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Oracle Life Sciences AI Agents

Oracle Life Sciences AI Agents (Life Sciences AI Agents) is a web-based, unified workspace for all users who design, plan, conduct, and monitor clinical trials across their lifecycle.

- Overview
  Life Sciences AI Agents provides centralized trial planning in a single, intelligent platform that eliminates redundant manual tasks, connects operational and clinical data, and provides AI-driven insights, risk scoring, and real-time monitoring to help you make faster, more confident decisions across the entire clinical trial lifecycle.
- New in this release
  This release of Life Sciences AI Agents provides multiple features for designing, planning, conducting, and monitoring clinical trials across their lifecycle.

Overview

Life Sciences AI Agents provides centralized trial planning in a single, intelligent platform that eliminates redundant manual tasks, connects operational and clinical data, and provides AI-driven insights, risk scoring, and real-time monitoring to help you make faster, more confident decisions across the entire clinical trial lifecycle.

Life Sciences AI Agents unifies the processes from separate source Oracle Life Sciences applications, which enable you to complete most of your trial oversight and decision-making workflows within the application.

The source Oracle Life Sciences applications can include:

- Oracle Life Sciences Clinical One Cloud Service
- Clinical Trial Management System (CTMS)
- Oracle Life Sciences Site Activate
- Oracle Life Sciences Site Select
- Electronic Trial Master File (eTMF)
- Portfolio and trial health monitoring
  Clinical operation leads and executives can use Life Sciences AI Agents to view real-time portfolio health across studies, sites, and countries with AI agents highlighting emerging risks in enrollment, timelines, or quality metrics. AI-generated alerts and suggested actions enable proactive intervention and resource optimization across programs.
- Trial categories and data integration
  There are two categories of trials in Life Sciences AI Agents: Ongoing Trials and Trials in Planning. How you integrate your trial data into Life Sciences AI Agents determines how the trials are categorized.
- Trial planning
  Study managers and clinical operations leads can use Life Sciences AI Agents to extract protocol elements and generate protocol-driven plans with budgets, milestones, enrollment assumptions, and ranked country and site scenarios.

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- User roles
  There are two user roles in Life Sciences AI Agents that determine your feature access: Administrator and Trials.

Portfolio and trial health monitoring

Clinical operation leads and executives can use Life Sciences AI Agents to view real-time portfolio health across studies, sites, and countries with AI agents highlighting emerging risks in enrollment, timelines, or quality metrics. AI-generated alerts and suggested actions enable proactive intervention and resource optimization across programs.

Trial categories and data integration

There are two categories of trials in Life Sciences AI Agents: Ongoing Trials and Trials in Planning. How you integrate your trial data into Life Sciences AI Agents determines how the trials are categorized.

There are two methods for integrating trial data into Life Sciences AI Agents. You can collaborate with Oracle to import existing trial data from an Oracle Life Sciences source application, such as Clinical One Cloud Service or CTMS. Or, you can manually upload protocols for new trials that are not yet active.

- **Ongoing trials**—Active trials that have had their data onboarded into Life Sciences AI Agents by Oracle from a source application.

  Ongoing Trials are active clinical trials with ongoing or completed patient enrollment and data collection. They can be at any stage of the clinical trial lifecycle and are supported in Life Sciences AI Agents through the provision of AI-driven insights and summaries.
- **Trials in Planning**—Inactive trials whose protocols you manually uploaded into Life Sciences AI Agents.

  Trials in Planning are not yet active, with all or some features (such as protocols, operational plans, and site identification) that are still being developed or refined. The trial data is in the process of being reviewed and validated.

Depending on your permissions, you may have access to only Ongoing Trials or to Ongoing Trials and Trials in Planning. You cannot have access to just Trials in Planning.

Trial planning

Study managers and clinical operations leads can use Life Sciences AI Agents to extract protocol elements and generate protocol-driven plans with budgets, milestones, enrollment assumptions, and ranked country and site scenarios.

Trial planning in Life Sciences AI Agents supports scenario comparison, customer-specific data sources, and complex multi-arm protocols. This eliminates manual feasibility and planning work, accelerating study startup while ensuring plans remain aligned with enrollment and budget targets.

You can begin operational planning as soon as a protocol synopsis is drafted by uploading the synopsis including the Schedule of Assessments into Life Sciences AI Agents. After the protocol processing is complete, Life Sciences AI Agents provides three trial plans by default with different objectives: cost efficient, fastest, and a balance of both. You can review the trial plans, make any necessary adjustments, and then select the best plan for your trial.

