

Expert Clinical Research Associate (eCRA)

Job ID
REQ-10080974

6月 25, 2026

Argentina

摘要

Planificación, ejecución e interpretación de la investigación de ensayos clínicos, actividades de recopilación de datos y operaciones clínicas. Puede interactuar con sitios de investigación, consultores clínicos, organizaciones de investigación por contrato y otros proveedores. Colabora con colegas médicos/ clínicos del país, equipos clínicos globales y dirige actividades para ejecutar y entregar los estudios asignados. Monitorea los datos de los pacientes y la información relacionada con el estudio relacionada con los sitios de estudio clínico y la participación en ensayos clínicos. Asegura que el investigador se adhiera a los protocolos de investigación, los requisitos reglamentarios y las buenas prácticas clínicas y proporciona información en el plan de validación de datos. Proporciona un monitoreo oportuno y preciso de los datos de los pacientes y la información relacionada con el estudio de los documentos de origen, los registros de investigación y las visitas al sitio cuando corresponda. Puede monitorear los sitios de estudio y la selección de las instalaciones de auditoría.

About the Role

Key Responsibilities

- Lead site relationship management to support successful execution of Phase I through Phase IV trials
- Serve as the frontline liaison between Novartis and assigned investigative sites
- Manage complex study sites according to monitoring plans, procedures, and regulatory requirements
- Conduct site initiation visits and ensure site staff are trained on trial requirements
- Perform ongoing onsite and remote monitoring to ensure compliance and data integrity
- Identify site risks, process gaps, and improvement opportunities to support trial quality
- Promote a strong compliance culture focused on patient safety and ethical trial conduct
- Partner with cross-functional stakeholders to support recruitment, site development, and data quality
- Support audit and inspection readiness, including timely implementation of corrective actions
- Mentor junior Clinical Research Associates and contribute to innovative monitoring practices

Essential Requirements

- Degree in a scientific or healthcare discipline or equivalent relevant industry experience
- Minimum four years of experience in clinical monitoring and site management within the pharmaceutical industry
- Strong knowledge of clinical trial processes and drug development lifecycle

- In-depth understanding of international regulations including Good Clinical Practice and regulatory requirements
- Proven ability to independently manage complex clinical trial sites and monitoring activities
- Strong risk identification and issue management capabilities with a proactive, solutions-oriented approach
- Excellent communication, stakeholder engagement, and influencing skills across cross-functional teams
- Fluency in written and spoken English and local language

Desirable Requirements

- Field monitoring experience across complex or innovative clinical trial designs
- Experience supporting or mentoring junior Clinical Research Associates in a global environment

Why Novartis: Helping people with disease and their families takes more than innovative science. It takes a community of smart, passionate people like you. Collaborating, supporting and inspiring each other. Combining to achieve breakthroughs that change patients' lives. Ready to create a brighter future together? <https://www.novartis.com/about/strategy/people-and-culture>

Benefits and Rewards: Learn about all the ways we'll help you thrive personally and professionally. [Read our handbook \(PDF 30 MB\)](#)

部门
Development

Business Unit
Development

地点
Argentina

站点
Ramallo (Argentina)

Company / Legal Entity
AR01 (FCRS = AR001) Novartis Argentina S.A.

Functional Area
Research & Development

Job Type
Full time

Employment Type
Regular

Shift Work
No

```
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false }, sources: { options: {}, startTime: 0 }, disableUserCache: "true", plugins: {}, sources: { options:
{}}, ui: { showCCButton: false, settings: { showQualityMenu: true, showSpeedMenu:
false }, components: { fullscreen: { disableDoubleClick: false } }, uiComponents: [ { presets:
['Playback', 'Live'], area: 'BottomBarRightControls', replaceComponent: 'Fullscreen', get:
kPlayer.ui.components.Remove } ] } }; // Check and add plugins only if they exist if
(kPlayer.plugins["download"]) { config.plugins.download = { disable: true }; } if
(kPlayer.plugins["transcript"]) { config.plugins["playkit-js-transcript"] = { position: "right", // Default:
bottom;('left', 'right', 'top', 'bottom') to enable transcript. expandMode: "over", // Default:
alongside;('alongside', 'hidden', 'over') expandOnFirstPlay: false, showTime: true, downloadDisabled:
false, printDisabled: false, disable: true }; } if (kPlayer.plugins["preventSeek"]) {
config.plugins.preventSeek = { preventSeekForward: false, preventSeek: false }; }
config.plugins.floating = { disable: true }; if (kPlayer.plugins["navigation"]) { config.plugins.navigation =
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(kPlayer.plugins["moderation"]) { config.plugins['playkit-js-moderation'] = { disable: true }; } if
(kPlayer.plugins["info"]) { config.plugins['playkit-js-info'] = { disable: true }; } if
(kPlayer.plugins["share"]) { config.plugins.share = { disable: true }; } config.ui.uiComponents = []; if
(kPlayer.plugins["googleAnalytics"]) { config.plugins.googleTagManager = {};
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config.plugins.googleTagManager.containerId = 'GTM-57RJQ5';
config.plugins.googleTagManager.customEventsTracking.custom = [];
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false, playlistEvents: false, castEvents: false } }; }
```

```
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window.kalturaPlayerVideos = window.kalturaPlayerVideos || []; try { var kalturaPlayer =
kPlayer.setup(config); // Add the player to the global array.
window.kalturaPlayerVideos.push(kalturaPlayer); // Load the Player for other media.
kalturaPlayer.loadMedia({entryId: "1_dgfvmafo"}); } catch (e) { console.error(e.message) }
```

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