

BIOCARD Logistics

BIOCARD'S HISTORY

1988 year of creation of Biocard, the first foreign trade company in the system of Soviet Russia's healthcare.

In 1989 Biocard managed the first in the Soviet Republic international clinical trial. In fact it received monopoly right from the Ministry of Health to undertake such studies in Russia.

In 1990 this thriving and rapidly developing business was taken by the company's management private, which led to creation of two entities: MPC Mirgom and MPC Pharma, responsible for clinical trials and logistical support of clinical trials respectively.

In 1993 MPC Mirgom opened an official representative office in Europe (Frankfurt/Main, Germany).

In 2002 Biocard's team received a legal right to use officially its historic and original name BIOCARD.

In 2006 logistical line of our services was reorganized into Biocard Logistics. Logistical Support.

Biocard is ready to offer distribution services for whole cycle of Clinical Trial in Russia starting with importation of any clinical materials and ending with its destruction or exportation.

Biocard Research owns state-of-the-art drug storage facilities:

- Company's own drug storage facilities, separate and secure storage areas equipped with temperature and humidity gauges and computer control.
- Compliance with all temperature requirements: storage at room temperature (+18 -+25 C), in refrigerators (+2 -+8 C) and deep freezers (-20 C), including storage at -86 C.
- Company's own staff (pharmacist) to manage storage, distribution & drug accountability.
- Monitored Access and top class security system.

Highly qualified personnel, extensive tacit knowledge accumulated by the organization through out the years of its existence.

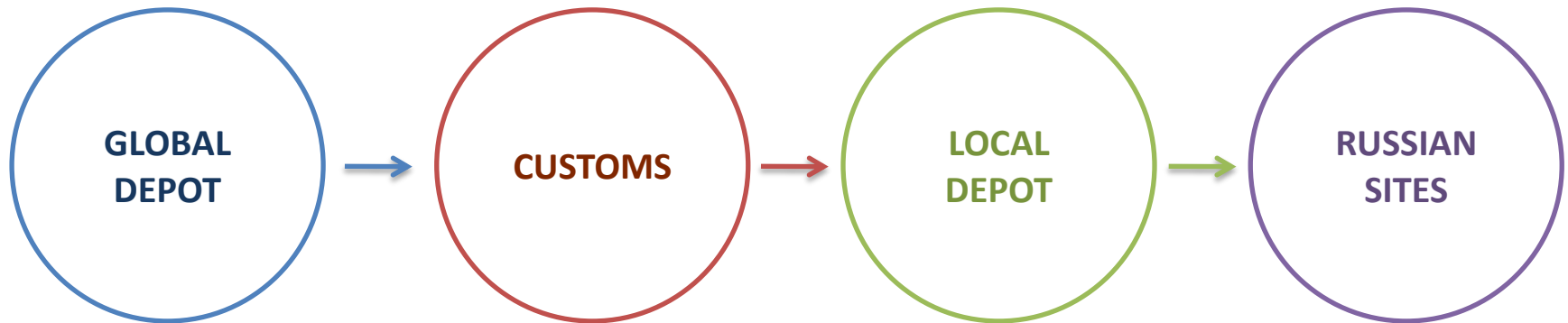
First-line supervision, highly structured internal procedures, minimal lead times and subcontracting to third parties.

Effective Quality Assurance System confirmed by International Standard ISO 9001:2000.

Our passion for quality, precision, consistency, self-education and constant drive for improvement!







WHY USE LOCAL DEPOT?

