













Consulting

Understanding target indications

Matching MOA to disease areas

Market evaluation

Clinical-regulatory strategy and guidance

Creating business plans

Full clinical development plans and outlines

Capital

Matchmaking

Asset evaluation to support fundraising

Utilizing network of pharma and VCs

Business planning to determine capital needs

Partnering and creative collaborations

Formulation

Full project management support

Vendor management

Large network of contract manufacturers

Consulting with formulation experts

Target product profile creation

Preclinical

Variety of disease and efficacy models

Model selection based on indication and MOA

Matching preclinical studies to clinical strategy

Full project management and study conduct

Strategic alliances with labs

Clinical trials

Phase 1-4 trials

Complete management, study conduct and support

Large database of study patients

Vast site network across the US

International CRO Partners

Data management and biostatistics partners

Regulatory

Filing for Pre-IND/Pre-IDE and IND/IDE meetings

Meeting briefing packages

FDA meeting requests and letters

Market

Phase IV post-marketing studies, patient registries,

Medical writing of abstracts, posters, podium presentations, scientific papers and trade publication messaging

KOL development and messaging planning

New product packaging concepts and website development

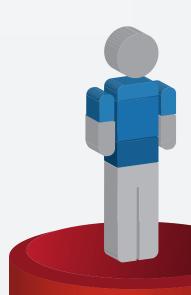
Mid-SizePharma

We offer a number of services to address Mid-Size pharma firms' needs, from market evaluation and consulting services to regulatory assistance in filing for Pre-IND/PRE-IDE and IND/IDE meetings. Our capital services expertise includes business planning to determine capital needs, as well as asset evaluation to support fundraising.

We provide formulation and preclinical services for early stage assets, full support and conduct of phase 1-4 clinical trials, full project management support for their programs, a variety of disease and efficacy models in preclinical and clinical stages, and data management and biostatistics expertise.

Learn More
Visit oraclinical.com

Call us: (978) 685-8900 email us: info@oraclinical.com



The Ora Advantages

- 1. On-time study completion
- 2. Knowledgeable staff
- 3. Strict protocol adherence



Ora stands apart from all other research groups in that we have conducted over 1,300 clinical trials and consulting projects within the ophthalmology category.

This deep knowledge and expertise has made Ora the preferred ophthalmic drug development partner for many of the top ophthalmic pharmaceutical companies in the world.