PROJECT SONAR (PS)

Advanced Alternative Payment Model

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Project Sonar Abstract

Project Sonar (PS) is a care management program developed by community-based physicians in partnership with a major payer to improve the management of patients with chronic disease. The key to the success of PS is the combined use of evidence based medicine coordinated with proactive patient engagement. The goal of PS is to move physicians from a dependency on fee for service medicine into value based practice. The initial chronic disease category chosen by PS was Inflammatory Bowel Disease (IBD), a family of disorders that are high cost and high risk with a frequency that has been increasing over the past few decades.

In addition to high cost and high risk, CD is also associated with a high variability in outcome and cost. We term this combination of factors as “High Beta” and believe that chronic illnesses can be stratified into high beta and low beta based upon an analogy from the financial industry.

The essential features of PS for the management of patients with chronic conditions, a PFPM, are:

- Evidence Based Guidelines are used to direct the course of care. These are embedded into the EMR through use of CDS tools
- All patients are risk assessed using a set of biopsychosocial measures
- All patients are enrolled in a web-based communication platform; if not web- or smartphone enabled, they are engaged by phone calls from the NCM
- Every patient is proactively ‘touched’ at least once a month; more frequently as needed
- A team based care model has been incorporated into the practice
- Clinical and financial data are analyzed
- The care pathway is continually refined through the development of care management algorithms
- We intervene before patients even realize they need care
PTAC Proposal – Project Sonar

1. Background and Model Overview

In the United States, chronic diseases account for an estimated 83% of total U.S. health spending and virtually all (99%) of Medicare’s expenditures are for beneficiaries with at least one chronic condition.\(^1\)

Project Sonar (PS) is a care management program developed by community-based physicians to improve the management of patients with chronic disease, using evidence based medicine coordinated with proactive patient engagement. Project Sonar was initially deployed in 2012 by the Illinois Gastroenterology Group (IGG), a 50-physician practice with locations in Cook, Lake, Kane, and DuPage counties. IGG is the largest single-specialty, non-academic, Gastroenterology practice in Illinois. As envisioned by the leadership of IGG, the goal of PS was to move physicians from a dependency on fee for service medicine into value based practice. As part of this exercise, IGG analyzed their practice demographics by conditions / diagnoses, procedures, age, payor mix, etc. and confirmed that the most significant chronic disease in a community-based Gastroenterology (GI) practice is Inflammatory Bowel Disease (IBD), which includes Crohn’s Disease and Ulcerative Colitis. When factoring in the costs of pharmaceuticals, whether physician or patient administered, IBD is responsible for almost 25% of the services generated by IGG and likely most other community-based GI Groups. IBD disorders are high cost (due to hospitalizations for complications and use of biologic medications) and high risk (loss of intestine, infections, development of cancers and extra-intestinal manifestations) with a frequency that has been increasing over the past few decades. After evaluating other chronic conditions managed by a community-based GI practice, such as GERD / Barrett’s esophagus, celiac disease, and chronic pancreatitis, the group determined that Crohn’s Disease (CD) was most appropriate for the development of a chronic care management program.

In addition to high cost and high risk, CD is also associated with a high variability in outcome and cost. We term this combination of factors as “High Beta”\(^3\) and believe that chronic illnesses can be stratified into high beta and low beta based upon an analogy from the financial industry. In finance, “beta” is a measure of a stock’s volatility in relation to the market\(^4\). The market has a beta of 1.0, and individual stocks are ranked according to how much they deviate from the market. A stock that swings more than the market over time has a beta above 1.0. As beta is a measure of risk, a beta greater than one generally means that the asset is volatile.

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\(^3\) Kosinski L, Brill J. The Promise of Patient Self-Monitoring: An App a Day Won’t Necessarily Keep the Doctor Away. Clinical Gastroenterology and Hepatology, December 2016; 1751-1752
This same analogy can be applied to most chronic illnesses. Examples of other high beta illnesses include asthma, chronic obstructive pulmonary disease, heart failure, end-stage liver disease, and type 1 diabetes, while examples of low beta illnesses include hypertension, diabetes, and hyperlipidemia. There are high beta periods in low beta illnesses.

IGG is contracted with every major commercial payor operating in Northeast Illinois, including Aetna, Blue Cross Blue Shield of Illinois (BCBSIL), Cigna, Humana, and United. Based on their own internal analysis of data, IGG met with all the payors, and BCBSIL was willing to work with the group to analyze this issue. A review of commercial claims for 21,000 patients with a ICD-9 diagnosis consistent with CD for the years of 2010 and 2011 revealed:

- The annual cost per patient with a diagnosis of CD was $11,000 (2011 data)
- The overall hospitalization rate for patients with CD was 17%
- Over 50% of the expenditures incurred were for inpatient care for the treatment of complications of CD
- Although 1/3 of the total spend was for physician services, only 3.5% of the total spend was for gastroenterology care.
- Over 2/3 of the patients who were admitted to a hospital for a complication related to CD had no identifiable contact (based on claims data) with a provider in the 30 days prior to the admission

Based on this data, we asked the following questions:

- Is there a way to decrease the cost of care of patients with CD by decreasing the complication rate through better medical management?
- Is there a way to identify the high-risk patient with CD before complications ensue?
- Is there a way to channel the care of these patients to those healthcare professionals who have the most knowledge, experience and expertise?
- Is there a better way to engage our patients so that their early warning signs can be assessed even before they realize they need intervention?

While IGG was analyzing and researching these questions, the American Gastroenterological Association developed and published a care pathway for the management of Crohn’s Disease. AGACDCP\(^5\). The recommendations in the pathway are as follows:

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To reduce potentially avoidable hospitalizations and to control the expense of those who are hospitalized, physicians must work with patients to control complications, which requires physicians to identify the high-risk patient and control their inflammation. This includes a biopsychosocial risk assessment that will identify the biological markers of risk and assess the patient for anxiety/depression along with their social environment.

The ensuing management requires a team-based approach deploying Clinical Decision Support (CDS) tools to guide healthcare professionals on optimal care, appropriate use of anti-tumor necrosis factor (TNF) and other biologic medications, and incorporation of Nurse Care Managers (NCMs) as the focal point in the team that was engaging with and managing the patient. Importantly it requires the engagement of the patients with the use of “hovering tools” to non-intrusively monitor the patient’s symptoms, in effect creating a “sonar system” to ping them in their usual environment on a periodic basis. Collectively, the goal of PS was to engage those patients who might otherwise minimize the severity of their disease, identify patients who were depressed and/or at risk of decompensation, and optimize patients on appropriate pharmaceutical treatment as early as possible, with the objectives of improving patient quality of life and decreasing costs through reducing potentially avoidable complications, emergency department (ED) visits, and inpatient (IP) admissions.