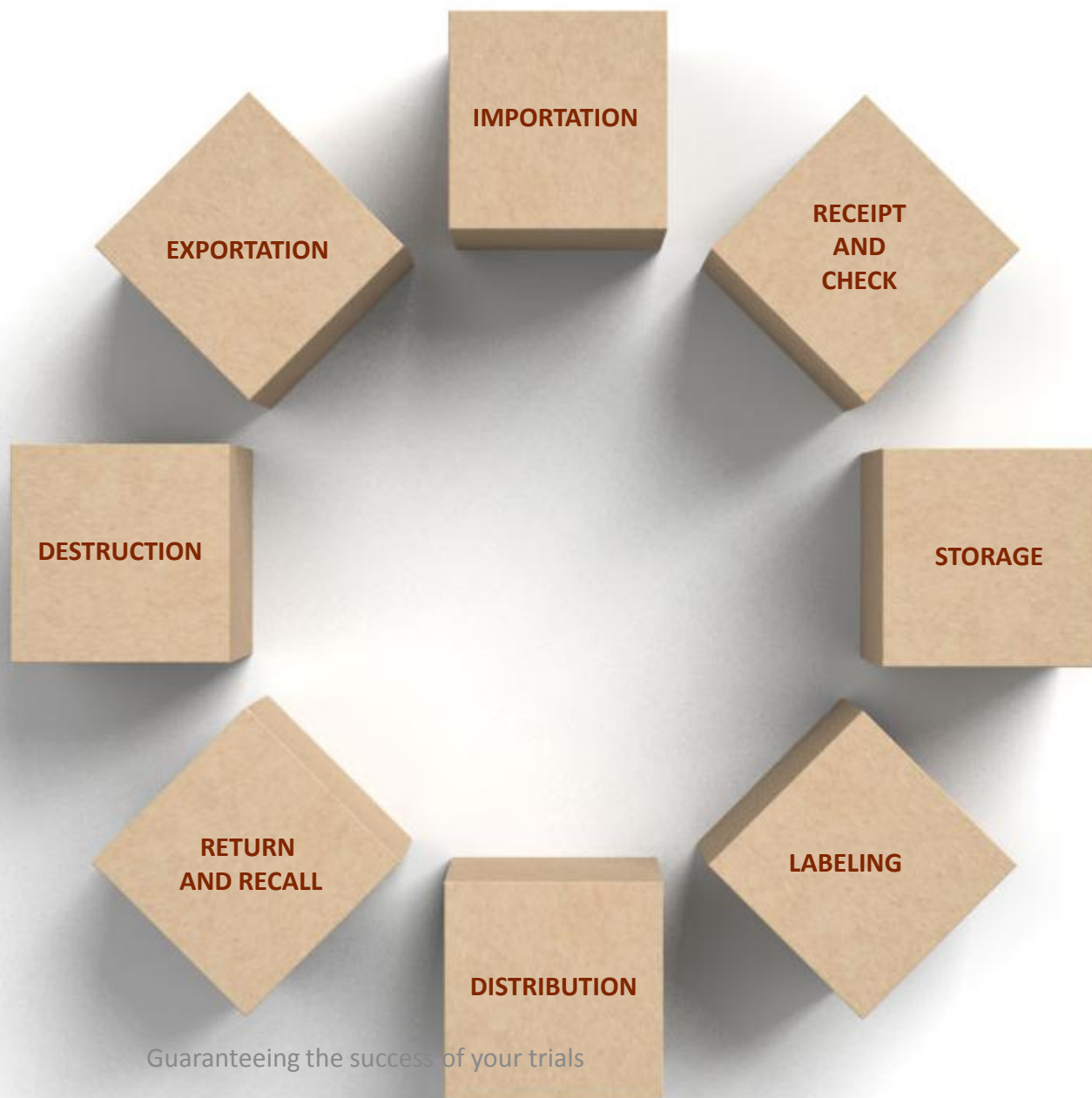
By leaving you to concentrate on what you do best...

- National culture and mentality
- Constantly changing regulations
- Administrative and bureaucratic barriers
- Harsh climate conditions

... we save your most precious asset – “time”!

- Initiation period
- Importation / exportation process
- Working across vast distances
- Numerous time zones
- Cash-flow management
(helping to evade cash breaks).





- Importation/exportation of all kinds of study materials (drugs, lab Kits, CRF's, ECG machines, etc.)
- Your consulting assistant in obtaining import/export licenses
- Process timelines do not exceed 3 working days.

- Own validated storage facilities designed and equipped for separate storage of different study materials
- Equipped with temperature and humidity computer management system
- Equipped with validated devices to provide different temperature conditions:

+18 ... +25°C / +2 ... +8°C / -20 ± 5°C / -86°C



- Special temperature and humidity controlled premises to ensure relabeling process
- Experienced and trained staff
- Accountability



- Special validated, temperature and humidity controlled premises for quarantine storage
- Accountability and corresponding reports



- Contracts with only certificated destruction companies
- Compliance with Russian and international standards
- Destruction of all kinds of clinical study materials
- Full package of necessary documentation.

