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A Revolution in R&D - Boston Consulting Group

Biopharmaceutical R&D is moving into a new era: almost every link in the value chain has the potential for tremendous boosts in efficiency or success But these advances are not assured Technological hurdles have yet to be overcome, particularly in the genet-ics wave Moreover, because the productivity boosts are likely to be unequal and

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Indian life sciences: Vision 2030 Expanding global relevance and driving domestic access 3 Executive summary 3 PAREXEL Biopharmaceutical R&D Statistical Sourcebook 2014 on "How to improve R&D productivity" in 2010, Team analysis

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R&D Spending: Pharmaceuticals 6 PAREXEL's Pharmaceutical R&D Statistical Sourcebook 2005/2006 New data continue to support the view that the pharmaceutical sector's center of gravity in terms of sales, R&D spending, and new product introductions has shifted from

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Maximizing Value and Quality in Phase IV Trials

Source: CTgov, PAREXEL Biopharmaceutical R&D Statistical Sourcebook 2011-2012 > Increased pressures from providers and payers for 'real-world' data generated outside of a controlled trial environment > Technology advancements and EHR adoption have made observational studies more cost-effective > Scientific methods for planning, analyzing and

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2012 Parexel Biopharmaceutical R&D Statistical Sourcebook 2012/2013; Medidata estimates Total Life Sciences IT spend (Software, hardware, IT services) = ~$45B Global clinical development spend (phase I – IV) = ~$85+B • Potential for Medidata to expand software capabilities to move into

areas currently addressed by IT services and hardware

Industry Perspective: The Challenges and Benefits in using ...

Industry Perspective: The Challenges and Benefits in using Expedited Regulatory Pathways Alan Poirier, Pfizer Inc Parexel Biopharmaceutical R&D Statistical Sourcebook (2013-2014) 12 - 14 years* for preclinical, clinical development, and regulatory review AAPS November 2016 4

CHI Policy Statement - Part D Rebates - May 2013

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