STEER Life

Your Partner for Potent & Difficult-to-Develop Generic FDFs

Changing the Way we Make and Take Medicines

Who we are

We are a specialised CRO/CDMO for pharma manufacturers and marketers

We offer:

- End-to-End Drug Development Services
- Dossier Licensing Opportunities
- Co-Development Opportunities

We focus on drug products going off-patent in the next 5-7 years: so that our partners can grow their product portfolio.

Our USP:

We gladly deliver complete technology transfer to our partners' manufacturing sites – or supply products as per the need.

Specialization

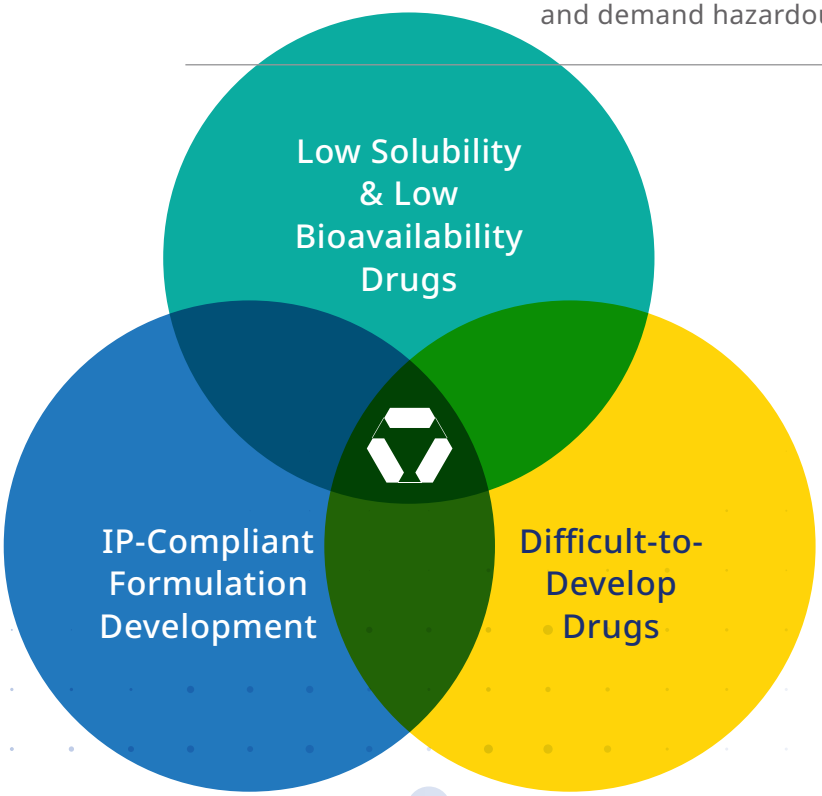
Oncology drugs, Anti-Retrovirals, Orphan drugs, High-Potent drugs, Implant-based drug products & any Difficult-to-Develop/Complex generics.

Core Strength

Unique integration of formulation expertise + know-how of equipment engineering.

More than just a CRO

The Problems We Solve
Poorly soluble drugs cannot be effectively formulated using conventional methods.
Oncology and Anti-Retroviral drugs require extremely precise manufacturing conditions that conventional granulation and batch processes cannot consistently deliver
Complex generics combining difficult-to-formulate APIs, bioavailability challenges, and multiple actives require integrated development expertise spanning both formulation science and advanced manufacturing technology.
Conventional methods cannot achieve the uniform dispersion and particle engineering needed for consistent drug performance and safety.
For poorly soluble APIs, conventional methods rely on the use of organic solvents which escalate manufacturing costs, inflate CAPEX and demand hazardous waste management.