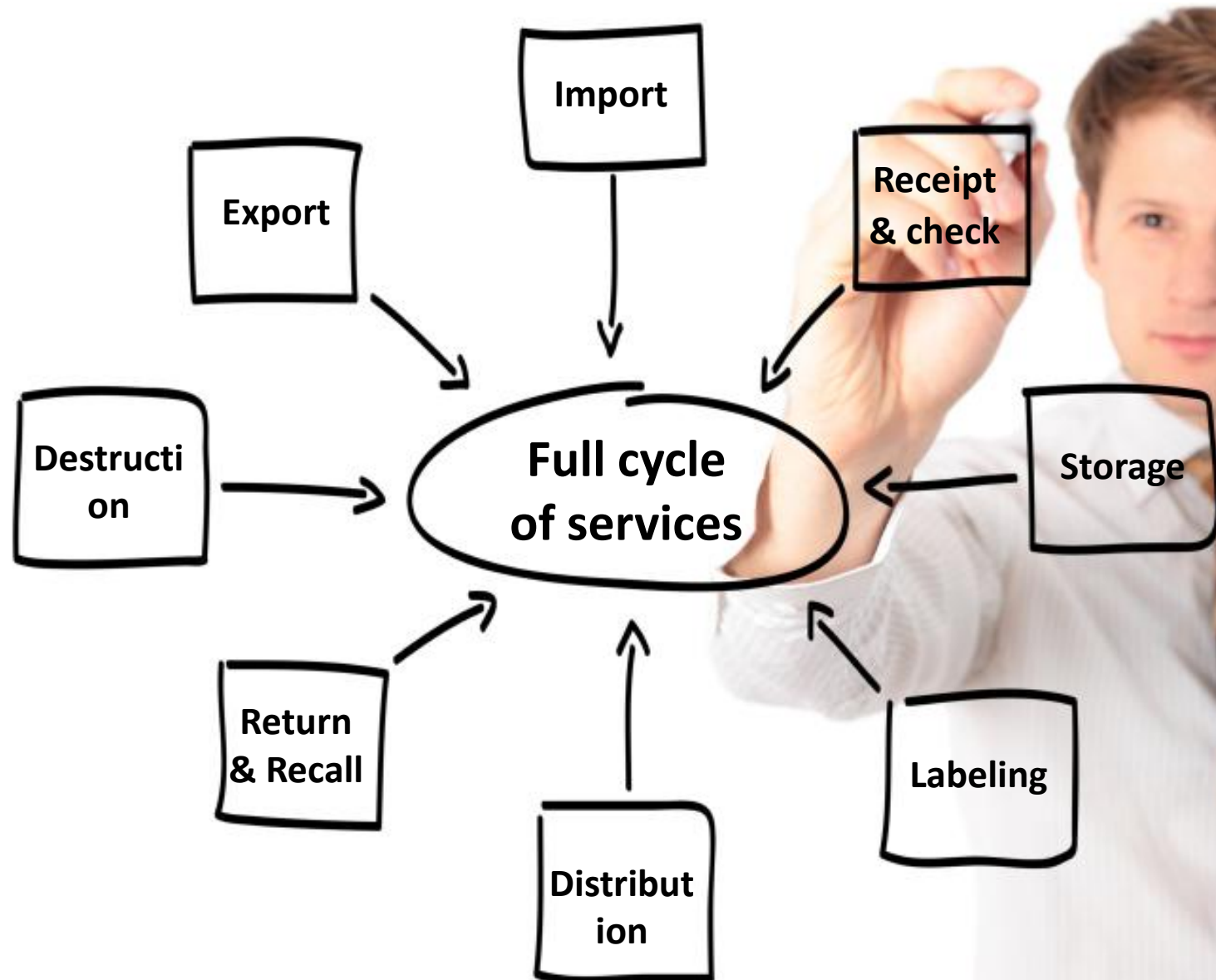


We save your money by providing timely service of a consistently high quality!



