Radio Frequency Identification (RFID) in Pharmaceuticals -Supply Chain Security Concerns Provide Impetus for RFID Adoption

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RFID is the most promising solution available to prevent counterfeiting

Radio Frequency Identification (RFID) in Pharmaceuticals - Executive Summary

GBI Research's report on the markets for RFID in the pharmaceutical industry provides a comprehensive analysis on the applications of RFID technology in the pharmaceutical supply chain. The market for RFID hardware, software and services is forecast for seven years and the key factors driving or restraining the market are analyzed. Analysis on regulations and mandates provides valuable insights into the key trends that drive the growth of RFID in the pharmaceutical industry.

Pharmaceutical Industry's Pressing Need to Regain Lost Reputation and Revenues Due to Counterfeit Drugs Will Drive the Growth of RFID Market

GBI Research finds that there is a growing need for the pharmaceutical industry to secure their distribution channels from counterfeit drugs. RFID solutions provide the ideal identification method by which pharmaceutical industry can counter fake drug issues. While pharmaceutical companies, distributors and wholesalers are on a continuous look out for these fake drugs, counterfeiters find innovative ways to introduce fake drugs into legitimate supply chains. The use of RFID solutions can avoid the loss of investor confidence and a consequent decline in share prices due to such counterfeiting incidents. Also, increased supply chain security and operational efficiency will help the pharmaceutical industry improve their reputation in the delivery of safe drugs. Hence, increasing concerns over the safety of supply chains and the health of patients have forced companies to think big and adopt RFID solutions. A steady increase in the adoption of RFID hardware, software and services in the pharmaceutical industry is expected to drive the growth of these solutions.



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In developing countries counterfeit drugs are estimated to account for approximately XX% of the total pharmaceutical market

5.6 Markets for Counterfeit Drugs

Counterfeit drugs are one of the key challenges facing pharmaceutical supply chains and the safety of patients. It is estimated that the global market for counterfeit drugs was \$XX billion in 2006 and is expected to grow to \$XX billion by 2010.

Counterfeit drugs are those drugs which are sold under a product name without authorization and which are sold with the intention of misleading the customer into believing that the drug is original. Counterfeiting is one of the major problems facing healthcare systems across the world. It is more prevalent in developing countries where there is limited control over the flow of drugs through the supply chain.

Counterfeiters find weak links in the supply chain to introduce fake drugs and so counterfeiting market thrives in developed countries where the movement of goods in the supply chain is not strictly regulated. In the developing countries of Africa, Asia and Latin America, counterfeit drugs constitute nearly XX% of the total pharmaceutical market.



7 Radio Frequency Identification (RFID) in the Pharmaceuticals – Market Characterization

7.1 Market Forecasts for RFID in Pharmaceuticals

The global market for RFID solutions in the pharmaceutical industry will grow rapidly due to the expected increase in awareness and the alleviation of concerns regarding implementation. The global market for RFID solutions in the pharmaceutical industry was valued at \$XXm in 2008. It is expected to grow to \$XXm in 2015 at a compound annual growth rate of XX%.



Table 7: RFID in Pharmaceuticals, Global, Revenues (\$m), 2008-2015									
Year	2008	2009	2010	2011	2012	2013	2014	2015	CAGR (%)
Revenue (\$m)									
Growth Rate (%)									
Source: GBI Research									

A lack of awareness about RFID technology and the lack of a proof-of-concept for returns on investments have hampered the growth of the market for RFID in pharmaceuticals. Although RFID technology has been implemented in industries such as the automotive industry, its implementation rate in the pharmaceutical industry has been slow. The pharmaceutical industry is reluctant to make large investments in RFID implementation as it is unsure about the efficiency of the technology in real-time functions. One of the key concerns raised is the lack of standardization across the supply chain that could lead to the inefficient functioning of RFID systems. Although RFID technology is being deployed in closed-loop systems, where tags can be reused, deployment in open loop systems will require an increase in awareness and proof-of-concept studies. The effectiveness and reliability of RFID systems in open-loop systems need to be demonstrated to encourage industry participants to adopt RFID technology.

The global market for RFID solutions was valued at \$XXm in 2008

9 Radio Frequency Identification (RFID) in the Pharmaceuticals – Regulatory Landscape

9.1 Regulations in the US Encourage RFID Adoption in the Pharmaceutical Industry

9.1.1 The Prescription Drug Marketing Act of 1987 (PDMA)

The Prescription Drug Marketing Act of 1987 (PDMA), as modified by the Prescription Drug Amendments of 1992, amended sections 301, 303, 503, and 801 of the Federal Food, Drug, and Cosmetic Act (the Act) to establish the requirements related to the wholesale distribution of prescription drugs. One of the primary purposes of the PDMA was to increase safeguards to prevent the introduction and retail sale of substandard, ineffective, and counterfeit drugs in the US drug supply chain.



9.1.2 California Pedigree Legislation's Deadline has been Postponed on Multiple Occasions Due to the Challenges Faced in RFID Implementation

According to the California pedigree legislation, a "Pedigree" means a record, in electronic form, containing the information regarding each transaction resulting in a change of ownership of a given dangerous drug, from the sale by a manufacturer, through to the acquisition and sale by one or more wholesaler, manufacturer or pharmacy, until the final sale to a pharmacy or other person furnishing, administering, or dispensing the dangerous drug.



Radio Frequency Identification (RFID) in Pharmaceuticals -Supply Chain Security Concerns Provide Impetus for RFID Adoption

12 Appendix

12.1 Market Definitions

Pharmaceutical manufacturer – Manufacturer's of traditional chemical drugs and biopharmaceuticals are classified as pharmaceutical manufacturers

Trading partners – Authorized wholesalers, secondary wholesalers, distributors, retail pharmacy chains and pharmacies are classified as trading partners.

