

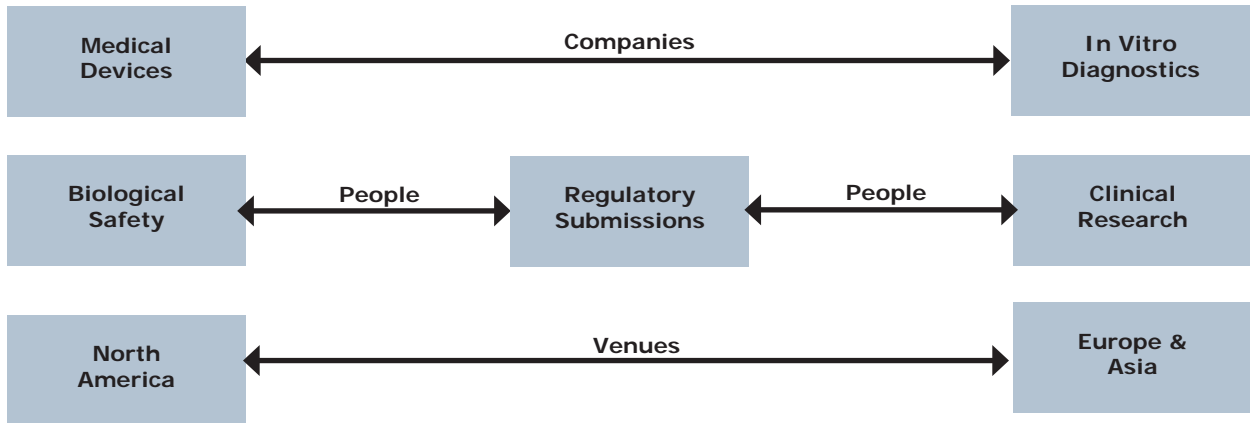


CLINICALDEVICEGROUP
Medical Device Evaluations

Rethink Your Expectations

About Us

Three Kinds of Customers



We do business with medical device and in vitro diagnostic companies from around the world, focusing our expertise on biological safety, clinical research, and regulatory submissions.

Four Kinds of Service



Cafeteria-style or turnkey, we are a full service contract research organization. We can do all or part of a clinical trial, consult on regulatory affairs, or prepare expert toxicology reports to address biological safety.

Our [eConferences](#), [Clinical Device Forum](#), on-site seminars, and [soon] on-demand training are cornerstones of our success. And our books and productivity tools for medical device good clinical practices are designed to help you work independently.

CDG serves the medical device industry, only. That's our love, speciality, and commitment.



The Leader



Nancy J Stark, PhD, President

Her major accomplishments include:

1990 - Founded CDG Inc.

2001 - Appointed US Co-chair to WG4/TC194, ISO 14155.

2002 - Founded the Clinical Device Forum.

2004 - Named one of 100 Notable People in Medical Devices by MD&DI Magazine.