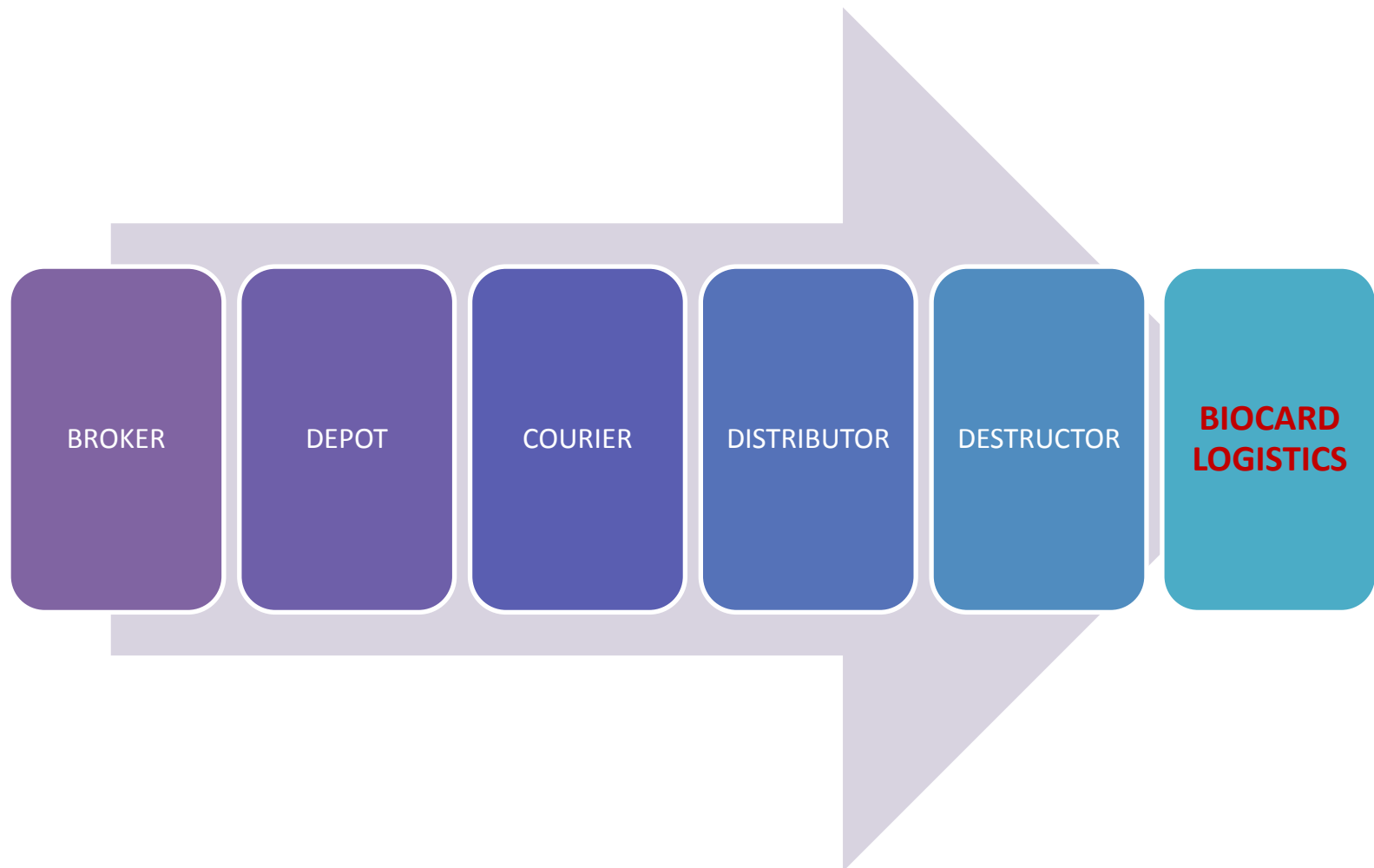
WHY WORK WITH US?





Having communication issues?