PS was deployed in IGG in 2013, initially in a pilot study population of 50 patients with CD. CDS tools designed around the AGACDCP and a hovering tool using the Crohn’s Disease Activity Index (CDAI) were deployed into IGG’s NextGen electronic medical record (EMR) system. Using the EMR practice portal, patients were sent a subset of questions derived from the CDAI at the beginning of each month. This subset of the CDAI returned a score (Sonar Score) that applied a numeric value to the patient. At the end of the first year, the hospitalization rate for these patients decreased from 17% to 5%.
IGG presented the data from this pilot to BCBSIL. Based on the findings from the pilot, BCBSIL partnered with IGG to create its first specialty-based Intensive Medical Home (IMH). Using a model similar to what it had deployed in its primary care PCMH, BCBSIL attributed 303 patients with CD to IGG of which 185 were enrolled, and the IMH project, which went live on Dec. 1, 2014 was based on the clinical structure described above. The practice receives a supplemental per member per month (PMPM) payment to cover the infrastructure for participating in PS. The PMPM payment is additional to fee for service payment and varies on an annual basis, adjusted based upon mutually agreed-upon goals for the clinical and financial performance of the patients enrolled. The practice receives one year of historical claims data on its attributed patients and quarterly medical and pharmacy claims data going forward. The practice is responsible for calculating the performance of physicians at an individual, strategic business unit (SBU), and practice level, including comparison of BCBSIL patients depending on their level of engagement (‘pinger’ vs. ‘nonpinger’) and comparison to patients enrolled in other commercial and government (Medicare, Medicaid, Tricare) programs.

Early on, it became clear that the EMR patient portal would not sustain the level of patient engagement required for adequate care. It was replaced with a web and mobile-based platform which includes interactive text messages to patients, and patient response data which drops directly into the EMR using a HIPAA-compliant HL7 interface. Patient engagement increased from 27% to 75-80% which has now been sustained for over 20 months.

The initial first full year of clinical, financial, and patient engagement data for PS are shown in the graphic below which was presented as a Distinguished Abstract at Digestive Disease Week 2016:

The data below are normalized to Medicare payment to emphasize true utilization rates. They show:

- A net decrease in cost of 9.87% even with an 8.97% increase in infusible biologics and Net of PMPM payments to the practices
- A 57.14% decline in inpatient costs driven by an equivalent decline in admissions/complications
These financial results are highly dependent upon the patient engagement. When we filter our patient population by those who respond to at least 50% of their monthly surveys (pingers) vs. those who do not respond (nonpingers), the cost differential of care falls 18% in the Pinger group whereas it rises 23% in the nonpinger group. This 41% difference in cost is shown in the figure below.

Thus, the cost savings is highly correlated to patient engagement, as it is derived from those patients who respond. Patient engagement is key to the success of PS and for the care of patients with chronic disease. Our results, which we believe are generalizable to other high-beta chronic conditions and to high beta periods in low beta conditions, lead to a model that must incorporate the use of evidence based medicine and, more importantly, must provide for ongoing patient engagement which is integrated into the workflow of care.

Key to the success of a management program for a chronic condition must be the use of appropriate risk measures. Every patient in PS is initially assessed using the risk assessment tool embedded in the AGACDP, which includes 26 biopsychosocial risk metrics in three categories: Inflammation Risk, Disease Burden Risk and Comorbidity Risk. Multiple linear regression analyses of each risk measure against the Crohn’s Related Cost of Care has helped us to identify which measures hold predictive value. This has led to further refinement of the processes within PS.

A process overview of PS is shown below:
The essential features of PS for the management of patients with chronic conditions, an APM, are:

- Evidence Based Guidelines are used to direct the course of care. These are embedded into the EMR through use of CDS tools
- All patients are risk assessed using a set of biopsychosocial measures
- All patients are enrolled in a web-based communication platform; if not web- or smartphone enabled, they are engaged by phone calls from the NCM
- Every patient is proactively ‘touched’ at least once a month; more frequently as needed
- A team based care model has been incorporated into the practice
- Clinical and financial data are analyzed
- The care pathway is continually refined through the development of care management algorithms
- We intervene before patients even realize they need care

2. **Scope of Proposed APM**

This proposal to the PTAC will expand CMS’s APM Portfolio in the following ways:

- It will move providers, especially specialists, from fee for service to reimbursement based on value based payment methodologies. It is addressing an issue in payment policy in several ways:
  - Moves patient care from reactive to proactive.
  - Focuses on early identification of potential problems and complications, encouraging ‘preventive’ management
  - Rewards physicians and other qualified healthcare professionals for ‘doing the right thing’ in a team-based manner, as opposed to reimbursement solely on an RVU-based methodology.
- It will promote “cost sensitivity” in specialist providers who are currently almost exclusively paid on a fee for service basis. This will promote the inclusion of APM entities for specialists whose opportunities to participate in APMs has been limited.
- It will link payment for specialist services to clinical, financial, and patient reported outcomes.
- It will allow specialists to participate in value based care outside of an ACO / MSSP / CRC+ model.
- It will allow physicians to participate in value based care for chronic conditions that are not triggered by a surgical procedure on an inpatient or outpatient basis.

Specialists want to be part of the value based solution but have not been fully able to exert the powerful force they have on cost containment and care improvement. The majority of CMS / CMMI initiatives to date have focused on ACOs, primary care based models, or conditions
triggered by a hospital procedure (orthopedic, cardiac). This has left the specialist out of the solution and kept them as part of the problem.

The patient has also been left out of the solution. Few value based models built around ACOs maintain a focus on patient engagement as a value-added solution. This is exacerbated by initiatives based on hospitalization triggers which do not have an outpatient focus on chronic management.

Most serious chronic illnesses are managed by specialists who are still paid under fee for service. **PS shows how a specialty group in partnership with a major payer can move from FFS to value-based care and demonstrate cost-savings with improved patient quality of life. We believe this model, which has been proven in patients with IBD, is applicable to other conditions.**

Although PS focused on CD, this model can be applied to other chronic illnesses. As noted earlier, we believe that chronic illnesses can be stratified into high beta and low beta. Some chronic diseases, like diabetes mellitus, hypercholesterolemia and hypertension, can be slow and indolent in the expression of their symptoms and morbidities, which we term as “low-beta” diseases. Other conditions, such as inflammatory bowel disease (IBD) are “high-beta” conditions that are not as forgiving in their disease progression. There are several high-beta conditions including asthma, heart failure, chronic obstructive pulmonary disease, end-stage liver disease, rheumatoid and psoriatic arthritis, cancers, malnutrition and other conditions where patients are at risk for frequent, potentially avoidable, emergency department visits and hospitalizations. Even diseases like diabetes may have high beta periods, such as the period following an admission for diabetic ketoacidosis or when patients have poor control of / high fluctuations in blood sugar / hemoglobin A1c levels. Although most of the chronic care of patients with low beta diseases can safely be provided by PCPs, the intervening high beta periods most often require the care of a specialist. The PS platform can be applied to high beta illnesses as well as the high beta periods embedded in the management of low beta chronic diseases.

We believe that PS has applicability to many patients with chronic disease and would be appropriate for use by primary care and specialist physicians and qualified healthcare professionals, both cognitive and procedural.