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User roles

There are two user roles in Life Sciences AI Agents that determine your feature access: Administrator and Trials.

- **Administrator**—As a user with the Administrator role you can manage all Life Sciences AI Agents users and assign their access to trials. You can also assign the permission that controls access to Trials In Planning and allows users to add trial protocols into the application.
- **Trials**—As a user with the Trials role you have access to all trials or a subset of trials. Optionally, this role gives you access to Trials In Planning and to add trial protocols into the application.

You can have the Administrator role, Trials role, or both. If you only have the Administrator role, you cannot access trial data.

New in this release

This release of Life Sciences AI Agents provides multiple features for designing, planning, conducting, and monitoring clinical trials across their lifecycle.

The application provides a Portfolio Health overview, centralized trials list, and a centralized alerts list for managing your trial data. A user management interface enables administrators to manage user accounts and grant trial access. Single sign-on (SSO) authentication, a navigation menu in the interface for seamless access to source application Home pages, and deep linking to source application records simplifies your access to Life Sciences AI Agents and between the application and any source applications without requiring re-authentication.

- Portfolio Health overview
  The Portfolio Health overview serves as the single-entry hub for portfolio oversight, enabling efficient monitoring, navigation, and decision-making for all Ongoing Trials.
- Trials list
  The Trials list serves as the central navigation hub for all the trials you have access to, and provides easy navigation between the Trials list and trial details.
- Alerts list
  The Alerts list serves as a central repository for all the alerts in the Ongoing Trials you have access to, and provides easy navigation between the Alerts list and alert details. Alerts are AI-generated notifications that provide important information or identify potential issues for the Ongoing Trials you can access.
- User management
  As a Life Sciences AI Agents user with administrator rights, you can manage user accounts and assign roles, trial access, and the permission that allows users to view Trials In Planning and add new trials.
- Single sign-on (SSO) authentication
  Life Sciences AI Agents uses single sign-on (SSO) authentication that allows you to use one set of credentials to sign in and to seamlessly navigate between the application and any onboarded source Oracle Life Sciences applications without needing to sign in to each one separately.
- Navigate to source application Home pages
  Life Sciences AI Agents provides a navigation menu in the user interface for seamless linking to the Home page in a source Oracle Life Sciences application.

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- Deep linking to source application records
  Life Sciences AI Agents provides deep linking directly from actionable insights, recommendations, or data anomalies to the relevant record in a source Oracle Life Sciences application.

Portfolio Health overview

The Portfolio Health overview serves as the single-entry hub for portfolio oversight, enabling efficient monitoring, navigation, and decision-making for all Ongoing Trials.

You can use the Portfolio Health overview to view the overall health of all the Ongoing Trials you have access to. This facilitates efficient oversight and management of a customer's clinical trial portfolio, enabling stakeholders to navigate and respond to changing circumstances effectively.

The Portfolio Health overview displays key performance indicators (KPIs), a health summary, an alerts list, and trial data in graph format that together provide:

- Portfolio-level visibility, unified analytics and insights, and seamless navigation across studies and source applications.
- Consolidated cross-study information and direct access to trial-level and application-level resources.
- Summary data of the overall health of the trials in the portfolio.

The centralized Portfolio Health overview serves as your Home page that opens by default when you sign into Life Sciences AI Agents.

- Portfolio health key performance indicators (KPIs)
  The portfolio health key performance indicators (KPIs) provide standardized metrics that measure study performance consistency across a customer's portfolio for the past 90 days to identify areas of operational risk.
- Portfolio health summary
  The portfolio health summary provides an AI-generated text summary of all Ongoing Trials in a customer's portfolio. The summary highlights top risks, study-level callouts with direct (deep) links, and cross-portfolio trends impacting efficiency and quality.
- Portfolio alerts
  The portfolio alerts provide an AI-generated list of the highest priority alerts (for example, insights and recommendations) for all Ongoing Trials in a customer's portfolio.
- Portfolio health graphs
  The portfolio health graphs display data for all Ongoing Trials in the portfolio in multiple selectable views, including Trials by Status, Trials by Phase, Trials by Therapeutic Area, and Enrollment by Region. Each graph format includes a timestamp. Hovering over the graphs displays statistic details.