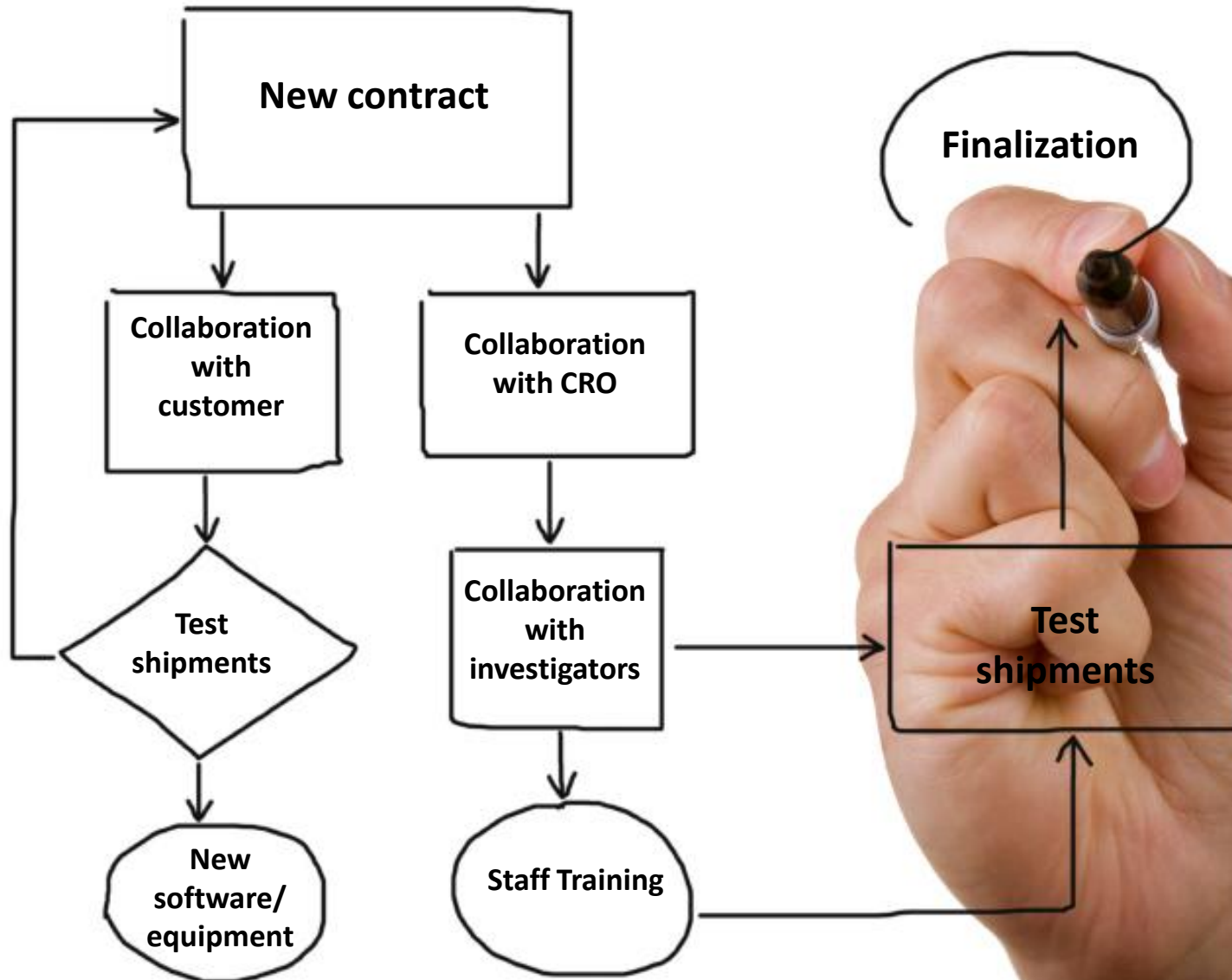




To give you an example, consider the number of audits we went through out the last several years.

January 2007	VENDER AUDIT	GLOBAL CRO
April 2007	POTENTIAL CUSTOMER AUDIT	GLOBAL CRO
June 2007	VENDER AUDIT	GLOBAL CRO
August 2007	ISO CERTIFICATION AUDIT	
August 2007	VENDER AUDIT	PHARMACEUTICAL COMPANY
September 2007	POTENTIAL CUSTOMER AUDIT	GLOBAL DEPOT
October 2007	VENDER AUDIT	PHARMACEUTICAL COMPANY
December 2007	ISO CERTIFICATION AUDIT	
January 2008	POTENTIAL CUSTOMER AUDIT	PHARMACEUTICAL COMPANY
Mach 2008	POTENTIAL CUSTOMER AUDIT	PHARMACEUTICAL COMPANY
April 2008	POTENTIAL CUSTOMER AUDIT	PHARMACEUTICAL COMPANY
April 2008	POTENTIAL CUSTOMER AUDIT	GLOBAL DEPOT
August 2008	VENDER	PHARMACEUTICAL COMPANY
September 2008	ISO CERTIFICATION AUDIT	
September 2008	VENDER AUDIT	GLOBAL CRO
January 2009	NATIONAL AUTHORITIES AUDIT	
January 2009	POTENTIAL CUSTOMER AUDIT	GLOBAL CRO
July 2009	VENDER AUDIT	GLOBAL CRO
August 2009	VENDER AUDIT	GLOBAL CRO
September 2009	ISO CERTIFICATION AUDIT	
September 2009	VENDER AUDIT	PHARMACEUTICAL COMPANY
January 2010	QA AUDIT	INTERNAL

NOT JUST A SIMPLE DISTRIBUTOR



- Main training plan for all employees
- Average staff working experience of at least 5 years
- Minimal staff turn-over!



Our basic strategy is a clear execution of customer's requirements.

But we aspire “to foresee customer's wishes”.

Give us a chance and present yourself with ease of mind!



- ISO 9001-2000 certificated quality system
- SOPs for ALL EXECUTED PROCEDURES
- Well structured disaster recovery plan
- Clear lines of reporting!



Give us a call, and our representative will
be glad to answer your questions!

Germany

Markstrasse 5, Frankfurt/Main
Germany, D-60388

Russia

Karamyshevskaya naberezhnaya
Moscow, Russia, 123423

