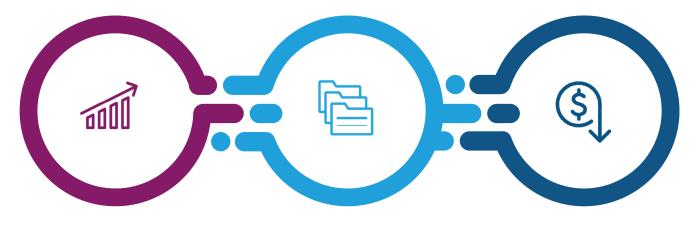


# **IQVIA** Technologies **Risk-Based Monitoring**

## **Orchestrate better outcomes with** IQVIA Technologies Risk-Based Monitoring



12.5

Days faster to database lock vs. non-RBM studies 31%

Less SDV backlog for RBM studies vs. non-RBM studies

21% **Reduction** in patient visit data entry lag



**Cost savings** Up to 25% cost savings when compared to traditional monitoring



Improved patient safety 67% of queries resulted in data base updates, with 74% of these related to patient safety

References:

1) Branch, E. (2016, April 30). Ways to Lower Costs of Clinical Trials and How CROs Help. Retrieved January 21, 2019, from https://www. americanpharmaceuticalreview.com/Featured-Articles/185929-Ways-to- Lower-Costs-of-Clinical-Trials-and-How-CROs-Help/

## Interoperable

Works seamlessly with IQVIA product portfolio and also 3rd party

## Superior data quality

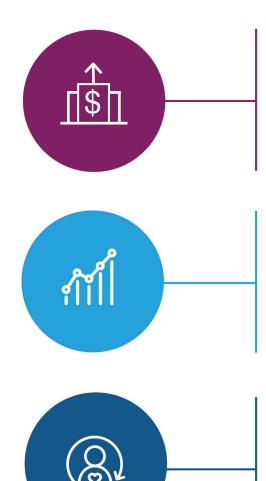
Allows actionable insights from many different data points

#### Identification

Allows early identification of trends, risks and outliers - with



## **Benefits** -



### **RBM reduces risk and cost**

- Intuitive workflow for faster, more informed decisions
- Intelligent AI/ML-enhanced application for better data analysis
- Interoperable with agile applications that work well on their own but even better together

## **RBM** improves study quality

- · Lowers error rate with real-time data entry and site communication
- · Drives timely actions with predictive and prescriptive analytics
- Leverages current and historical data and suggest "next best action"

### **RBM** enhances patient safety

- Predictive and advanced analytics help resolve issues immediately at site and patient level
- · Focuses on higher risk sites with timely communication and compliance
- Allows patient-level data reviews to identify trends and assure medical accuracy and congruence

## Features -



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Automated risk identification and assessment: IQVIA Knowledge Base; automated mitigation plan; configurable KRIs, CDPs; KRI and **CDP** library

Advanced & predictive analytics on aggregated site & subject data at study level; perform study reviews and compare to historical outcomes

Dashboard & drill downs for site status based on core KRIs and site data; automated & on-demand review generation; automated alerts and tasks based on user configuration

Subject comparison across the study; early signal detection to flag outliers and inconsistencies; analytics & trend analysis for more precise subject reviews

Operational reports; outcome reports

VISIT

iqvia.com/OCT



**TECHNOLOGIES** 



**Reports and insights** 

Study oversight

Site risk monitoring

Subject data surveillance