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3. Quality and Cost

Value can be defined as Quality/Cost. Since Quality = Outcome, value can be further defined as Outcome/Cost. The perception of care is essential to the measure of value, so the most appropriate definition of value is: Value = (Patient Outcome + Patient Satisfaction)/ Patient Cost.

\[
\text{VALUE} = \frac{\text{Patient Outcome} + \text{Patient Service}}{\text{Patient Cost}}
\]

The triple aim’s value today has refocused us on Population Health. So, the previous patient based equation applied to a population would be:

\[
\text{VALUE} = \frac{\text{Population Outcome} + \text{Population Service}}{\text{Population Cost}}
\]

PS applied to patients with chronic disease can provide this value through improving quality and lowering cost. Quality can be improved through:

- Use of Evidence Based Guidelines to support a focus on outcomes.
  - PS has demonstrated a significant decrease in complications, hospital admissions and emergency department visits using a model which is applicable to other chronic conditions.
- Improved Patient Satisfaction through targeted patient engagement.
  - Patients engaged through PS maintain a very high level of satisfaction with the program. They are benefitting from “Concierge Medicine” without the extra expense.
- Proactive Patient Engagement

Proactive patient engagement coupled with clinical surveillance and intervention is critical to the successful management of patients with chronic disease. Patients with chronic disease often surface only when they recognize they are in trouble and realize they cannot repair their situation themselves. Patients with high-beta diseases can rapidly deteriorate, resulting in hospitalization and complications.\(^9\)\(^10\). Patients with CD frequently minimize their own


deteriorating symptoms; they may present too late at which time morbidity has occurred and significant medical and/or surgical care must be provided. PS has demonstrated that patient engagement is critical to avoiding these occurrences.

4. Payment Methodology

The payment model of PS represents negotiation of a transition from fee-for-service to fee-for-value for physicians who may not ready or able to take on full risk, but are capable and should be ready to accept accountability for the care they provide. PS will facilitate the development of accountability for the care of the patient with chronic disease and will promote a decline in the variability in the cost of care of these patients.

The Structural Model of PS is detailed below:

**Project Sonar Structural Model**

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<thead>
<tr>
<th>Provided by Payer</th>
<th>Provided by the Practice</th>
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<tbody>
<tr>
<td>• Attribution list of patients</td>
<td>• Clinical Team</td>
</tr>
<tr>
<td>• Data</td>
<td>o Medical Leadership</td>
</tr>
<tr>
<td>o Baseline claims data for 12 months prior</td>
<td>o Nurse Care Managers: One NCM FTE per 200 patients</td>
</tr>
<tr>
<td>o Quarterly claims data going forward</td>
<td>o Patient Care Coordinator</td>
</tr>
<tr>
<td>• Payments</td>
<td>• Initial Care requirements</td>
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<tr>
<td>o Enrollment Visit: $200 (NCM Visit)</td>
<td>o Every patient to have an Enrollment Visit – NCM</td>
</tr>
<tr>
<td>o Care Management Payments $70 PMPM</td>
<td>o Program Goals and their Barriers</td>
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<tr>
<td>• Performance Measurements – Annually assessed</td>
<td>o Anxiety Depression Assessment</td>
</tr>
<tr>
<td>o Outpatient Medical Costs</td>
<td>o Action Plan</td>
</tr>
<tr>
<td>o Inpatient Cost</td>
<td>o Physician Visit</td>
</tr>
<tr>
<td>o ER Visit Cost</td>
<td>• Review of Enrollment Visit</td>
</tr>
<tr>
<td>o Biologic Cost</td>
<td>• Risk Assessment</td>
</tr>
<tr>
<td>o Infused Biologic Costs</td>
<td>• Review of Action Plan</td>
</tr>
<tr>
<td>o Injected Biologic Costs</td>
<td>• Ongoing Care requirements</td>
</tr>
<tr>
<td>o Other Cost</td>
<td>o Every patient must be “touched” monthly</td>
</tr>
<tr>
<td></td>
<td>o NCM reviews each patient’s monthly ping and reports to MD</td>
</tr>
<tr>
<td></td>
<td>o MD must respond to needed care</td>
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In the PS APM we are proposing a prospective payment model with retrospective reconciliation. PS is based upon a chronic care management (CCM) model combined with proactive patient engagement. Physicians who voluntarily choose to participate with the model would continue to have their services reimbursed through the MPFS. Target prices are compared to the actual cost of the care provided. Payment adjustments are based on quality and financial performance. In the case of cost savings, the shared savings component of the payment would be paid to the physician entity. In a situation where the physician’s attributed
costs overrun the expected target, the physician would be required to repay losses up to the agreed upon limit in its contract with CMS. To protect against catastrophic losses, the model will build in stop-loss provisions and outlier protections.

Participating providers would also be required to contribute based upon their agreement with PS as the APM entity. A monthly payment for non ‘face-to-face’ services by clinical staff, overseen by the physician, is essential for the successful deployment of the model to cover the infrastructure costs required.

There are initial and ongoing costs for dedicated NCMs and ping coordinators (clinical staff), IT expenses for incorporation of CDS tools into the EMR workflow and deployment of the patient engagement platform and data management. These are detailed below:

- NCM Cost: $25 PMPM
- Ping Coordinator $12 PMPM
- IT Platform $20 PMPM
- Medical Director $2.50 PMPM
- Total Cost $59.50 PMPM

The CCM payment is designed to cover the cost of deployment of the care management infrastructure not to provide supplemental income to the practice. Incentives to the practice will be generated by the sharing of cost savings derived from a retrospective adjust. A Performance Based Adjustment (PBA) is made based upon this retrospective cost reconciliation. If savings are favorable, the PBA will represent a supplemental payment to the practice. This will be capped at 10%. If performance is poor and costs are excessive based on risk adjusted norms, then the PBA would be negative up to a cap of 5%.

This payment structure should be viewed as one designed to promote a continuous process of value generation over time. We anticipate that going forward many more value-based initiatives will be necessary to continue to move the cost curve down. Episode and procedural bundled payments will be necessary as well as consolidation in NCM functions. The goal will be to gradually push cost down until a convergence to an ideal mean can be accomplished.

5. Value over Volume

Volume based drivers have dominated physician practices for the last 50 years. Since “every system is perfectly designed to get the results it gets”[^11], medical practices are designed to “do more”. In primary care practices this correlates to the metric: patients seen and RVU generated per hour. The obvious challenge is to maintain quality of care when the physician or qualified healthcare professional is seeing 4 or more patients per hour, especially if the patient has multiple co-morbid conditions, complex medication regimens, psychosocial issues, etc. There is

[^11]: Conway E, Bataiden P. Institute for Healthcare Improvement; August 21st, 2015
a limit to the productivity of a practice, whether primary care or specialist, which cannot be solved through use of qualified healthcare professionals.

While specialists may be more procedural, they are equally as volume driven. Data from IGG shows that over 54% of the revenue of a community-based GI practice comes from screening (G0105, G0121), diagnostic (45378), and procedural colonoscopy (45380, 45384, 45385), primarily related to colorectal cancer (CRC) screening, diagnosis, and surveillance. Many Gastroenterologists have developed a very efficient focused-factory model for performing this procedure.

Why should a GI move from this model to one based on value? Colonoscopy is a mature service; CMS has adjusted procedure wRVU downwards in CY 2016 and 2017, revenue is declining and costs are rising, compressing margins. Alternative methods for CRC screening are available which, in a population based environment, might lead to a shift from procedural to diagnostic colonoscopy with a corresponding decline in the volume of colonoscopy. Anticipating this trend, Project Sonar was developed to provide GI physicians with an option to improve care based on value based chronic disease management, rather than attempting to perform more procedures.

