



Case Study

Geisinger Leverages Standards for Inventory Visibility

Healthcare leader utilizes UDI information and enriched data for inventory innovations.

Challenge

Geisinger Health, like many large-scale healthcare institutions, used manual processes to store and replenish medical supplies. Because manual processes can be a source of human error, Geisinger made an investment in technology that uses the latest identification standards in medical devices to eliminate manual processes, beginning in areas that use life-saving implants, such as heart catheterization labs.

Solution

Geisinger teamed with Owens & Minor to implement QSight®, a cloud-based inventory management system, to leverage the data in barcodes affixed to medical devices for the U.S. Food and Drug Administration (FDA) Unique Device Identification (UDI) Rule. Geisinger can easily capture and store a product's Global Trade Item Number® (GTIN®) - the most widely used UDI standard - and other critical data with a simple scan.

Benefits

- **Accuracy, Efficiency, Productivity.** Multiple clinical professionals within the healthcare system are no longer tasked with manual record-keeping rife with error. A simple barcode scan places a product into inventory. Subsequent scans track its lifecycle throughout the healthcare system whether it is used (and where), disposed of, returned, or recalled.
- **Profitability.** Inventory management eliminates waste, guaranteeing supplies are not over-ordered and are used prior to expiration. It also delivers fiscal benefits by enabling automated centralized purchasing on behalf of large institutions like Geisinger.
- **Patient Safety.** Inventory management assures patients that any tissue or implant used in their treatment is readily available. It allows clinicians to interface with both the electronic medical record (EMR) and an individual's electronic health record (EHR) with detailed record-keeping for more accurate documentation.
- **Future-Proof Planning and Implementation.** Through its inventory management initiative, Geisinger has established a baseline for system-wide data quality that will enable future operations and initiatives, including accurate and efficient recall management and advanced analytics related to cost and patient outcomes.

Geisinger Health has been a leader in healthcare innovation, seeking solutions to modern healthcare challenges and adopting technologies in service to its patients, caregivers, students, and community. This commitment to innovation is one of the reasons Geisinger Health can boast more than a century of service to its Pennsylvania communities, and why research and consulting firm Gartner consistently ranks Geisinger in the Healthcare Supply Chain Top 25.

What truly sets Geisinger apart is its exceptional due diligence in leveraging every possible capability from its supply chain. At Geisinger, any new technology implemented to meet a challenge is scrutinized for other features that can introduce efficiencies to other departments, other practices, other facilities.

“Technology right out of the box might have more capability than you initially need, but you need to create a roadmap for enabling other features that will increase the performance of the business,” says Kevin Capatch, director of process engineering at Geisinger Health.

A case in point is Geisinger’s comprehensive data and inventory management program that uses increasingly critical supply chain data to drive patient safety within its areas of care. Geisinger leverages the same codes required by the U.S. Food and Drug Administration (FDA) regulations for unique device identification (UDI) and automatic identification and data capture (AIDC) encoded on medical devices to identify products in inventory.

Pencil and Paper to Keyboard

“Over a decade ago, I watched as a well-intentioned cardiac catheterization technician managed his inventory storage area using paper and pencil, then keyed his shopping list into an enterprise resource planning (ERP) system template to reorder,” says Capatch.

Manual processes like these were rife with potential for error and were occurring in every procedural area of the Geisinger system. Moreover, hospital technicians might be tempted to over-order or unintentionally pass over products close to expiration, costing the institution valuable resources – or worse yet – facing an out-of-stock on a critical device needed by a patient.

And with no system to manage the dates, expirations were inevitable. “The combination of capturing information with a barcode scan and having a system that can use and act on that captured information has allowed Geisinger to turn the tide on managing expirations in our procedural areas.”





“You must enforce [standards] in all disciplines; internally in your technology, in your business processes; externally with your vendor community and your solution providers.”

Kevin Capatch
Director of Process Engineering, Geisinger

After this encounter, Geisinger opted to introduce technology that would eliminate manual processes governing inventory and eliminate the three “nevers” in healthcare supply chains: never run out of a critical product, never waste inventory due to product expiration, and never fruitlessly search for recalled items.

Geisinger teamed up with Owens & Minor, a global healthcare solutions company, to use its QSight® inventory management platform.