Portfolio health key performance indicators (KPIs)

The portfolio health key performance indicators (KPIs) provide standardized metrics that measure study performance consistency across a customer's portfolio for the past 90 days to identify areas of operational risk.

The KPIs enable clinical operation leads to monitor study performance, identify potential risks, and make informed decisions to mitigate those risks.

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The portfolio health KPIs appear across the top of the centralized Portfolio Health overview page below the page title. They are refreshed every 24 hours at 2 am UTC and are only calculated and shown if there is enough data for each.

> **Note**
>
> You can hover over any KPI to see details about how its value was calculated, including the minimum and maximum values, the number of trials used, and the calculation date.

The KPIs can include:

- **Needs Attention**—Count of trials with alerts (insights). The count depends on the number of active alerts.
- **Active Trials**—Number of trials users have access to in the Life Sciences AI Agents application.
- **Site Activation**—Percent of activated sites that have met their activation target dates. This metric is a measure of start-up efficiency. A high variance signals bottlenecks.
- **Enrollment vs Plan**—Actual enrollment divided by the planned enrollment (uses the enrollment forecast). This metric shows if the portfolio is meeting recruitment goals.
- **Data Entry**—Percentage of expected CRFs completed (regardless of query status). This metric serves as an early warning of operational lag. Low data entry completeness correlates to monitoring backlog and slow data entry timeliness.
- **Deviation Rate**—Number of protocol deviations reported per 100 subjects across the portfolio or subset of trials. This metric represents the universal compliance indicator.
- **SAE Rate**—Serious adverse event rate per 100 subjects enrolled. This is a key portfolio safety metric that normalizes by exposure (compares safety risks across different studies).
- **TMF Completeness**—Percentage of expected documents collected. This metric provides leadership a sense of inspection readiness across the portfolio.
- **Trials on Track** —Percentage of trials hitting enrollment targets within ±10% of plan. This executive indicator shows the percentage of studies that are progressing as planned.
- **Dropout Rate**—Percentage of enrolled subjects that have dropped out or were terminated early. Appears only if you are not using a source application.
- **Data Entry Timeliness**—Average number of days from subject visit dates to when the data is entered into eCRFs in EDC systems. This metric provides an early warning signal for downstream delays in cleaning, reporting, and analysis. Appears only if you are not using a source application.

Portfolio health summary

The portfolio health summary provides an AI-generated text summary of all Ongoing Trials in a customer's portfolio. The summary highlights top risks, study-level callouts with direct (deep) links, and cross-portfolio trends impacting efficiency and quality.

The portfolio health summary provides executives, study managers, or clinical operations leads with access to all trials a consolidated view of the portfolio health by surfacing the most critical risks, trends, and study-level callouts.

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The portfolio health summary appears below the KPIs on the centralized Portfolio Health Overview page.

> **Note**
>
> The portfolio health summary appears on the Portfolio Health overview page only if you have access to all trials.

Portfolio alerts

The portfolio alerts provide an AI-generated list of the highest priority alerts (for example, insights and recommendations) for all Ongoing Trials in a customer's portfolio.

The portfolio alerts provide a concise and regularly updated view of the most critical issues in a customer's clinical trial portfolio, enabling you to quickly respond to emerging issues or trends.

The portfolio alerts appear at the bottom left on the centralized Portfolio Health overview page.

The portfolio alerts:

- Show one row per trial and up to three top ranked issues for each trial (each alert provides a link to the Alert details).
- Prioritize trials based on risk severity (Critical, High, Moderate, or Low). The default sort of the trials in the list is by risk severity.
- Refresh every 24 hours at 2 am UTC. If no trials need attention, a "No trials currently require attention." message appears in the Alerts section.

From the portfolio alerts list, you can:

- Open the Alerts list to view the list of alerts for all the Ongoing Trials you have access to.
- Open the Alerts list filtered for the study where the alert originated.
- Open the details for an alert. For every alert you can view (if applicable) a proposed protocol amendment and any supporting information, the AI reasoning steps for the alert, and the history of the alert. You can also snooze or dismiss an alert.
- Open the details for the study where the alert originated.

Portfolio health graphs

The portfolio health graphs display data for all Ongoing Trials in the portfolio in multiple selectable views, including Trials by Status, Trials by Phase, Trials by Therapeutic Area, and Enrollment by Region. Each graph format includes a timestamp. Hovering over the graphs displays statistic details.

The graphs enable you to tailor the visualization of your trial data to suit your analytical requirements, potentially enhancing your ability to understand and act upon the insights provided by the data.