Authorized wholesalers – Authorized wholesalers buy drugs directly from the pharmaceutical manufacturers and have an ongoing relationship with the manufacturer

Secondary wholesalers – Secondary wholesalers buy from authorized wholesalers and sell to retail, hospital and individual pharmacies

Radio Frequency Identification – it is an automatic identification technology used to store and remotely retrieve data using devices called RFID tags or transponders

RFID Solutions Market – The scope for RFID solutions market includes the markets for RFID hardware, software and services.

Service providers – RFID service providers include the players who offer services such as installation, integration, maintenance, IT support and training

One-stop solutions – This type of RFID solution includes a single industry participant or a collaboration of industry participants which offers a combination of hardware, software and services for complete implementation and integration

12.2 Abbreviations

ADR	Authorized Distributors
API	Active Pharmaceutical Ingredient
ASN	Advance Shipment Notice
CAGR	Compound Annual Growth Rate
ССРА	Coalition for Community Pharmacy Action
DTP	Direct-to-Pharmacy
EAN	European Article Number
ECJ	European Court of Jurisprudence
EMEA	European Medicines Agency
EPC	Electronic Product Code
ETSI	European Telecommunication Standards Institute
EU	European Union
FDA	Food and Drugs Administration
HDMA	Healthcare Distribution Management Association
HF	High Frequency
IS	Information Services
ISO	International Standards Organization
LBT	Listen before Talk
LF	Low frequency
MAH	Market authorization holder
ONS	Object Naming Service
PDMA	Prescription Drug Marketing Act of 1987
PhRMA	Pharmaceutical Research and Manufacturers of America

Radio Frequency Identification (RFID) in Pharmaceuticals -Supply Chain Security Concerns Provide Impetus for RFID Adoption

RAM	Random Access Memory
RFID	Radio Frequency Identification
ROM	Read Only Memory
UHF	Ultra high frequency
ICE	Immigration and Customs Enforcement
EFPIA	European Federation of Pharmaceutical Industry and Association
DS	Discovery Service
GPOs	Group Purchasing Organizations
WHO	World Health Organization
NHS	National Health Service
ROI	Return on investment
UAE	United Arab Emirates
3IS	Internet Infrastructure and Identity Services
FSA	Fluidic Self Assembly
GSC	Global Security Consulting
IAS	Identity and Authentication Services
IC	Integrated Circuits
NABP	National Association of Boards of Pharmacy
ROI	Return-on-Investment
SSL	Secure Socket Layer

12.3 Research Methodology

GBI Research's dedicated Research and Analysis Teams consists of experienced professionals with a pedigree in marketing, market research, consulting background in the medical devices industry and advanced statistical expertise.

GBI Research adheres to the Codes of Practice of the Market Research Society (www.mrs.org.uk) and the Society of Competitive Intelligence Professionals (www.scip.org).

All GBI Research databases are continuously updated and revised. The following research methodology is followed for all databases and reports.



12.3.1 Coverage

The objective of updating GBI Research's coverage is to ensure that it represents the most up to date vision of the industry possible.

Changes to the industry taxonomy are built on the basis of extensive research of company, association and competitor sources.

Company coverage is based on three key factors: revenues, products and media attention/innovation/ market potential.

- The estimated revenues of all major companies, including private and governmental, are gathered and used to prioritize coverage; and
- Companies which are making the news, or which are of particular interest due to their innovative approach are prioritized.

GBI Research aims to cover all major news events and deals in the pharmaceutical industry, updated on a daily basis.

The coverage is further streamlined and strengthened with additional inputs from GBI Research's Expert Panel (see below).

12.3.2 Secondary Research

The research process begins with exhaustive secondary research on internal and external sources being carried out to source qualitative and quantitative information relating to each market.

The secondary research sources that are typically referred to include, but are not limited to:

- Company websites, annual reports, financial reports, broker reports, investor presentations and SEC Filings;
- Industry trade journals, scientific journals and other technical literature;
- Internal and external proprietary databases;
- Relevant patent and regulatory databases;
- National government documents, statistical databases and market reports;
- Procedure registries; and

 News articles, press releases and web-casts specific to the companies operating in the market.

12.3.3 Primary Research

GBI Research conducts hundreds of primary interviews a year with industry participants and commentators in order to validate its data and analysis. A typical research interview fulfills the following functions:

- It provides first-hand information on the market size, market trends, growth trends, competitive landscape, future outlook etc;
- Helps in validating and strengthening the secondary research findings; and
- Further develops the Analysis Team's expertise and market understanding.

Primary research involves e-mail correspondence, telephone interviews as well as face-to-face interviews for each market, category, segment and sub-segment across geographies.

The participants who typically take part in such a process include, but are not limited to:

- Industry participants: CEOs, VPs, marketing/product managers, market intelligence managers and national sales managers;
- Hospital stores, laboratories, pharmacies, distributors and paramedics;
- Outside experts: investment bankers, valuation experts, research analysts specializing in specific medical equipment markets; and
- Key opinion leaders: physicians and surgeons specializing in different therapeutic areas corresponding to different kinds of medical equipment.

12.3.4 Expert Panel Validation

GBI Research uses a panel of experts to cross verify its databases and forecasts.

GBI Research expert panel comprises marketing managers, product specialists, international sales managers from medical device companies; academics from research universities, KOLs from hospitals, consultants from venture capital funds and distributors/suppliers of medical equipment and supplies etc.

Historic data and forecasts are relayed to GBI Research's Expert Panel for feedback and adjusted in accordance with their feedback.

12.4 Contact Us

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