“QSight gives us product visibility from the moment a product comes into Geisinger, until it is either used with a patient, expired, disposed of, returned, or recalled,” Capatch says. “It brings inventory control, it brings functionality for recalls, it brings expiration management, it brings lot and serialization control – the inherent benefits we set out to acquire by instituting an inventory management program.”

A Standard of Supply Chain Care

Data standards development had been underway for a few years when Geisinger and O&M began working together. In fact, data standards development continues to this day throughout health care and adjacent industries. As Capatch points out: “You must enforce [standards] in all disciplines: internally in your technology, in your business processes; externally with your vendor community and your solution providers.”

GS1 Standards – including Global Trade Item Numbers (GTINs) and Global Location Numbers (GLNs) – are the preferred and prevalent standards used by the healthcare community, pharmaceutical firms, medical device manufacturers and distributors, and are accredited by the FDA to support UDI implementation for all classes of medical devices. Eighty-seven percent of all UDI submissions utilize GS1 Standards, which work in conjunction with standards unique to health care¹. But many entities in the healthcare ecosystem have yet to adopt GTINs and GLNs, which adds a level of complexity to any inventory management effort.

“We are proponents of GS1 Standards. If a vendor is on the fence, or deciding about upgrading business systems, we strongly encourage them to move to GS1 Standards. Theirs are more global. It’s a lot easier to interpret [GS1] application identifiers. This was an integral part in choosing our solution.” says Capatch.

¹ Health Industry Bar Code (HIBC) standard is used to produce uniform data transfer for patient safety and unique device identification (UDI). An ISBT code is an international standard for the transfer of information associated with blood transfusion, cellular therapy and tissue transplantation.

Quality of Data Care

Data quality is imperative for optimal performance: it allows for seamless digital communications among systems, allows UDI data supplied by manufacturers to be used with confidence, and it powers inventory management.

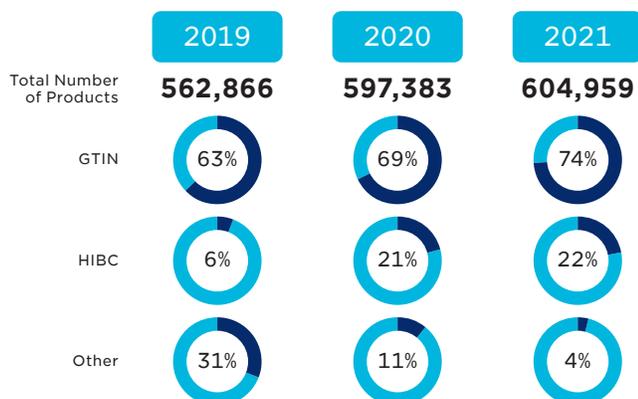
“The importance of data quality is incalculable. Without clean data and a ‘single version of truth’ for all products coming into your system, you are hamstrung in what you can accomplish,” Capatch says. “That is why we benefitted from teaming up with a solution partner that made an investment in managing the item master data using standard identifiers where available, and we did not have to duplicate their efforts, just leverage them.”

“When working with a new client, the first thing we do is data cleansing,” says Vicky Lyle, vice president of Industry Associations at Owens & Minor. “We go into a [customer] department, scan each instance of every single product in that department to make sure the product exists in the global item master. In the instance where a product is not in the database, we utilize the GTIN to source the data from the GUDID [Global Unique Device Identification Database] or directly from the manufacturer’s site.”

Affiliates within the Geisinger system use QSight’s highly enriched item master database. A scan of GS1 barcodes on any given facility’s stockroom shelf will yield a product match of up to 95 percent. Once a product is added in the global item master, it is available for any customer that scans GS1 barcodes.

In the age of Big Data, hospitals are faced with collecting and interacting with massive amounts of data for thousands of products, most of which carry enriched attributes that go beyond manufacturers’ production information. A standards-based inventory management system, provides a secure repository and communications nucleus, bringing value to healthcare facilities like Geisinger needing to leverage the data in several vital ways, including patient safety.

Distribution of Device Identifiers in QSight®



Currently, QSight has over 600,000 stock keeping units (SKUs) in its cloud-enabled database, which is available to all customers, with 74 percent of SKUs tied to a GS1 Global Trade Item Number® (GTIN®). QSight allows for flexibility so

that these alternate identifiers can be “tied” to the GS1 GTIN, when necessary, and supports the industry transition from proprietary identifiers to the standards-based UDI. QSight enhances its databases constantly when additional products are scanned by customers. And because it is cloud-based, the collective information is available to all customers as the scanning takes place, consistently recognizing a product via the GTIN encoded as the unique device identifier. That the constant evolution of QSight data is shared among all users benefits everyone.