The portfolio health graphs appear at the bottom right on the centralized Portfolio Health overview page.

- **Trials by Status (bar graph)**—Shows the number of Ongoing and In Progress trials.

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- **Trials by Phase (pie graph)**—Shows the percentage of trials in each clinical trial phase (I to IV).
- **Trials by Therapeutic Area (bar graph)**—Shows the number of trials by therapeutic area (for example, Cardiology/Vascular Diseases, Dermatology, and Endocrinology).
- **Enrollment by Region (bar graph)**—Shows the number of subjects enrolled by country.

Trials list

The Trials list serves as the central navigation hub for all the trials you have access to, and provides easy navigation between the Trials list and trial details.

You can filter the list by the operational lifecycle stage: All Trials, Ongoing Trials, Other Trials (trials that are completed, suspended, or terminated), and if you have access, to Trials in Planning. You can use the Trials list as a convenient and organized way to access and navigate through the trials you can access.

You can select the Trials tab at any time to view the complete list of trials you can access. The first time you open the Trials list it defaults to the All Trials view.

Use the Trials list to:

- View the status, study name, study title, phase, indication and therapeutic area, and the dates the trials were created or last updated.
- Search for and sort the trials in the list.
- Filter the list based on Status, Therapeutic Area, Phase, and Indication to easily locate the list of relevant trials for your role.
- Navigate to a trial details page to view details of a specific trial and then return to the Trials list while keeping the trial list filters in place.
- Add a new trial. If you have the required permission (access to Trials in Planning), you can add new trials into Life Sciences AI Agents (upload the trial protocols, review and validate the data, and then choose from three AI-generated trial plans) or modify existing protocols by uploading a different version and regenerating existing trial plans.
- Export all trial data (filtered for All Trials, Ongoing Trials, Other Trials, or Trials in Planning) to a CSV file.
- Trial details
  The trial details provide the trial name, trial key performance indicators (KPIs), trial health summary and insights, a trial alerts list, and trial data graphs for a specific trial on a separate page.
- Trial health key performance indicators (KPIs)
  The trial health key performance indicators (KPIs) provide standardized metrics that measure trial performance consistency for the past 90 days to identify areas of operational risk.
- Trial health summary
  The trial health summary provides an AI-generated text summary for an active Ongoing Trial that highlights the top risks impacting efficiency, quality, and trial success. The summary provides you with a consolidated view of the trial health by surfacing the most critical risks, trends, and study-level callouts.
- Trial alerts
  The trial alerts provide a prioritized list of AI-generated alerts for the specific trial.

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Trial details

The trial details provide the trial name, trial key performance indicators (KPIs), trial health summary and insights, a trial alerts list, and trial data graphs for a specific trial on a separate page.

The trial details provide a centralized location for you to access and review critical information about a specific clinical trial, enabling informed decision-making and effective trial management.

Trial health key performance indicators (KPIs)

The trial health key performance indicators (KPIs) provide standardized metrics that measure trial performance consistency for the past 90 days to identify areas of operational risk.

The trial health KPIs provide standardized metrics and consistent measurements over time, which facilitates data-driven decision-making and supports the successful execution of clinical trials.

The trial health KPIs appear across the top of a trial details page below the page title. They are refreshed every 24 hours at 2 am UTC and are only calculated and shown if there is enough data for each.

> **Note**
>
> You can hover over any KPI to see details about how its value was calculated, including the minimum and maximum values and the calculation date.

The KPIs can include:

- **Enrollment Progress**—Actual enrollment accrued in CTMS at the protocol level. This metric shows the total number of subjects enrolled for the trial at any time.
- **Enrollment vs Plan**—Actual enrollment divided by the planned enrollment (uses the enrollment forecast). This metric shows if the trial is meeting its recruitment goals.
- **Site Activation**—Percentage of activated sites that have met their activation target dates. This metric is a measure of start-up efficiency. A high variance signals bottlenecks.
- **Data Entry**—Percentage of expected CRFs completed (regardless of query status). This metric serves as an early warning of operational lag. Low data entry completeness correlates to monitoring backlog and slow data entry timeliness.
- **Deviation Rate**—Number of protocol deviations reported per 100 subjects across the trial. This metric represents a universal compliance indicator.
- **SAE Rate**—Serious adverse event rate per 100 subjects enrolled. This is a key portfolio safety metric that normalizes by exposure (compares safety risks across different studies).
- **Query Rate**—Number of queries raised at the site divided by the number of items on any started eCRF (for example, for data that has been collected so far).
- **Requery Rate**—Number of re-opened queries divided by the total number of queries. This metric shows when the site's initial response was insufficient, incorrect, or generated a new query. This can occur if:

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  - The initial site response did not close a query, resulting in a "re-query" to the site (the query state changed from Answered to Opened).
  - The initial site action (data change) did not close a query, resulting in a "re-query" to the site (the query state changed from Opened to Answered to Opened).