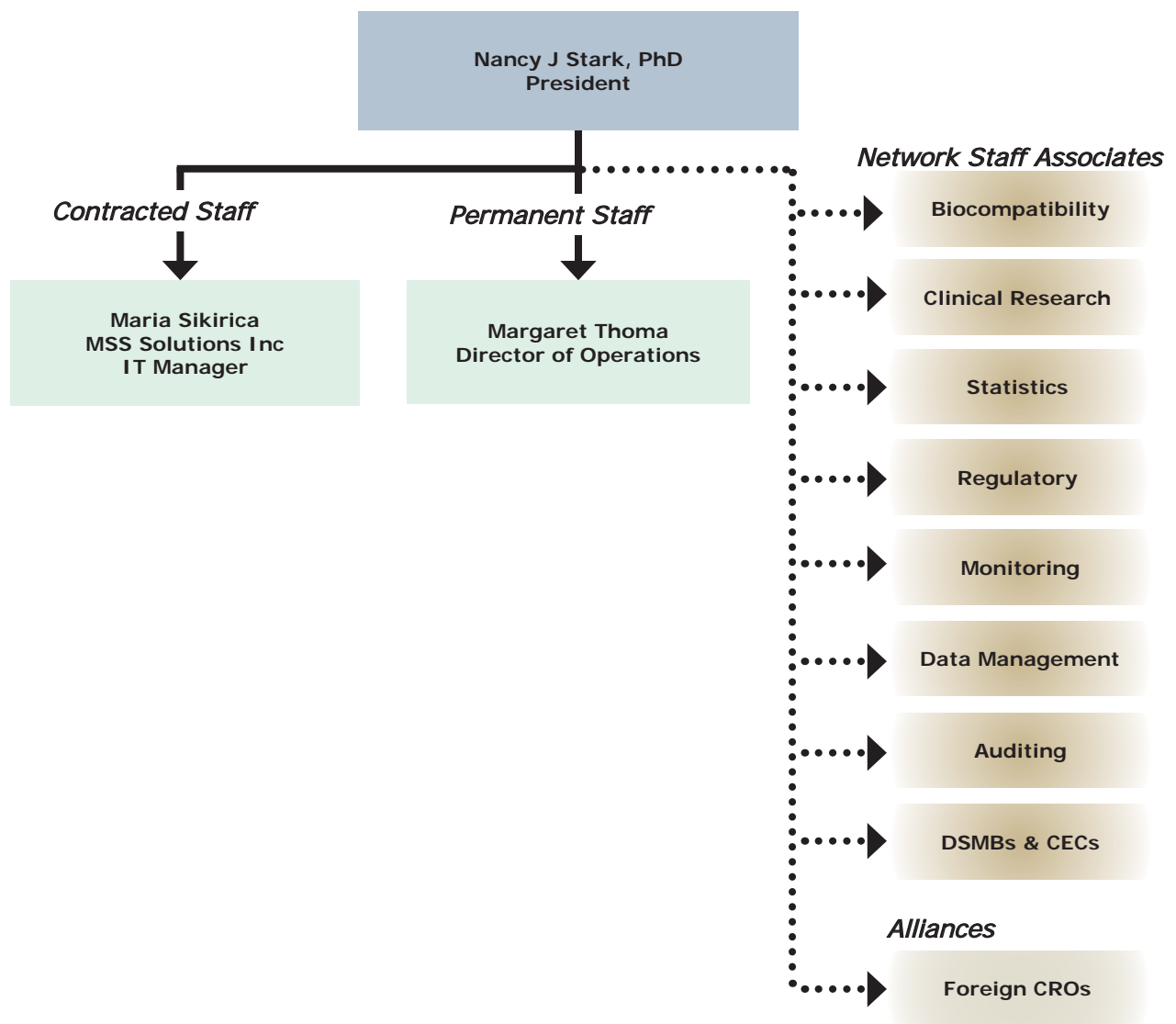
2007 - Invited to Editorial Advisory Board of MD&DI.

You can download her current curriculum vitae at www.nancyjstark.com.

Stark publishes regularly in peer-reviewed journals and is viewed as a world expert on good clinical practices for medical devices.

Organization

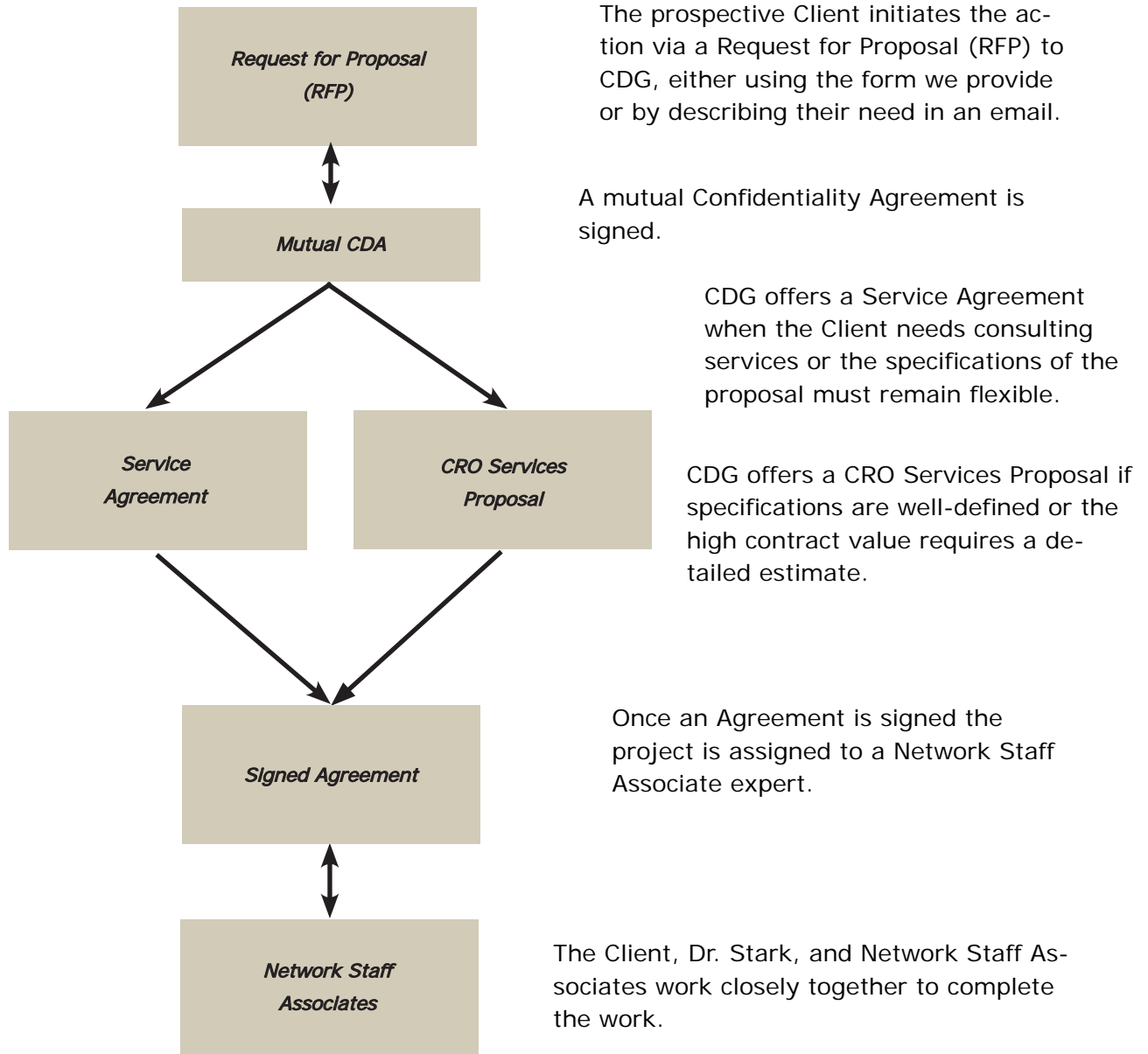
Clinical Device Group was founded in 1990 by our President, Nancy J Stark, PhD. She is supported full time by a Business Manager, Office Assistant/Corporate Artist, and Executive Assistant/Bookkeeping and as needed by an orchestra of Network Staff Associate and CRO Alliance sub-contractors. The Associates are expert independent professionals, the Alliances extend our service to you worldwide. We only work with medical device specialists.