Expertise in Niche Technologies

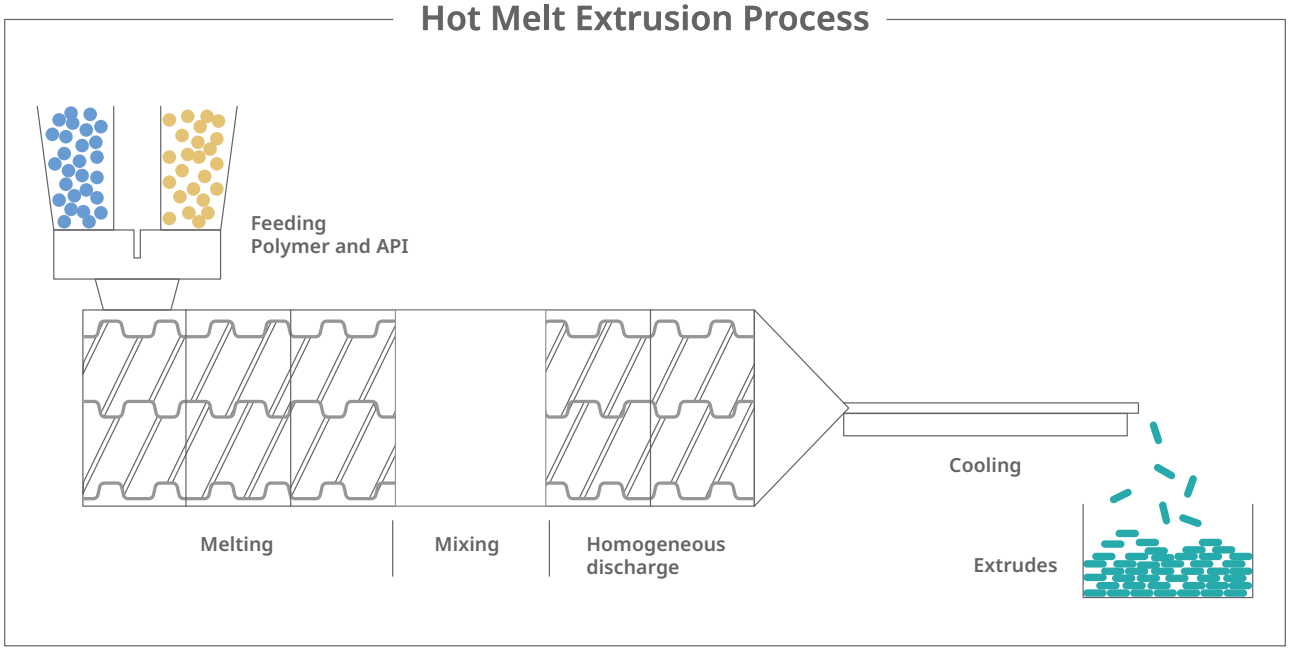
A. **Poorly Soluble Drugs:** Hot Melt Extrusion (PureMelt™)

B. **Complex Generic Drug Products:** Continuous Twin-Screw Granulation (Integraal™)

C. **Heat-sensitive & Shear-sensitive Drugs:** FragMelt™

D. **Effervescent Drug Products:** EXIMIOUS®

We enable you to achieve robust scalability with all technologies

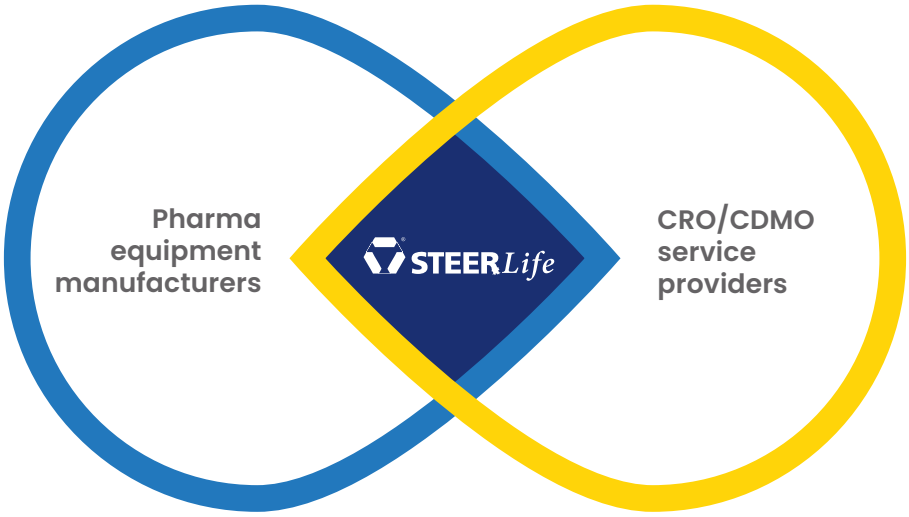


The unique integrated advantage

Traditional OEMs
excel at equipment engineering but lack formulation expertise.

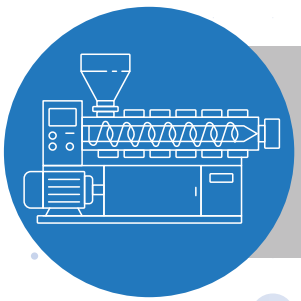
Traditional CDMOs
understand formulation development but lack deep equipment knowledge.