The following trial KPIs may appear on the page if you are not using a supported source application:

- **Dropout Rate**—Percentage of enrolled subjects that have dropped out or were terminated early.
- **Data Entry Timeliness**—Average number of days from the date of a subject visit to the date the data is entered into an eCRF in EDC. This metric provides an early warning signal for downstream delays in cleaning, reporting, and analysis.

Trial health summary

The trial health summary provides an AI-generated text summary for an active Ongoing Trial that highlights the top risks impacting efficiency, quality, and trial success. The summary provides you with a consolidated view of the trial health by surfacing the most critical risks, trends, and study-level callouts.

Trial details provide executives, study managers, or clinical operations leads with a concise, AI-driven overview of the health of the clinical trial, enabling stakeholders to quickly identify potential issues and make informed decisions to mitigate risks and ensure trial success.

The trial health summary appears below the KPIs on a trial details page.

Trial alerts

The trial alerts provide a prioritized list of AI-generated alerts for the specific trial.

The trial alerts provide a concise and prioritized view of potential issues for a specific clinical trial, enabling you to quickly respond to emerging issues or trends.

The trial alerts appears at the bottom left on a trial details page.

The trial alerts:

- Show one row per alert and the top seven alerts for the trial.
- Are prioritized in order of risk severity (Critical, High, Moderate, or Low).
- Refresh every 24 hours at 2 am UTC.

From the trial alerts list, you can:

- Sort the alerts by Severity and Date Created.
- Navigate to the Alert details, where you can perform various actions on the alert as needed.
- Navigate to the Alerts list with filters preset to the trial.

Alerts list

The Alerts list serves as a central repository for all the alerts in the Ongoing Trials you have access to, and provides easy navigation between the Alerts list and alert details. Alerts are AI-

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generated notifications that provide important information or identify potential issues for the Ongoing Trials you can access.

You can use the Alerts list to efficiently manage and respond to alerts across multiple trials, streamlining your workflow and improving your ability to address potential issues or opportunities. By default, the alerts are prioritized based on risk severity (Critical, High, Moderate, or Low). You can view alerts details and provide feedback on every alert. You can also view the AI reasoning steps for an alert, view the history of the alert, snooze or dismiss alerts, and view details for the study where the alert originated.

Use the Alerts list to:

- Search for and sort the alerts in the list.
- Filter on Trials, Status, Severity, and Date Modified.
- Navigate to and view details of a specific alert and then return to the Alerts list while keeping the Alerts list filters in place.
- Navigate to the study details for the study where the alert originated.
- Alert details
  The alert details provide a summary of the issue and recommendation, the supporting information for any insight or recommendation that comes from the AI agent, and if the alert is a proposed protocol amendment, provides the details of the amendment with a summary, rationale, and justification.
- Alert feedback
  On every alert details page, you can provide real-time feedback on the alert to report whether the AI-generated insights and suggestions were helpful, correct, or actionable.

Alert details

The alert details provide a summary of the issue and recommendation, the supporting information for any insight or recommendation that comes from the AI agent, and if the alert is a proposed protocol amendment, provides the details of the amendment with a summary, rationale, and justification.

The alert details help you to understand the alert and its underlying reasoning, as well as any proposed changes or supporting information. This enables you to make informed decisions and take appropriate actions in response to the alert.

Use the alert details to:

- View the AI reasoning for the alert to understand how insights and recommendations were rationalized.
- View the history of the alert to see its chronological history. The history includes the date the alert was first detected and the actions taken on the alert including the date first created, severity changes, and status changes.
- Snooze the alert to pause its notifications to you. When you snooze an alert it is immediately removed from the Alerts list. The notifications for a snoozed alert appear to you again if the criticality of the alert increases at a later time or if you undo the snooze action. You snooze your own alerts, and you can view a list of the alerts your snoozed.
- Dismiss the alert to stop the alert notifications to you. When you dismiss an alert it is immediately removed from the Alerts list. The notification for a dismissed alert will not appear to you again unless you undo the dismiss action. You dismiss your own alerts, and you can view a list of the alerts you dismissed.

