

Careers in Medical & Development

By applying your medical knowledge and skills at Novartis you have the opportunity to impact millions of lives around the world. It's one of the key reasons many of our colleagues joined from the clinical environment and together we are building the leading medicines company, powered by data and digital.

Global Drug Development

We in the Global Drug Development (GDD) organization oversees the development of new medicines discovered by our researchers and partners. We transform the lives of people by developing innovative and life-changing medicines.

To help lead the world in biopharmaceutical development, we have built our strategy on the Novartis strategic pillars: Delivering breakthrough innovation, going big on data and digital, building trust with society, embracing operational excellence, and unleashing the power of our people.

11 000

GDD associates globally

90+

Projects in clinical development

300+

Ongoing clinical trials in GDD

50+

Major submissions planned for lead & new indications (2020-24).

Meet the teams:

Global Development Operations

Global Development Operations (GDO) is the bridge that turns great science into great medicine. We provide clinical trial operations services to the divisional development teams, taking the compounds developed by our colleagues and test those compounds in clinical trials.

We have more than 5,000 colleagues working across more than 60 countries, working in specialized core functions for end-to-end trial operations management:

- Trial Management
- Trial Monitoring
- Data Operations
- Regulatory Writing and Submissions (RWS)
- Strategic Planning and Feasibility

And dedicated capabilities to drive continuous improvement:

- Transformative Business Operations
- Risk Management
- External Development Operations (EDO) for vendor management
- Strategy and Integration
- Clinical Innovation and Technology
- data42

These focus areas, combined with our use of advanced technology and data analytics allow us to make informed decisions, potentially resulting in new, life-changing therapies for patients globally.

Patient Safety

The Novartis Group's Chief Medical Office and Patient Safety organization is an integrated team which provides Novartis-wide oversight and governance of patient focused activities including Patient Safety, Pharmacovigilance, Device-vigilance as well as Ethical, Regulatory and Reputational issues. We support our company mission by further strengthening our focus on patients and providing patient-centric solutions and outcomes by:

- Establishing a single Chief Medical Office for Novartis for all patient related policy and standards
- Creating one single Safety/pharmacovigilance organization across all divisions and one harmonized Product Stewardship team
- Integrating all country level Pharmacovigilance and Device vigilance into a single organization
- Implementing and deploying the Novartis Patient Declaration across the Group

□

4 years with Novartis. From an intern to HR Analyst and Patient Safety Specialist. Novartis offers us the flexibility to drive our own career. There are many growth opportunities here, and innovation and thinking-outside-the-box mindset is actively encouraged, even when it comes to your career development.

Giselle, Patient Safety Specialist

Regulatory Affairs

Regulatory Affairs

Regulatory Affairs (RA) is a unified, diverse, and strategic company-wide team. We are passionate about delivering regulatory excellence and innovation beyond traditional boundaries to enable timely and sustainable access to superior products for patients worldwide.

Careers in RA offers us the ability to gain real breadth of experience across many therapeutic areas but also depth of experience and knowledge as well.

Technical Research & Development

We in the Technical Research & Development (TRD) team are responsible for developing innovative technical product designs and robust, scalable manufacturing processes.

We are responsible for the manufacture and supply of all pre-clinical and clinical study materials across all stages of clinical and pharmaceutical development. We add value by working with and enabling a broad spectrum of cross-divisional partners, including early Research teams (see Research) late phase Development and registration teams, our commercial manufacturing teams, (see Technical Operations) and our Generics development teams (Sandoz.com).

Given this huge breadth of areas covered by TRD, we as associates benefit from one of the strongest possible learning environments and professional, career development opportunities.

We have an impact on a variety of other Novartis functions too. As a team of 2500 associates we work across 4 continents and 8 countries, developing practically the entire Novartis portfolio, across all Disease areas, including cutting edge Cell and Gene therapies.

Portfolio Strategy and Management

We in the Portfolio Strategy and Management team provide innovative methods and systems that enable informed decision making across Development in order to advance the Science of Medicine and the Science of Operations. Additionally we help to identify and lead strategic initiatives and programs to drive innovation and new capabilities across the Development team.

Digital & Data

The newly created Data & Digital team within GDD aims to use the power of data and digital to enhance the value of our innovative and impactful medicines. For instance, by reducing the time it takes to develop medicines, and improving our processes, or to bring down costs which can be reinvested back into R&D.

A growing number of data science and digital technology solutions are already in play aiming to:

- Leverage all available data to generate insights about diseases, patients and treatments
- Deploy technologies to improve the patient experiences, help trial recruitment and retention and enhance the benefit of our drugs
- Embed tools e.g. predictive analytics to improve our end-to-end trial processes

Click to find out about [careers in Digital and Data](#):

Clinical Development & Analytics

Clinical Development & Analytics (CD&A) is an integrated global line function delivering clinical, biostatistical, advanced analytical and functional excellence in integrated drug development.

We are organized around the following 4 key areas:

- Therapeutic Areas (TAs) across the Development Units and Sandoz Biopharmaceuticals
- Analytics function comprised of the Biostatistics and Pharmacometrics groups
- China Development and Japan Clinical Development
- Central Team comprised of Clinical Sciences, Pediatric Centre of Excellence, PRO Centre of Excellence and Strategy & Operations groups

Why Development at Novartis?

At Novartis, we recognize career development stems from personal growth, and personal growth stems from experiences. We are 'unbossed' to drive our own personal growth, to nourish our curiosity, and to challenge, stretch and enhance our skills and knowledge.

GDD is one such team that allows us to truly grow. In addition to the innumerable [learning & mentoring opportunities](#) available, we have access to the:

- Global Development University
- New hire training programs
- New platform programs
- Disease specific lectures
- Capabilities building
- Talent and Inclusion development programs
- And access to the OneCollaborate platform that enables internal sabbaticals/rotations and project assignments across the wider R&D community.

CONEXTS

CONEXTS is a solutions partner that creates value for the larger enterprise through our highly specialized/ skilled teams. Within CONEXTS the Medical and Clinical Solutions team support internal business stakeholders with:

Medical Communications Services

We in the Medical Communication Service team provide end-to-end support Medical Affairs, Regulatory and Clinical groups internal to Novartis through publications and supporting scientific events and congresses. Expert support is provided across franchise and therapeutic areas.

Medical Information Services

By joining the Medical Information Services team you will provide support to the Medical Affairs team sat in the Commercial division of Novartis. We provide them with accurate information to support health care professionals (HCPs) treating patients with Novartis products. These range from preparing responses to medical information enquiries and global guidance documents, to supporting congress booths.

Post-Launch Scientific Support

The Post-Launch Scientific Support service provides end-to-end support to internal clients conducting medical affairs studies. Our services cover: clinical project management, medical writing, data management and biostatistics and statistical programming.

Real-World Evidence

The Real-World Evidence team supports Medical affairs and Medical Access teams in generating and analyzing real world evidence data about the safety and efficacy of

medicines in daily clinical use.

Why CONEXTS?

In addition to the thousands of training opportunities available, as part of smaller team there is greater visibility for your knowledge, skills and deliverables while working on valuable medical projects and services.

Find your life-changing career at Novartis

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