In our initial review of commercial payer claim data from 2010 and 2011, the average cost/patient with CD was $11,000, which includes professional, laboratory, imaging, hospital / facility, and pharmaceutical costs. The average yearly reimbursement to a GI for managing these patients was $385, which includes only professional fees for office visits, procedures, and supervision of physician-administered infusions. Similar to oncology, physician administered infusions represent an additional source of revenue to the practice, while prescribing a patient self-administered drug would not generate any practice revenue.

The problems in today’s FFS model are obvious and include:

- Physicians are compensated more to do more. Physicians are incented to perform more procedures where the revenue per RVU is higher. By revising the 2017 fee schedule to pay a facility more for colonoscopy procedures (e.g. 45380, 45385) and less for screening / diagnostic procedures (e.g. G0105, G0121, 45378), CMS has potentially created an incentive for physicians who own an ASC to perform more procedural services.
- As the per-minute compensation for cognitive services is less per RVU compared to procedural services, many of these visits have been relegated to qualified healthcare professionals (nurse practitioners, physician assistants) in the practice who might not detect early signs of deterioration in a patient. This creates a paradoxical situation where the most knowledgeable member of the healthcare team is busy performing the most repetitive and least cognitive task (procedures) while the more complex cognitive services are performed by less-trained professionals.
- Prescribing infused biologics to increase practice revenue.
o The patient is seen every 6-8 weeks when they come in for their infusion. If that infusion is performed in an office setting where the physician sees the patient, our data suggests that the hospitalization rate is much lower than if the infusion is performed in a setting (hospital outpatient or, for commercial patient, free-standing infusion center) where the patient is not seen by their provider involved in the ongoing management of their condition.

o Patients do not want to disrupt their lives every six to eight weeks to come in for an infusion. This results in lost productivity at work, use of PTO days and time away from family. For these and other reasons, patients may prefer self-administered drugs, which creates a potential management issue as the patient is not being monitored as they would if they were coming in for an infusion.

o On a total cost basis when looking at the cost of drug and infusion, the cost of infusible biologics is higher than the cost of self-administered agent, which does not consider the lost productivity cost. This must be balanced by the higher hospitalization rate with self-administered agents due to a loss of ongoing communication with the provider.

PS was developed to address these issues. How does PS move this to value?

- The provider in PS is compensated a prospective payment fee in addition to their usual FFS compensation. In our commercial model, this represents approximately $600 per year to the provider, which is not significantly different from the approximately $490 per year were the physician to bill CPT code 99490 (chronic care management). The physician is encouraged to continue office visits for the patient, and to provide procedural services when clinically indicated, such as for CRC screening / surveillance.

- The physician practice is responsible for paying the NCM and ping coordinator and data management out of the monthly CCM payment.

- Expenses for all procedures are paid under a bundled payment methodology.

- Physicians are encouraged to utilize anti-TNF and other biologics based upon the most efficacious agent, not the one that generates the most revenue to the practice. The PS system facilitates patient engagement and communication regardless of the site of service.

Performance data using PS in a commercial population has shown a “Net” 9.87% savings. Based upon the $11,000 cost per patient, this amounts to a savings of over $1,000 per patient. This is net of the monthly CCM payments and is a normalized savings based upon Medicare Payment Rates. The combination of prospective CCM payments which support care infrastructure of NCMs combined with the use of the Sonar Platform facilitate this overall savings and its improvement in the quality of life of its patients.
6. Flexibility

PS represents a flexible and scalable platform upon which to expand its use to most all practices. To date, we have installed the common PS platform in 20 GI practices across the country representing approximately 600 physicians in 12 states, including community and academic practices.

The “glue” that binds the PS practices is the use of the web and mobile-based platform, which represents a single database of provider driven and patient driven metrics. A web-based platform which is both EMR and smartphone device agnostic allows PS to apply changes centrally and distribute automatically to all sites. The results of Sonar Scores are pushed into EMRs as lab data using a HL7 interface.

The PS platform has been used for research on clinical and pharmacological outcomes and effectiveness. For example, the network of PS practices has been leveraged to study the effects of oral therapies for irritable bowel syndrome as well as IBD, and to assess whether assessment of small bowel mucosa correlates with clinical and CDAI findings.

PS requires the practice to change its focus from ‘passive-reactive’ to proactive population health. Instead of a “one patient at a time” passive focus, waiting for the patient to contact the physician with new or intensifying symptoms, the PS practice is focused on the population of patients. The practice, and the patients with a disease process, are actively engaged to improve the care of the patient. This requires infrastructure changes such as the incorporation of NCMs and the ping coordinator.

We believe that a similar care management infrastructure is applicable to other illnesses. We further believe that PS is agnostic to the specialty and is applicable to other chronic illnesses, and can be used by primary care and specialist physicians involved in the care of patients with high beta chronic disease or the high beta periods in patients with low beta conditions.

7. Ability to be evaluated

Evaluation of any new care model is essential to validate its value. PS has been under evaluation since its inception. The following financial metrics are monitored on a regular basis:

- Average total cost/patient
- Average inpatient cost/patient
- Average Emergency Room cost/patient
- Average biologic cost/patient – both infused vs injected

Total cost from a commercial plan is a misleading metric as it must be normalized to Medicare Payment Rates to minimize the site of service differential. That is not to say the site of service is not important, it must be minimized as well. By using a single payment methodology, utilization can be better assessed.
Ultimately the most important outcome is the improvement in the quality of life of the patients with the disease in question. Financial measures alone may not reveal the entire patient benefit. We must always place higher value on the costs incurred for disease morbidity. An example is the cost of inpatient care vs the cost of biologic therapy. Despite the expense of the latter, it does not come with the suffering of the morbidity of disease.

Outcomes driven quality metrics, difficult to develop, have slowed the movement from volume to value. Since PS has a unique access to both quality as well as claims data it has been able to develop predictive models based on how changes in quality metrics effect cost. Utilizing the 26 risk assessment metrics in the AGACDCP, PS ran multiple linear regressions of each metric against the Crohn’s Disease Cost of Care. This has allowed for the development of a mathematical model which assigns a relative strength to each metric with respect to its ability to control the cost of care.

<table>
<thead>
<tr>
<th>CDCP Risk Factor</th>
<th>Odds Ratio</th>
<th>95% Confidence Limits</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td>Lower Limit</td>
</tr>
<tr>
<td>Inflammation risk: Albumin</td>
<td>19.4</td>
<td>3.9</td>
</tr>
<tr>
<td>Inflammation risk: Joint pain</td>
<td>5.7</td>
<td>2.2</td>
</tr>
<tr>
<td>Comorbidity risk: Inflammation</td>
<td>11.5</td>
<td>1.5</td>
</tr>
<tr>
<td>Comorbidity Risk: Restricting</td>
<td>5.4</td>
<td>2.2</td>
</tr>
</tbody>
</table>

Each risk assessment metric has a relative value. Patients can be profiled with this scoring methodology and placed accurately in risk categories. An example would be the use of serum albumin levels as a predictive value for risk. We have shown that changes in serum albumin are predictive of 19.4% of the variation in cost of care for CD\textsuperscript{12}. Patients are monitored quarterly for serum albumin level changes.