“When the standard is present, a match happens almost effortlessly. You don’t have people looking at descriptions because part numbers don’t matter.”

Kevin Capatch
Director of Process Engineering, Geisinger

“That’s where we’re starting to see some gains in using systems that are built on GS1 Standards,” Capatch says. “We can use standards to figure out what items we all have in common.”

The expansiveness of GS1 Standards provides a means of capturing specialized attributes that are crucial in tissue and bone implant procedures. A hip replacement appliance must include the side of the human body into which it is intended for use, for instance; tissue grafts may require information taken from a visual inspection or preparation of the biological material. The need for greater specificity to populate medical practice records as well as individual medical histories is satisfied using standards-based inventory management.

From Dock to Doc

The enterprise resource planning (ERP) system that purchases products is primarily quantity-based. ERP confirms receipt of a carton of 12 stents, for example, but does not note the 12 individual serial numbers or differing expiration dates. By making the connection via QSight between the quantity received and the enriched data connected to those products, visibility and inventory integrity is achieved.

For example, in the cath area, the receiving dock creates a delivery ticket using ERP data; QSight matches the ERP data to the requesting specialty inventory analysts in the cardiac catheterization lab expecting delivery of the stents. Each stent is scanned individually – each with its own serial number along with its lot number and expiration date. From that moment on, Geisinger has visibility into each stent until it is used on a patient, is expired, or is returned to inventory because it was ordered in a case, but a larger stent was needed during the medical procedure.

A scan of the barcode will have captured everything about that stent and Geisinger’s inventory will reflect every step in the lifecycle of that product: ERP uses the incoming

shipment's GTINs to match the source to the order placed. The cath lab personnel captures critical attributes - size, expiration data, lot/batch number - in addition to the unique identifier to link back to the manufacturer in the event of a recall. And nurses who scan the individual stent's barcode in the procedure room allow the metadata to automatically populate the electronic medical record (EMR) and an individual's electronic health record (EHR).

“We are proponents of GS1 standards. If a vendor is on the fence, or deciding about upgrading business systems, we strongly encourage them to move to GS1 Standards. There are a more global. It's a lot easier to interpret [GS1] application identifiers.”

Kevin Capatch
Director of Process Engineering, Geisinger

Although the industry continues to advance in standards adoption, exceptions still arise, so support of master data management by allowing for the recognition of products that are new to Geisinger is crucial. Product procurement can bypass ERP systems entirely, for instance. Stents from a new supplier might be ordered directly by the specialty department needing them (i.e., the cath lab). When the stents arrive at the point-of-care, the lab technicians scan the GS1 barcodes for supporting data for the devices, effectively introducing them into the master data system and placing them in the inventory management system.

Not only will the stents automatically be recognized at every subsequent juncture when scanned within the hospital, the data is automatically available within QSight where its cloud-based model of data distribution makes it available to all customers simultaneously, consistently linking the same unique identifier or GTIN encoded in the barcode to the product.

“When a barcode scan matches, almost everything else happens effortlessly,” Capatch says. “You don't have people looking at descriptions, and other identifies on the packaging, you know the product is in your catalog, and can become trackable in inventory.”

The Proof is in the Inputting

Geisinger has successfully integrated inventory management fully into all non-OR procedural areas - including the catherization (cath labs), electrophysiology, and interventional radiology labs.

“The cath lab was first, but we didn't have to do a lot of convincing after the first implementation: ‘That's so much better than my paper-and-pencil grocery list inventory system!’ and ‘I don't have to worry about expirations anymore because the system tells me you're 90 days out on this item’ were common reactions from other departmental areas within Geisinger. Departments lined up. It was virtually self-marketing for the next installation,” Capatch says.

While the three initial practice areas are using QSight for 100 percent of their product tracking, inventory management, and data governance, Geisinger is now concentrating on bringing the operating rooms managing tissue and implants into full utilization.

“ORs are a little bit more challenging. You're getting into a world where the identifier may have been on the packaging of the non-sterile item, and that identifier gets removed when it's put into the sterilizing case. So, we are figuring out how we're going to capture data on things like non-sterile implants,” Capatch says.

