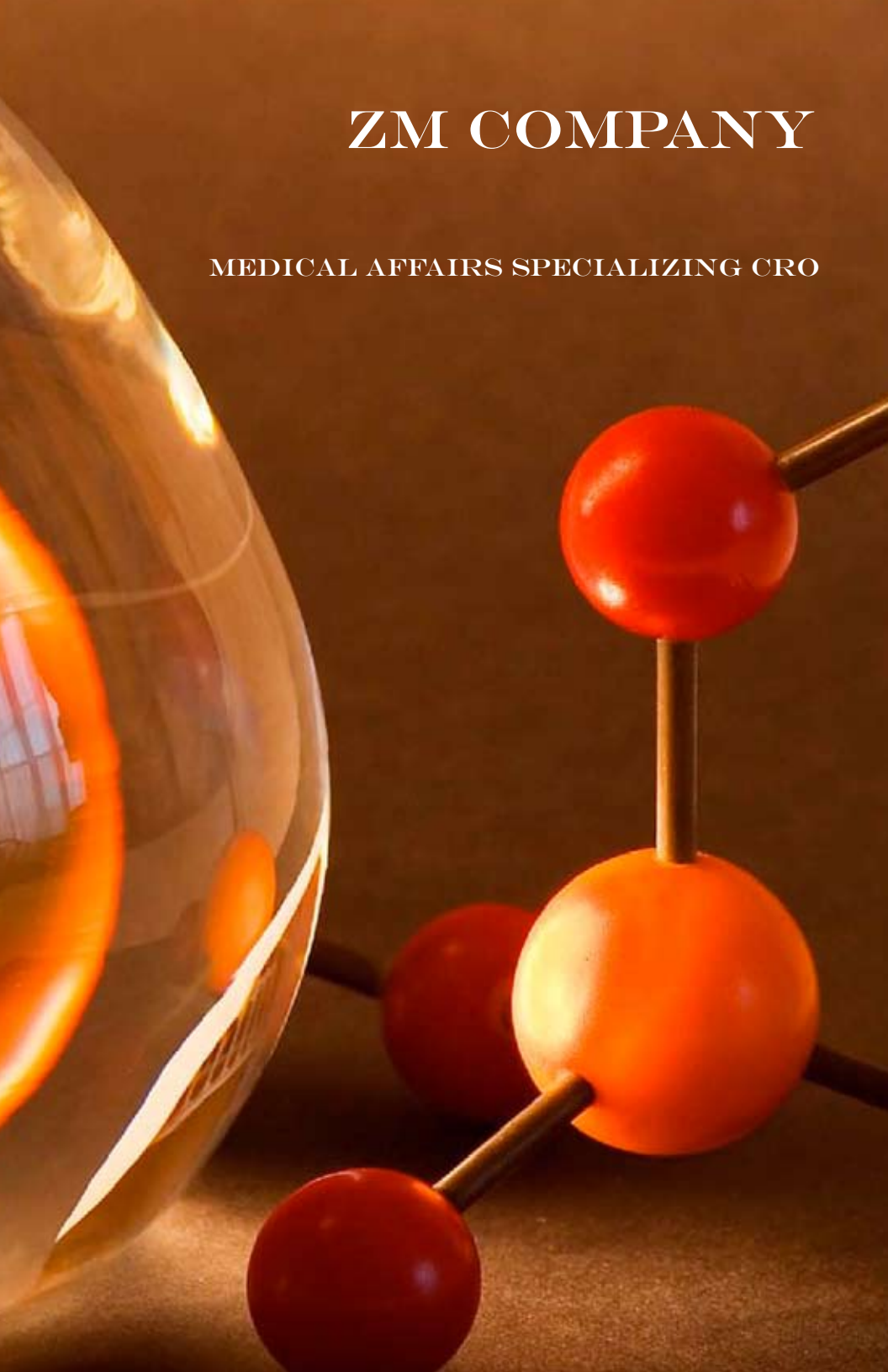


# ZM COMPANY

MEDICAL AFFAIRS SPECIALIZING CRO



**TAILS I WIN. HEADS YOU LOSE.**



**There's no luck involved in the selection of a first-rate CRO.**

**In a highly regulated industry, such as Clinical Trials, there is no room for error in service provider selection.**

**The pharmaceutical and biotechnology companies are looking for special knowledge, thorough understanding of regulations, quality and high standards.**

**THAT'S WHERE WE COME IN.**

**SUPERB QUALITY BEGINS WITH EXPERIENCE.**

**Our team consists of  
qualified and experienced  
physicians.**

**We know industry standards, methodology and regulations.**

**Our expertise covers a broad array of therapeutic areas in all phases  
of Clinical Trials.**



## BRIDGING THE GAP.

**We have a unique, unparalleled in the industry combination of medical knowledge, clinical trials experience, and skills in clinical data management.**

**We bridge the gap between clinical and data management aspects to ensure the medical accuracy and integrity of clinical trial data.**

A close-up, artistic photograph of a clock face. The focus is on the Roman numeral 'IV' (4) and the 'III' (3) below it. The clock's hands and the metal casing are visible, with a warm, golden-brown color palette. The background is softly blurred, showing more of the clock's mechanism.

**EVERY SINGLE-MINUTE DELAY  
IN DRUG R&D COSTS \$1.800 IN LOST REVENUE.**

**When it comes to fast-turnarounds, often you need it done by tomorrow. We will at least aim for today, without sacrificing quality.**

**Your goals become our goals. And your success becomes the only true measure of our success.**

**COST A PRETTY PENNY.**




**We bring together a combination of skills that ensure that the clinical trial delivers high quality data at a competitive and affordable price.**

## **LIST OF SERVICES.**

**24/7 Medical monitoring**  
**Qualified Person for Pharmacovigilance in Europe**  
**SAE management and reconciliation**  
**Case narrative writing**  
**Safety data review & medical review of CRFs**  
**Safety data & Pharmacovigilance audit**  
**Medical writing**

**Medical data coding**  
**Coding consistency review**  
**CRF design and development**  
**Medical data validation /edit**  
**checks development**  
**Training**  
**Consulting**

## **CONTACT US**



**Let us make a world of  
difference for you.**

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