Our orchestra of Network Staff Associates and CRO Alliances gives you instant access to a broad range of professional experience.



Getting Started with CDG

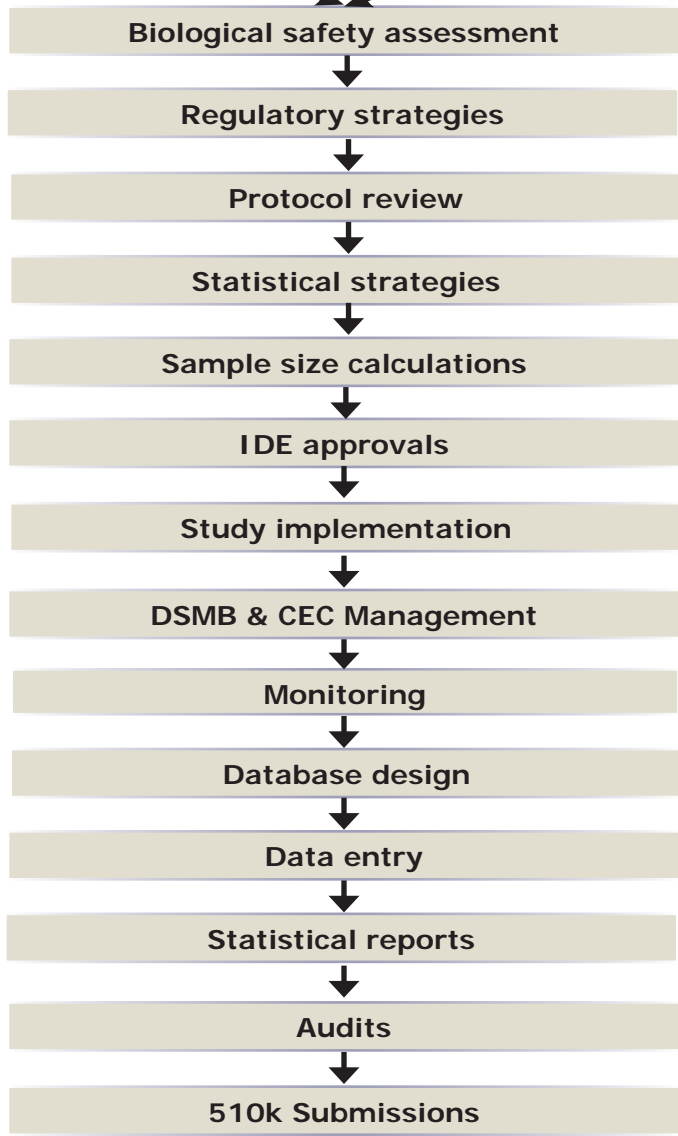


We work closely with Clients to assure your managerial control of projects. All Agreements are fee-for-service, so you pay only for what you use.

Our Capabilities



Product Development Process



Cafeteria style, or turnkey,

we can help you with biological safety,

clinical research,

or regulatory submissions.

Our commitment is to medical devices.

Market Entry!



Throughout the Year



Since 1990

We are as solid as the 1890 Victorian flat we live in. Visit us anytime. The first day of capabilities assessment, project presentation, consulting, and lunch, is on us.



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