Capatch adds, “Without end-to-end inventory management, a nurse in the OR might have to employ multiple tools to scan multiple implants, which is not just bothersome, it's potentially dangerous. And without standards, none of the automatic identification of products is possible.”

“Without end-to-end inventory management, a nurse in the OR might have to employ multiple tools to scan multiple implants, which is not just bothersome, it's potentially dangerous. And without standards, none of the automatic identification of products is possible.”

Kevin Capatch
Director of Process Engineering, Geisinger

Geisinger Health's Advice on Inventory Management

From Kevin Capatch, Director of Process Engineering, Geisinger Health



Resources

Organizations like GS1 and GS1 US® are working to eliminate pain points. This is not a situation where you can't get there because something hasn't been figured out.

It's been figured out.



Collaboration

Having a technology partner that's **willing to adapt** and modify an offering as you encounter new challenges is essential.



Timing

We will all get there, it's just a timing issue. **And the time to get started is now.**

The Wide View

Enabled by standards-based master data and the advanced data analytics available in QSight, Geisinger can drill-up to gain supply chain insights or drill-down to the case level to get the information the healthcare system needs, aggregated in the way it chooses.

Geisinger's extensive inventory management efforts provide the underpinning for a system-wide recall process. The system's assimilation of GS1 Standards facilitates visibility – not just into individual products in inventory – but into the network's entire operations by supplying reliable parallel comparisons – the kind of analytics needed to improve performance, which in the case of health care, can have profound consequences such as patient safety, expense management, and reimbursements. Standards open the door even wider: multiple hospitals can team-up to conduct life-saving research into health outcomes when apples-to-apples analyses are achieved through standards. These forward-thinking initiatives on Geisinger's roadmap will all be enabled by leveraging data made available by scanning the barcodes on medical devices.



About the Organizations



About Geisinger

Geisinger Health is a healthcare system located in central Pennsylvania serving one million people. Founded more than 100 years ago by Abigail Geisinger, the system now includes 10 hospital campuses, a health plan with more than half a million members, a Research Institute and the Geisinger Commonwealth School of Medicine. With nearly 24,000 employees and more than 1,600 physicians on staff, Geisinger boosts its hometown economies in Pennsylvania by billions of dollars annually. www.geisinger.org



About Owens & Minor

Owens & Minor, Inc. (NYSE: OMI) is a Fortune 500 global healthcare solutions company integrating product manufacturing and delivery, home health supply, and perioperative services to support care through the hospital and into the home. Owens & Minor drives visibility, control and efficiency for patients, providers and healthcare professionals across the supply chain with proprietary technology and solutions, an extensive product portfolio, an Americas-based manufacturing footprint for personal protective equipment (PPE) and surgical products, as well as a robust portfolio of products and services for patients managing chronic and acute conditions in the home setting. Operating continuously since 1882 from its headquarters in Richmond, Va., Owens & Minor is a 140-year-old company powered by more than 20,000 global teammates. www.owens-minor.com

About GS1 Healthcare US

GS1 Healthcare US® is an industry group that focuses on driving the adoption and implementation of GS1 Standards in the healthcare industry in the United States to improve patient safety and supply chain efficiency. GS1 Healthcare US brings together members from all segments of the healthcare industry to address the supply chain issues that most impact healthcare in the United States. Facilitated by GS1 US, GS1 Healthcare US is one of over 30 local GS1 Healthcare user groups around the world that supports the adoption and implementation of global standards developed by GS1. www.gs1us.org/healthcare



About GS1 US

GS1 US®, a member of GS1® global, is a not-for-profit information standards organization that facilitates industry collaboration to help improve supply chain visibility and efficiency through the use of GS1 Standards, the most widely used supply chain standards system in the world. Nearly 300,000 businesses in 25 industries rely on GS1 US for trading-partner collaboration that optimizes their supply chains, drives cost performance and revenue growth while also enabling regulatory compliance. They achieve these benefits through solutions based on GS1 global unique numbering and identification systems, barcodes, Electronic Product Code-based RFID, data synchronization, and electronic information exchange. GS1 US also manages the United Nations Standard Products and Services Code® (UNSPSC®). www.gs1us.org

Proprietary Statement

This document contains proprietary information of GS1 US. Such proprietary information may not be changed for use with any other parties for any other purpose without the expressed written permission of GS1 US.