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- Deep link directly from an actionable insight, recommendation, or data anomaly to the relevant record in a source Oracle Life Sciences application.

The details for each alert open on a separate page.

Alert feedback

On every alert details page, you can provide real-time feedback on the alert to report whether the AI-generated insights and suggestions were helpful, correct, or actionable.

Life Sciences AI Agents uses your feedback to improve the quality, accuracy, and relevance of the AI-generated insights, to refine future reasoning, and to improve the AI-driven guidance.

Your feedback helps Life Sciences AI Agents to iteratively learn and improve, leading to better performance and more accurate insights over time. The alert feedback feature enables the application to adapt to your needs and preferences, ultimately providing more effective support for clinical trial management.

For every alert, you can:

- Select the thumbs up or thumbs down icon on an alert details page as your feedback. Life Sciences AI Agents logs your username, timestamp, Alert ID, and Trial ID.
- Change your feedback on an alert at any time.

User management

As a Life Sciences AI Agents user with administrator rights, you can manage user accounts and assign roles, trial access, and the permission that allows users to view Trials In Planning and add new trials.

This feature provides comprehensive control over users and their trial access, enabling administrators to ensure the system is used securely and efficiently, with users having the necessary access to perform their designated roles.

As an administrator, you can:

- Add, remove, and update users.
- View existing users and user details.
- Assign, remove, and update user roles and trial access. You can assign access to all trials or to only selected trials.

Single sign-on (SSO) authentication

Life Sciences AI Agents uses single sign-on (SSO) authentication that allows you to use one set of credentials to sign in and to seamlessly navigate between the application and any onboarded source Oracle Life Sciences applications without needing to sign in to each one separately.

SSO authentication simplifies access between Life Sciences AI Agents and any source Oracle Life Sciences applications making it easier for you to work efficiently and effectively.

You can navigate from Life Sciences AI Agents to a source Oracle Life Sciences application using the navigation menu or deep links without re-authenticating as long as there is an active session in the source application and you have a valid account in the source application.

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- Before you can sign in to Life Sciences AI Agents, you must have an Oracle Cloud (IDCS) account (created by an Oracle administrator) and be added as a Life Sciences AI Agents user.
- Life Sciences AI Agents uses secure verification with multi-factor authentication (MFA) to verify your identity when you activate your account and sign in. All sign ins to Life Sciences AI Agents and navigations to source Oracle Life Sciences applications are logged for auditability.
- Life Sciences AI Agents automatically signs you out of the system after the amount of continuous use or inactivity that is set in your Oracle Cloud account. You see a notification message when your inactivity timeout is about to occur; you can extend your session if necessary. After your session ends for either reason you must sign in again.
- Signing out of Life Sciences AI Agents terminates your Oracle Cloud account session. It does not sign you out of any source Oracle Life Sciences applications.

Navigate to source application Home pages

Life Sciences AI Agents provides a navigation menu in the user interface for seamless linking to the Home page in a source Oracle Life Sciences application.

The navigation menu simplifies your access to source Oracle Life Sciences applications to enhance productivity.

- The navigation menu shows a list of the source applications that are onboarded to Life Sciences AI Agents that you have access to.
- Selecting a source application navigates you to the Home page for the application, as long as you have a valid account.
- When you navigate to or from Life Sciences AI Agents to a source application, you are not required to re-authenticate as long as you have access to the source application and both applications have active sessions.

Deep linking to source application records

Life Sciences AI Agents provides deep linking directly from actionable insights, recommendations, or data anomalies to the relevant record in a source Oracle Life Sciences application.

Deep linking allows you to quickly investigate or take action on a record without manually searching for the record in one or more source application systems. This feature enables study managers or clinical operations leads to facilitate efficient and effective decision-making by providing direct access to relevant records in source Oracle Life Sciences applications.

- When you have a valid account in a source Oracle Life Sciences application, you can link from an insight, recommendation, or metric in Life Sciences AI Agents directly to the relevant page, form, or record in the source application with the context pre-filtered (for example, the subject, site, or query ID). You can only access the pages you have the rights to in the source application.
- When you select a deep link URL to open a record in source application, you are not required to re-authenticate as long as you have access to the source application and both applications have active sessions.

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