



Professional services and technology solutions

TMF Migrations from any source to any eTMF

TMF Audit Services



Multi-Channel Marketing for samples distribution

[www.drscorp.com](http://www.drscorp.com)

**2011:** The development of HTML program language and SaaS (Software as a Service) applications were becoming the norm. No matter which mobile device was used by our client base, there was a solution. We deployed our Rx sampling solution on Windows (PC, CE, SVS) in North American GSK sales and replaced their legacy systems with SaaS products to enable real time sampling, HCP ordering and regulatory reporting.

**2012:** DRS launched the latest version of DRS eTMF software, offering new techniques and options to improve scalability and provide even better proactive reporting processes. DRS worked with the Bill and Melinda Gates Foundation to expand their worldwide studies related to Tuberculosis. At the same time we expanded our TMF product group to include TMF Audit Services, conducting audits and reporting on the integrity of sponsor studies while ensuring inspection readiness of studies managed by CRO vendor partners.

**2018:** Our three decades of experience enable us to develop products and services that fill in the blanks of a life science organization and their corporate strategy. Agile software development and forward thinking methods help our client partners to realize their goals no matter how difficult they may seem. DRS will develop the strategy that has kept us on a successful path to long term partnerships with our customers for three decades. We feel that trust in a partnership helps to maintain the core strength of the relationship. This is one of our company and building on that strength is our goal for the next years to come.

**2014:** DRS introduced the ideal electronic sample ordering system, DRS 360 Plus. A system where Sales and Marketing could select when and how drug samples would be delivered. Health Care Professionals and product development encompasses the latest regulatory requirements for monitoring of Terminal Distributed Dangerous Drugs (TDD). The legislation assists with monitoring of Opioid distribution, a topic that has become very important to many science organizations.

**2017:** With background of 30 years document management, DRS was selected to complete migration of over 20 million clinical documents for sponsor organizations who need to convert legacy paper, microfilm and electronic formats into one common platform. Our Life Science customer base for both clinical studies and commercial operations continues to grow. This is surely a testament to DRS' well-designed technology but also to its service professionals who integrate solutions and services with our client base.

What will tomorrow bring?

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