Improvements

Improvements and changes are periodically made to publications by GS1 US. All material is subject to change without notice. Please refer to GS1 US website for the most current publication available.

Disclaimer

Except as may be otherwise indicated in specific documents within this publication, you are authorized to view documents within this publication, subject to the following:

1. You agree to retain all copyright and other proprietary notices on every copy you make.
2. Some documents may contain other proprietary notices and copyright information relating to that document. You agree that GS1 US has not conferred by implication, estoppels, or otherwise any license or right under any patent, trademark, or copyright (except as expressly provided above) of GS1 US or of any third party.

This publication is provided “as is” without warranty of any kind, either express or implied, including, but not limited to, the implied warranties of merchantability, fitness for a particular purpose, or non-infringement. Any GS1 US publication may include technical inaccuracies or typographical errors. GS1 US assumes no responsibility for and disclaims all liability for any errors or omissions in this publication or in other documents which are referred to within or linked to this publication. Some jurisdictions do not allow the exclusion of implied warranties, so the above exclusion may not apply to you.

Several products and company names mentioned herein may be trademarks and/or registered trademarks of their respective companies. GS1 US does not, by promulgating this document on behalf of the parties involved in the creation of this document, represent that any methods, products, and/or systems discussed or recommended in the document do not violate the intellectual property rights of any third party. GS1 US has not performed a search to determine what intellectual property may be infringed by an implementation of any strategies or suggestions included in this document. GS1 US hereby disclaims any liability for any party's infringement of intellectual property rights that arise as a result of any implementation of strategies or suggestions included in this document.

This publication may be distributed internationally and may contain references to GS1 US products, programs, and services that have not been announced in your country. These references do not imply that GS1 US intends to announce such products, programs, or services in your country.

GS1 US shall not be liable for any consequential, special, indirect, incidental, liquidated, exemplary, or punitive damages of any kind or nature whatsoever, or any lost income or profits, under any theory of liability, arising out of the use of this publication or any content herein, even if advised of the possibility of such loss or damage or if such loss or damage could have been reasonably foreseen.

GS1 US HEREBY DISCLAIMS, AND YOU HEREBY EXPRESSLY RELEASE GS1 US FROM, ANY AND ALL LIABILITY RELATING TO YOUR COMPLIANCE WITH REGULATORY STANDARDS AND LAWS, INCLUDING ALL RULES AND REGULATIONS PROMULGATED THEREUNDER. GS1 US MAKES NO WARRANTIES OF ANY KIND RELATING TO THE SUITABILITY OF THE GS1 STANDARDS AND THE SPECIFIC DOCUMENTS WITHIN THIS PUBLICATION TO COMPLY WITH ANY REGULATORY STANDARDS, LAWS, RULES AND REGULATIONS. ALL INFORMATION AND SERVICES ARE PROVIDED “AS IS.”

*GS1 US employees are not representatives or agents of the U.S. FDA, and the content of this publication has not been reviewed, approved, or authorized by the U.S. FDA. The following information contained herein is for informational purposes only as a convenience, and is not legal advice or a substitute for legal counsel. GS1 US Inc. assumes no liability for the use or interpretation of the information contained herein.

No Liability for Consequential Damage

In no event shall GS1 US or anyone else involved in the creation, production, or delivery of the accompanying documentation be liable for any damages whatsoever (including, without limitation, damages for loss of business profits, business interruption, loss of business information, or other loss) arising out of the use of or the results of use of or inability to use such documentation, even if GS1 US has been advised of the possibility of such damages.

IAPMO

In this publication, the letters “U.P.C.” are used solely as an abbreviation for the “Universal Product Code” which is a product identification system. They do not refer to the UPC, which is a federally registered certification mark of the International Association of Plumbing and Mechanical Officials (IAPMO) to certify compliance with a Uniform Plumbing Code as authorized by IAPMO.

*If applicable

GS1 US Corporate Headquarters

Princeton South Corporate Center, 300 Charles Ewing Boulevard
Ewing, NJ 08628 USA
T +1 609.620.0200 | E info@gs1us.org
www.gs1us.org

Connect With Us



© 2022 GS1 US All Rights Reserved
GDTI: 0614141028993v1.0