We continue to refine this mathematical model, which will benefit from a larger “n” of patients. This model enables rating of each physician and NCM performance on a risk adjusted basis. Ping response rate is critical to the outcome of PS; provider and NCM specific ping response rates can be used as a metric for performance.

We have developed an upside risk revenue share model which creates the proper incentives for physician behavior. Working with physician groups in a variety of practice and geographic settings, we have arrived at the following formula:

- 33%: based on number of patients followed
- 33%: based on the ping response rate
- 34%: based on the risk adjusted cost of care

With additional data and additional participating practices, we believe this formula can and will be adjusted in the future to promote optimal value-based behaviors.

8. Integration and Care Coordination

The success of PS rests on the deployment of a team-based care infrastructure. MD can no longer stand for “My Decision”. To successfully manage the patient with a chronic condition, physicians and qualified healthcare professionals, NCMs, Ping Coordinators and ancillary personnel such as behavioral health and pharmacists are required.

At present, the current care management model for PS centers around the NCM, who is the most important component of the team. Based on data from the Case Management Society of America, each FTE NCM should be able to follow 150-200 patients\(^\text{13}\). Our data shows that using PS’s web and mobile patient engagement platform, this number can be safely increased to 250 patients per FTE NCM. Practice management data indicates that one NCM can provide support to multiple physicians and qualified healthcare professionals in the practice. For solo and smaller practices that might not have the patient volume to support a dedicated NCM, we believe that a shared-service model could support such practices.

The NCM relationship begins at enrollment when the patient comes in for a “Supervisit”. At this visit, patients are introduced to the NCM and queried as to their personal goals for the program. Barriers to accomplishing these goals are identified and an action plan generated. Patients are rated with respect to depression/anxiety using a PHQ-2 tool as these components affect all chronic diseases. Diet histories are generated with action plans for nutritional support. At the conclusion of the initial visit, the patient is seen by the physician or qualified healthcare professional who reviews the details of the Supervisit and all parties sign off on the action plan.

After the initial Supervisit, ongoing management begins. The PS platform provides a desktop to the NCMs where their patients can be monitored together as a population. The NCM follows the Sonar scores of the patients and contacts all patients whose scores fall out of standards, or who does not respond to their ‘ping’ within a predetermined amount of time. Based on the patient responses, if required the NCM goes into the practice EMR and initiates a communication with the physician or qualified healthcare professional. Patients whose scores indicate that they are deteriorating are contacted and engaged with the physician or qualified healthcare professional, either in-person or telephonically. Precision care is being managed proactively and constantly. Patients are engaged and physician services are initiated as needed by the NCM.

All care, from the Supervisit through the ongoing care, is contained in the same relational database. Reports can be generated from queries developed by the PS leadership team. Interactions between the members of the team can be monitored and rated. The platform provides the ability to rate NCMs separately from providers.

Each practice should have a clinical staff member who functions as a “ping coordinator”, monitoring the ping process on a regular basis. This person monitors who is entering and exiting the program.

The issue of integration with PCPs and other specialists needs to be discussed. We see this model as being integrated into the structure of the entire care team. In pure high beta illnesses like CD, the specialist may be the only member of the team that receives the PMPM payment. In high beta periods of low beta conditions, the specialist will receive the PMPM only during the high beta period. The bottom line is that the provider managing the care of the chronic illness in question receives the payment. When comorbidities exist which require management by multiple specialists, then multiple management fees may be necessary.

9. Patient Choice

PS is focused around the needs of the patient. In fact, the focus of PS is the patient. Our data and results have shown that the patients prefer the engaged infrastructure of PS - their NCM functions as their own personal care coordinator, or “concierge care within a managed environment.” Patient satisfaction is high; PS has maintained a 75-80% sustained patient ping response rate over 2 years.

We learned this that hard way. In the first six months of PS, we used a patient portal for our patient communication. This was not ideal and the patients did not like logging in to the portal to obtain their surveys. The EMR portals are cumbersome, inflexible, and slow to align with advancements in technology (e.g. new smartphones and devices used by consumers). In June 2015, we switched to a web and mobile-based platform developed by PS that utilizes smartphone technology, which has been very well received by the patients and is clearly their choice of communication.

For the 20-25% of the patients who either do not have a smart phone or choose not to use one, we use more conventional means of communication. A small number of patients prefer the portal. Telephone calls from the NCM asking the patients the questions in the survey are used when other means are not possible. As we serve a population in whom English might not be the primary language for some patients, PS is in process of translating the questions into Spanish and using ATT Language Line for others. In the end, we “touch” every patient every month.

Approximately 24% of patients are deemed depressed or anxious at enrollment. These patients require a higher level of service which can be provided through PS. As our data shows that
patients with behavioral health issues is associated with a 13% higher cost of care, they are a major focus for the NCM. These patients may receive multiple calls and touches during the calendar month. In the end, PS is the patient’s choice. High levels of patient satisfaction exist.

10. Patient Safety

PS must be considered a Sonar System for patients with chronic disease. It is like looking out at the ocean. Think of the patients as submarines, submerged and running silent and deep. They only surface when they are in trouble which means two things will need to happen for them to surface: they recognize they are in trouble, and realize they can’t get out of trouble on their own. Unfortunately, patients make mistakes on both issues and their safety can be compromised. In high beta diseases like Crohn’s Disease or end-stage liver disease, there is not much room for error. They surface in deep trouble and must be brought in to port (the emergency department) which usually results in hospitalization.

Our data shows a significant improvement in patient safety with implementation of PS, as demonstrated by the significant decrease in emergency department use and hospitalization rate. The close patient engagement built into PS minimizes patient safety issues.

The algorithms which are built into PS are a key component of our success. The NCM desktop is alerted when a patient’s Sonar Scores fall out of safety ranges. These ranges are based upon a continuous process of refinement / development of algorithms:

The SonarMD Algorithm Development Cycle
11. Health Information Technology

PS is built on an HIT platform which utilizes both web and mobile-based technology to communicate with providers, their staffs and their patients. It is cloud-based and agnostic to the operating system of the handheld device as well as the practice EMR. Although not an EMR, the PS platform integrates with all EMRs by pushing sonar scores through to the EMR as lab data using an HL7 Interface.

Patients using the PS platform receive monthly “pings” on the device of their choice: smartphone, tablet, PC, etc. Some patients who do not have access to this technology receive their communication via telephone.

The strength of the mobile platform lies in its ability to provide immediate feedback to the patient. Since it is a web-based platform communicating to a SQL database, algorithmically generated responses can provide patients with immediate feedback. If the patient’s Sonar Score triggers one of the algorithms, the patient is instructed to contact the office of the practice. Concurrently, the NCM in the practice receives the same notification and most often contacts the patient before the patient does. This provides not only an excellent communication tool, it provides the patient with a sense of security that someone is constantly monitoring their condition.

On the practice side, the NCMs utilize a set of desktop templates where they can monitor the performance of each of their patients as well as their assigned patients in aggregate. Sonar Scores are color coded based on algorithmic rules. See Appendix VII for full images. Unlike an EMR where patient charts are opened one patient at a time, the Sonar Platform is designed for population health providing the NCM and the physician with the ability to see all of their PS patients in a user-friendly interface.