STEER*Life* combines both
with deep expertise on both sides.



a. Specialists in IP-Compliant Formulation Development
for BCS class II & IV molecules.

b. Leadership born in drug development
Decades of leading global R&D at industry giants like Viatris & Dr. Reddy's. With 80+ patents & publications and various novel manufacturing platforms developed, they don't just understand drug development, they've lived it.



STEER is among a handful of global players with an understanding of manufacturing technologies and their application to developing new drug products

What this Means for our Partners

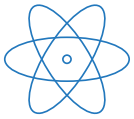
- Enable Local Manufacturing or Product Supply
- Overcome Complex Formulation Barriers
- Accelerate Time-to-Market with Integrated Development & Scale-Up
- Create Defendable Intellectual Property
- Achieve Sustainable, Solvent-Free Manufacturing
- De-Risk Scale-Up and Achieve Cost Parity

Scale-Up Assurance

We don’t just develop formulations, we transfer proven processes to commercial scale.

R&D formulations translated into manufacturing-ready processes

Technology transfer support ensuring success at your facility



Continuous manufacturing systems that maintain formula integrity and consistency

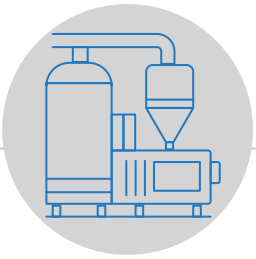
Compliance-ready scaled solutions ready for market deployment

Our Services

End-to-End CRO Solutions

R&D (or) Development:

- Feasibility trials
- Patent Landscaping
- Process Development
- Formulation Development / ASD (Amorphous Solid Dispersion) Development (Optimization)
- Analytical method development and validation



Scale-Up & Manufacturing:

- Technology Transfer & Scale-Up
- Execution of Exhibit batch campaigns for regulatory submissions (Pilot & Pivotal BE studies)
- Contract Manufacturing and Product Supply (FDFs)

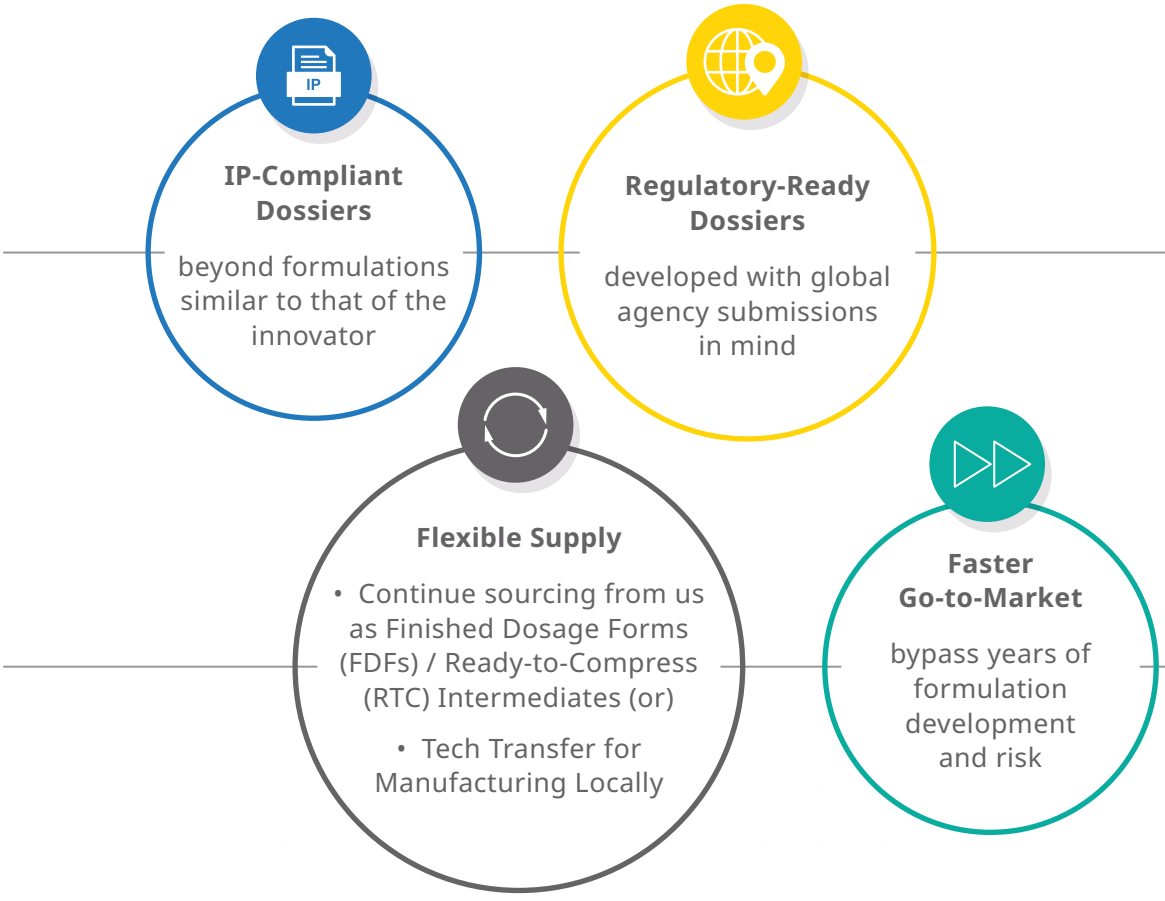
Other Integrated Services:

- Bioequivalence Studies – Coordination and Partner Selection
- Support for Dossier Data Compilation
- Support in API Sourcing
- RLD Procurement
- IP Support

Our Services
Dossier Licensing

Skip the development risk. License our ready dossiers and fast-track your path to market authorization.

Why License from STEER*Life*?



Flexible engagement models

Choose the partnership structure that best fits your needs:

End-to-End Drug development (CRO) solutions

We custom develop dossiers – end-to-end so that you can take them to any market globally. This can be followed by technology transfer for local/in-house manufacturing (or) direct product supply.

Dossier-Licensing Model (CDMO) solutions

Results in the partner licensing our ready dossiers, followed by either technology transfer (or) direct product supply.

Co-Development

Collaborative development with shared expertise

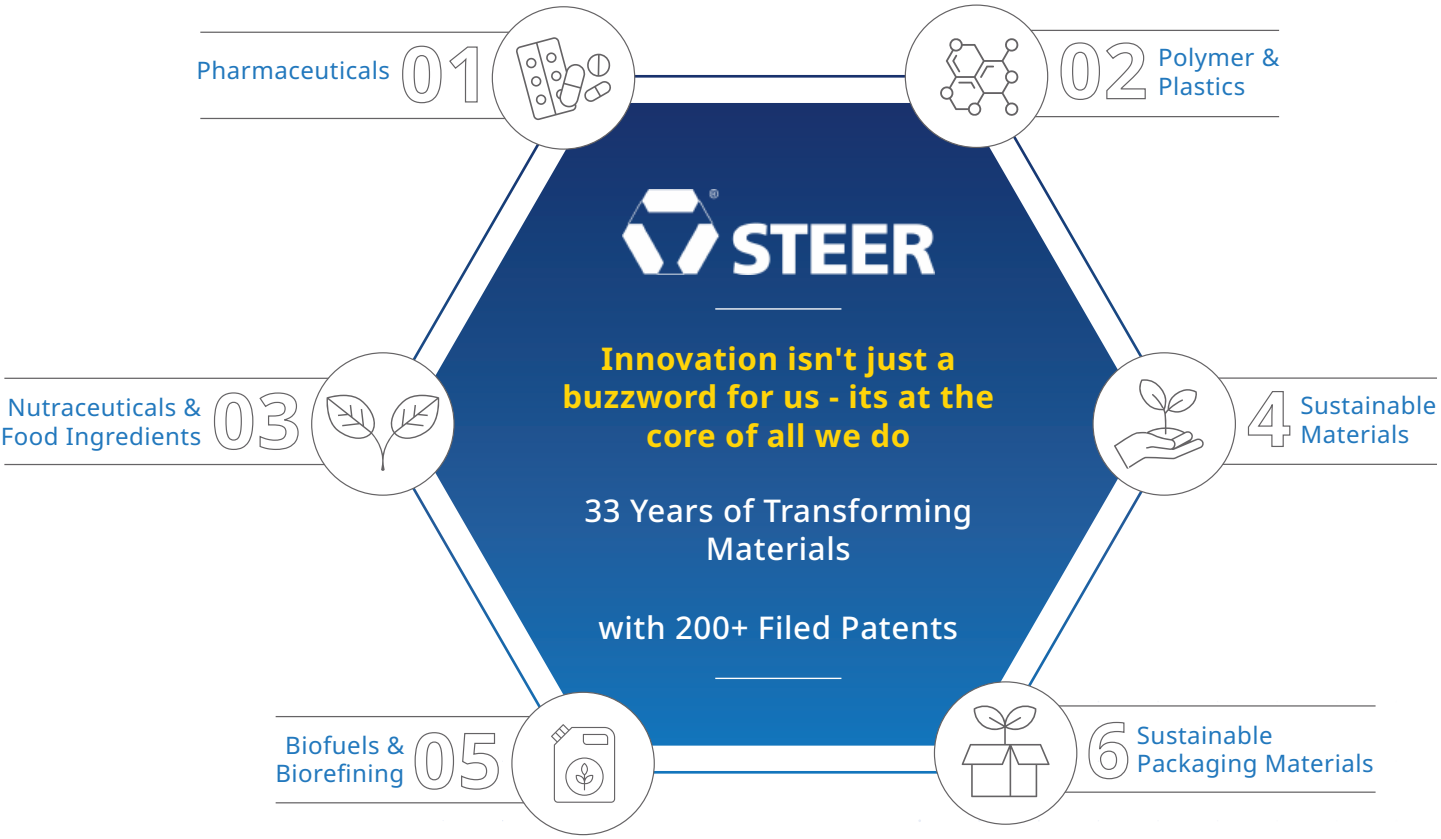
Complete Turnkey Solutions

Complete end-to-end solutions with development + technology transfer + equipment to enable our partners to manufacture locally for their markets

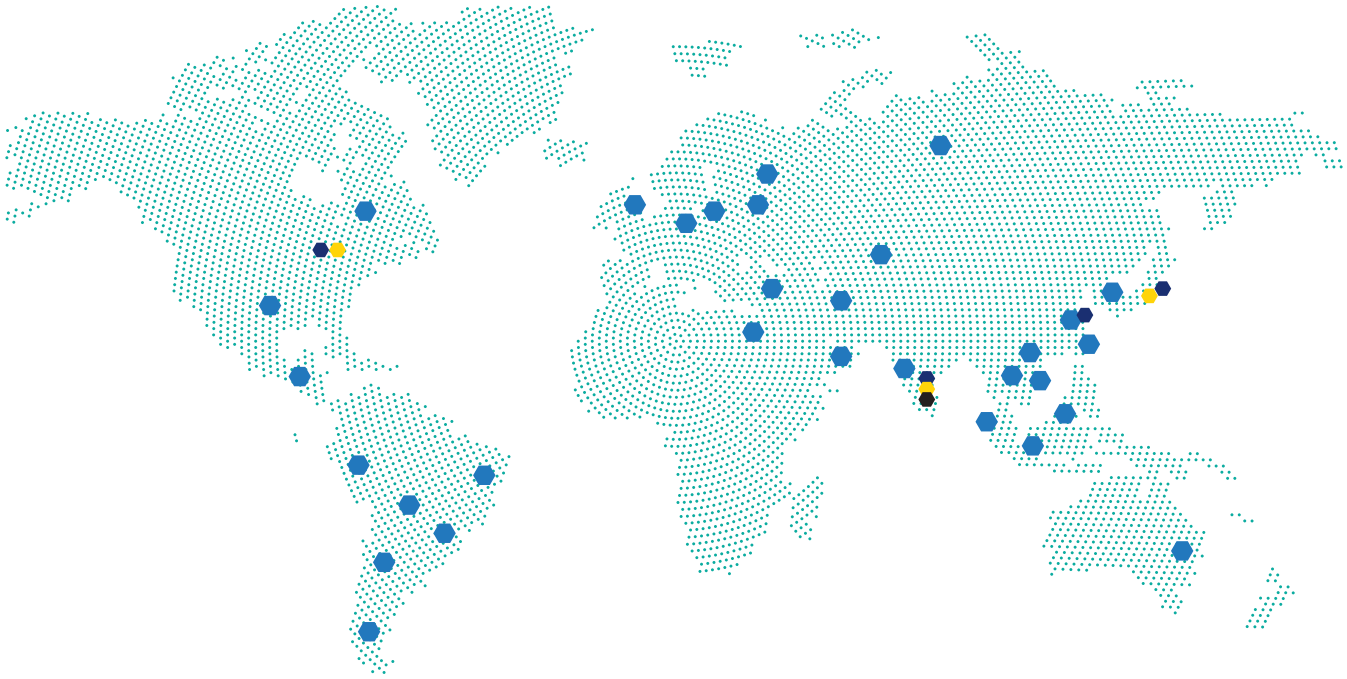
The Lifesciences Vertical of STEER World

A Global Leader in Continuous Flow Manufacturing Technology

STEER World is a global leader in pioneering continuous flow manufacturing innovation across diverse industries such as pharmaceuticals, polymers, plastics, nutraceuticals, food, sustainable materials, biofuels and biorefining.



Let's Grow Your Portfolio, Together.



Global Offices Channel Partners

Application Development Centers Manufacturing Facility

STEER Life

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Process. Product. Performance.