead CRA.

Clinical Research Associate   8/1/2007 - 2/1/2010
Trioex Inc.   New Cityland, CA

- Screened and recruited subjects based on eligibility criteria for project participation.
- Served as point-of-contact for Clinical Research Professionals.
- Conducted pre-site visits and performed site evaluations.

Education

High School Diploma   XXXX - XXXX
New Parkland High School   New Parkland, CA

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There are plenty of opportunities to land a Clinical Research Associate job position, but it won't just be handed to you. Crafting a Clinical Research Associate resume that catches the attention of hiring managers is paramount to getting the job, and LiveCareer is here to help you stand out from the competition.

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