Each practice participating in PS has its own sub-platform, thus they cannot comingle data from other practices. At a central level, PS’s “Sonar Central” can pull data into a central repository for query creation and reporting. The ability to aggregate and analyze data from practices
across the United States has in effect created a GI Registry for IBD. PS is able to monitor trends across the population, study clinical factors related to outcomes, and provide guidance and recommendations back to the practices. It is also our vision to use these capabilities to further drive clinical decision support at the time of engagement and evidences based medicine across the specific disease state(s).

Safety and security are maintained through the appropriate use of encryption and use of firewalls. This is shown in the diagram below.

The PS’ IT infrastructure lends itself to being expanded and scaled as necessary to accommodate multiple practices and disorders. Safety and security are maintained through the appropriate use of encryption and use of firewalls. Sonar has a full HIPAA compliance program in place to safeguard access and use of the PS data.

There are currently over 600 in 20 large GI practices across the country who have implemented the SonarMD platform. All of the data from these practices is contained in the above structure in a safe/secure environment. This infrastructure is perfectly positioned to be expanded to other disease processes and practices in other specialties.
12. Supplemental Information

The expansion of Project Sonar beyond the Illinois Gastroenterology Group necessitated the formation of an entity to provide the necessary structure for a national project. SonarMD, LLC is a Delaware LLC founded in 2014 by physicians dedicated to the development of chronic care management tools. Through development of its provider focused CDS tools and its patient engagement digital platform SonarMD provides medical practices with a patient friendly infrastructure for population health.

The medical practices currently using the SonarMD digital platform constitute the SonarMD Medical Group (SMG). Each practice is contracted with SonarMD under identical contractual arrangements, and nominates a member to the SMG advisory board. We anticipate that as PS grows, advisory groups will be organized around specialty and condition.

We have considered whether SonarMD, LLC could be the APM entity that contracts directly with CMS. In this scenario, the risk arrangement for the APM entity including, total risk and stop-loss provisions would be agreed upon in the APM entity’s contract with CMS. Infrastructure expenses would include:

- Development of a CCM payment designed to cover the infrastructure necessary to implement the PS platform: NCMs, Ping coordination, IT Platform
- Mechanisms for claims processing under prospective bundled vs. retrospective payments.
- Claims data reporting
- Quality reporting based upon mutually agreed upon measures and metrics. These measures will include MIPS measures as well as PS derived measures. An example for IBD would be as follows:

<table>
<thead>
<tr>
<th>Category</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>NQF 128</td>
<td>Preventive Care and Screening: Body Mass Index (BMI) Screening and Follow-Up Plan</td>
</tr>
<tr>
<td>NQF 226</td>
<td>Preventive Care and Screening: Tobacco Use: Screening and Cessation Intervention</td>
</tr>
<tr>
<td>NQF 271</td>
<td>Inflammatory Bowel Disease (IBD): Preventive Care: Corticosteroid Related Iatrogenic Injury – Bone Loss Assessment:</td>
</tr>
<tr>
<td>NQF 275</td>
<td>Inflammatory Bowel Disease (IBD): Assessment of Hepatitis B Virus (HBV) Status Before Initiating Anti-TNF (Tumor Necrosis Factor) Therapy</td>
</tr>
<tr>
<td>PHQ 2 Q1</td>
<td>Little Interest or Pleasure in doing things?</td>
</tr>
<tr>
<td>PHQ 2 Q2</td>
<td>Feeling Down Depressed or Hopeless</td>
</tr>
<tr>
<td>Lab</td>
<td>Serum Albumin</td>
</tr>
<tr>
<td>Lab</td>
<td>Serum Hb</td>
</tr>
<tr>
<td>Lab</td>
<td>Fecal Calprotectin</td>
</tr>
<tr>
<td>Lab</td>
<td>C-reactive protein</td>
</tr>
<tr>
<td>Lab</td>
<td>Endoscopic Assessment of Disease Activity</td>
</tr>
<tr>
<td>Sonar</td>
<td>Ping Response Rate</td>
</tr>
<tr>
<td>Sonar</td>
<td>Average Sonar Score</td>
</tr>
</tbody>
</table>

- Financial reporting measures as previously addressed
Appendices

I. Abstract: Validation of American Gastroenterological Association’s Crohn’s Disease Care Pathway

Project Sonar: Validating Predictive Algorithms and Risk Scores for IBD Outcomes Using a Community-Based Registry and Patient Engagement Data

Background

Project Sonar is a community-based registry and disease management program developed to improve clinical and economic outcomes in Inflammatory Bowel Disease (IBD). This cloud-based program integrates monthly patient-reported symptoms and health-related quality of life (HRQoL) information with clinical data delivered through electronic medical record derived Clinical Decision Support tools (CDS). These fields are then combined with payer provided claims data to provide comprehensive, real-time information to physicians and patients on current symptoms and health status, as well as composite ‘Sonar’ scores. The CDS tools were developed using the American Gastroenterological Association’s Crohn’s Disease Care Pathway (CDCP), and Ulcerative Colitis Care Pathway, both of which determine treatment guidelines for Crohn’s Disease and Ulcerative colitis (UC), respectively. The objective of this project was to verify and measure the variation of data within Project Sonar, validate the CDCP and UCCP guidelines, and Sonar risk scores using the real world data collected via Project Sonar and healthcare claims of a subset of patients. The objective of this presentation is to detail the methods used in this process.

Methods

The verification and validation project consisted of three major steps: (1) verification and comprehensive quality check of all Project Sonar data to assure its quality mirrors that of commercially available patient databases; (2) creation of an analytic dataset that merges disparate patient level data from electronic medical records, healthcare claims data, and patient reported outcomes; and (3) validation of CD and UC algorithms and Sonar risk scores. Assignment of patients into correct risk category was confirmed by verifying the elements of the three sub-scores (i.e., burden categories) that were important to the treatment algorithm and development of the overall risk score, and assuring that each element and sub-score were weighted appropriately. Then, the drivers for risk categorization were identified (i.e. the variables most commonly associated with patients moving to a higher risk category). Finally, convergent validity was assessed for the three sub-scores, risk categories, and Sonar scores by assessing the correlations between risk categories and other indicators of disease severity (e.g., healthcare resource use, health-related quality of life).

Results/Conclusions

Project Sonar provides a unique opportunity to combine electronic medical records, healthcare claims/resource utilization data, and patient reported outcomes to predict treatment failure and target appropriate therapy in a community-based setting. With the completion of this
comprehensive verification and validation process, the Project Sonar database provides a rich source of IBD patient data for understanding the natural course of IBD, identifying practice patterns, comparing treatment options and outcomes, and identifying potential therapeutic cost savings.

II. Project Sonar: Improvement in Patient Engagement Rates Using a Mobile Application Platform

Project Sonar: Improvement in Patient Engagement Rates Using a Mobile Application Platform

Background

Project Sonar (PS) is a community-based registry and disease management program developed by the Illinois Gastroenterology Group (IGG) to improve clinical and economic outcomes in patients with Inflammatory Bowel Disease (IBD). PS integrates monthly patient-reported health-related quality of life (HRQoL) information using a subset of questions from the Crohn’s Disease Activity Index (CDAI) sent via a Patient Portal (PP) producing a monthly ‘Sonar Score’. These scores are then joined by clinical data fields delivered through electronic medical record derived Clinical Decision Support tools (CDS). This combined data is analyzed against payer provided-claims data to provide comprehensive, real-time information to physicians and patients on current symptoms and health status, as well as composite ‘Sonar’ scores. In an effort to improve portal-based patient response rates PS deployed the SonarMD Platform (SMDP), a cloud based platform developed by SonarMD, LLC which uses smartphone technology to improve patient engagement instead of a PP.

Methods

Patient surveys were developed using five questions derived from the Crohn’s Disease Activity Index in an effort to obtain HRQoL scores. The sum of the values on these questions results in a Sonar Score (SS) which produces a quantifiable assessment of HRQoL. SSs are monitored for individual scores as well as the slope of change over time, which allows for the development of care management algorithms that drive interventions. The PP was used to send these surveys from January 2014 through May 2015. Due to an unacceptable PP response rate, the SMDP was deployed in June 2015. Patient response rates were compared for each of the two methods.

Results

The patient response rate for the PP averaged 27.6% over the 17 months of its use. Due to the structure of the PP, SSs needed to be individually calculated upon their receipt resulting in significant staff expense and a delay in patient feedback. The patient response rate for the SMDP was 66.27% for the three months it has been deployed, far exceeding the rate of the PP. Since the SMDP platform automatically calculates the SS at the time the patient answers the survey, patients receive immediate algorithm derived responses. Staff time is significantly reduced as a result.
Conclusion

Project Sonar provides a unique opportunity to combine electronic medical records, healthcare claims/resource utilization data, and patient reported outcomes to predict treatment failure and target appropriate therapy in a community-based setting. Since patient participation is critical to the success of PS, the cloud-based SMDP's superior performance improved the patient response rate over the patient portal from 27.6% to 60%. It also resulted in significant savings in staff time. More time of deployment will be necessary to confirm that these findings will stand the test of time.

III. Project Sonar: Validating a Cost Normalization Methodology in a Community-based Registry

TITLE: Project Sonar: Validating a Cost Normalization Methodology in a Community-Based Registry

AUTHORS: Kosinski, Lawrence1, 2; Sorensen, Michael2; Brill, Joel2, 3; Landsman-Blumberg, Pamela4; Turpin, Robin5; Baum, Charles5

INSTITUTIONS (ALL):
1. Illinois Gastroenterology Group, Elgin, IL, United States.
2. SonarMD, LLC, Elgin, IL, United States.
3. Predictive Health, Paradise Valley, AZ, United States.
4. Xcenda, LLC, Palm Harbor, FL, United States.
5. Takeda Pharmaceuticals USA, Inc, Deerfield, IL, United States.

BACKGROUND

- Project Sonar, a community-based registry and disease management program developed to improve clinical and economic outcomes in Inflammatory Bowel Disease.
- Project Sonar uses a cloud-based platform combining health-related quality of life (HRQoL) information with clinical data delivered through electronic medical record derived Clinical Decision Support tools.
- These data fields are then combined with payer provided-claims data obtained from the Intensive Medical Home with Blue Cross Blue Shield of Illinois (BCBS-IL) database.

OBJECTIVES

- Develop and validate a normalization methodology to mitigate the variation of cost data between actual payments made vs a normalized payment structure derived from Medicare Payment information to allow for meaningful assessment of Project Sonar’s economic impact.

METHODS

Patient Sample
- Patients participating in Project Sonar and continuously enrolled in BCBS-IL from January 1, 2014 through September 30, 2015.

Costing Method
Crohn’s related claims-based payments were identified for each patient in the BCBS-IL medical claims with ICD-9-CM 555.X or select ICD-9-CM symptom codes.

Claims-based payments were calculated and presented as total Crohn’s-related payments and by type of service (i.e., infusible injection, diagnostic services, inpatient services, and physician services).

All claims-based payments were adjusted to 2015 US dollars using the medical component of the Consumer Price Index.

Normalization Method

Normalized Crohn’s-related payments were calculated using standardized provider payments as displayed in Table 1.

Table 1. Factors to Normalize

<table>
<thead>
<tr>
<th>Service Type</th>
<th>Factors to Normalize</th>
</tr>
</thead>
<tbody>
<tr>
<td>Physician Services</td>
<td>CPT Code Payment on 2015 Outpatient Prospective Payment System</td>
</tr>
<tr>
<td>Diagnostic Services</td>
<td>2015 Clinical Diagnostic Laboratory Fee Schedule</td>
</tr>
<tr>
<td>Infusible Biologics</td>
<td>2015 office based payment rates</td>
</tr>
<tr>
<td>Inpatient Hospital Payments</td>
<td>DRG Payment</td>
</tr>
</tbody>
</table>

To remove bias from site of service all infusible biologics were normalized to office based infusion rate of $85/unit compared to $158/unit.

For inpatient Hospital Payments:
- The DRG Payment was calculated using a base rate derived as the sum of the Operating Base Payment and the Capital Base Payment.
- The base rate was then multiplied by the DRG Weight derived from the Medicare Severity Diagnosis-Related Groups Relative Weighting Factors.

Analysis

We examined the absolute difference and percentage difference between actual and standardized payments for total Crohn’s-related payments and by type of service.

RESULTS

Sample Description

185 patients participating in Project Sonar and continuously enrolled in BCBS-IL were included in the analysis.

Figure 1. Crohn’s-Related Payments, Total and by Type of Service
- Normalized Crohn’s–related payments are consistently lower than claims-based payments (see Figure 1), although the percentage difference varies by service type
  - Total: -18%
  - Physician services: -30%
  - Inpatient services: -47%
  - Diagnostic services: -56%
  - Infusible biologics: -46%
- Diagnostic
- Inpatient
- Physician
- X% of infusions were administered in the hospital outpatient department which may contribute to the large difference in normalized payments compared to claims based payments

LIMITATIONS
- ICD-9-CM codes for Crohn’s related symptoms were selected based on clinical opinion after review of a sample of medical claims for BCBS-IL Crohn’s patients enrolled in Project Sonar and the published literature. Therefore, costs could be underestimated if relevant ICD-9-CM codes were omitted and overestimated if symptom claims included were related to alternative diagnoses.
- Claims data are collected for the purpose of billing, not research. Therefore, diagnoses may be coded incorrectly or under-coded, possibly introducing measurement error.

CONCLUSIONS
- Project Sonar represents a unique opportunity to manage care using the AGA Crohn’s Care Pathway and analyze the results on payments.
- For meaningful comparison, it is essential that payments are normalized against Medicare Payments in order to mitigate the site of service differential in payment reporting.
- The normalization methodology presented has broad applicability for those seeking to analyze complex real-world clinical practice data and cost.
IV. Project Sonar Abstract of Distinction: Digestive Disease Week 2016

Project Sonar: Reduction in Cost of Care in an attributed cohort of patients with Crohn’s Disease.

Lawrence Kosinski,1 Michael Sorensen,1 Joel V. Brill,2 Pamela B Landsman-Blumberg,3 Robin Turpin,4 Charles Baum4


Background: Project Sonar (PS), a joint venture between Illinois Gastroenterology Group and Blue Cross Blue Shield Illinois, is a community-based registry and disease management program developed to improve clinical and economic outcomes in Inflammatory Bowel Disease (IBD). PS integrates patient-reported symptoms and health-related quality of life information with clinical data delivered through electronic medical record derived Clinical Decision Support tools (CDS) using a cloud-based program developed by SonarMD, LLC. CDS tools are based on the American Gastroenterological Association’s Crohn’s Disease Care Pathway (CDCP) to determine treatment guidelines for Crohn’s Disease (CD). Clinical data is combined with payer claims data to provide comprehensive, real-time information to physicians and patients on current symptoms and health status, as well as composite ‘Sonar’ scores. This study reports performance of Project Sonar in a subset of continuously enrolled CD patients.

Methods: 152 attributed patients with CD were continuously enrolled in PS between Jan 1, 2014 and Sept 30, 2015. The baseline period was Jan 1 through Nov 30, 2014, and the study period was the PS go-live of Dec 1, 2014 through Sept 30, 2015. Total Costs were corrected for the difference in time periods. CD-based claims were identified using ICD-9 555 codes plus additional codes identified as Crohn’s-based due to their relationship with CD. Payments for physician and diagnostic services were normalized using a methodology based on Medicare payment rates; DRG payments for inpatient stays were adjusted to eliminate site of service differentials in payment rates.

Results

<table>
<thead>
<tr>
<th>Crohn’s Payments</th>
<th>Total Crohn’s Payments</th>
<th>Average Per Patient Payment</th>
<th>Total Normalized Crohn’s Payments</th>
<th>Normalization Difference</th>
<th>Inpatient Payments</th>
<th>Emergency Room Payments</th>
<th>Infusible Biologics</th>
<th>Injectable Biologics</th>
<th>Total Biologics</th>
</tr>
</thead>
<tbody>
<tr>
<td>Pre-Period</td>
<td>2,118,308.65</td>
<td>$13,936.24</td>
<td>1,932,069.37</td>
<td>$196,339.28</td>
<td>210,967.47</td>
<td>$52,363.19</td>
<td>892,443.30</td>
<td>560,980.84</td>
<td>1,453,424.14</td>
</tr>
<tr>
<td>Study Period</td>
<td>1,884,758.63</td>
<td>$12,399.73</td>
<td>1,741,326.02</td>
<td>$143,432.62</td>
<td>90,410.85</td>
<td>$24,465.00</td>
<td>972,485.87</td>
<td>423,561.72</td>
<td>1,396,047.59</td>
</tr>
<tr>
<td>Difference</td>
<td>$233,550.02</td>
<td>$1,536.51</td>
<td>$190,743.36</td>
<td>$42,806.66</td>
<td>$20,556.62</td>
<td>$27,898.19</td>
<td>80,042.57</td>
<td>137,419.12</td>
<td>$57,376.55</td>
</tr>
<tr>
<td>Percentage Difference</td>
<td>-11.03%</td>
<td>-11.03%</td>
<td>-9.87%</td>
<td>-22.98%</td>
<td>-57.14%</td>
<td>-53.28%</td>
<td>8.97%</td>
<td>-24.50%</td>
<td>-8.95%</td>
</tr>
</tbody>
</table>

Findings: Total payments for PS patients with CD declined by 11.03% driven by > 50% declines in hospital and emergency room payments. Utilization of physician administered biologics rose 8.97% but total biologic costs declined 3.95% driven by a 24.5% reduction in the use of injectable biologics. Normalization of payments to eliminate site of service payment differentials resulted in a 9.87% savings in payments.

Conclusions: PS demonstrates value-based improvement in care for CD patients in a community-based setting through integration of clinical data, patient reported outcomes and healthcare claims/resource utilization data. Over 50% declines in hospital admissions and emergency room visits were achieved through CDS tools promoting adherence to AGA CDCP combined with intense patient engagement. PS represents an innovative model of population health likely applicable to other chronic conditions.
V. Guide to the Use of the Crohn’s Disease CDS Tool

The use of CDS tools is critical to our success in the management of our patients with chronic disease. They also provide us an opportunity to bring together our PQRS requirements and serve as a platform for participation in research studies. Accordingly, the Project Sonar Physician Advisory Board, with the approval of the IGG Board of Managers, has decided to make the use of these CDS tools mandatory. This document will provide you clarity in how to enter data into this template.

The Crohn’s Disease CDS Tool is a “Demographic” template and therefore only has to be populated only once. It will remain in its populated state from visit to visit and only requires updates. It will be launched automatically when you enter a chief complaint of Crohn’s Disease.

The template has several sections as shown below:
VI. AGA Crohn’s Disease Care Pathway Risk Assessment
25 metrics are captured in three categories: Disease Burden, Inflammation Burden, and Comorbidity Burden. This assessment is updated annually on each patient. SonarMD is developing a mathematical model behind the relative strengths of each metric. Currently due to the size of our sample population, we are not able to reach statistical significance. The implementation of this APM on a national basis would provide the necessary patient population.
VII. SonarMD Nurse Care Manager Dashboard

The demo site shown below with fictitious patients demonstrates the user interface that the NCMs and Physicians use each month to monitor their patients. Patients are listed according to the assigned list of each NCM and Physician. Their raw Sonar Scores as well as the slope of their scores over time are listed with color coordination based upon algorithmic rules. The NCM or Physician can drill down on each one for further details. See VIII below.
VIII. SonarMD Patient Survey
This template allows the reviewer to drill down on individual surveys. Each patient receives the following questions every month via their device of choice. This template is an example of what the NCMs utilize to monitor the answers and scores.

<table>
<thead>
<tr>
<th>Question</th>
<th>Date Sent: 11/1/2016</th>
<th>Date Received: 11/1/2016</th>
<th>Date Sent: 09/28/2016</th>
<th>Date Received: 09/28/2016</th>
<th>Date Sent: 09/01/2016</th>
<th>Date Received: 09/07/2016</th>
</tr>
</thead>
<tbody>
<tr>
<td>How many loose stools per day have you had in the last seven days?</td>
<td>One (1)</td>
<td>One (1)</td>
<td>Two (2)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>For the last seven days how would you describe any abdominal pain or cramps?</td>
<td>Mild</td>
<td>Mild</td>
<td>Mild</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Over the last seven days how would you describe your general well being?</td>
<td>Generally well</td>
<td>Poor</td>
<td>Slightly Under</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Select any of the symptoms you have had over the last seven days.</td>
<td>Eye Pain</td>
<td>Eye Pain</td>
<td>Eye Pain</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Are you currently taking any medication for diarrhea?</td>
<td>Yes</td>
<td>No</td>
<td>Yes</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Do you have any questions or comments for your care team?</td>
<td>Completed by M.Sorensen</td>
<td>Mike testing</td>
<td>Survey taken by M.Sorensen</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Are you currently taking one of the following medications?</td>
<td>Entyvio</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Are you currently taking one of the following medications?</td>
<td>None of these</td>
<td>None of these</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>
IX. Examples of SonarMD Ping Reports

The report corresponds to the answers for the first question in the survey shown in the vertical axis. For the last seven days how would you describe any abdominal pain or cramps? The figure can be filtered by the items shown on the right side: age, gender, depression rating and by responder percentage.

The following figure demonstrates the same findings for the second question: “How many loose stools per day have you had in the last seven days”
X. SonarMD Cost Report
The following figure shows a steady fall in average aggregate cost of care over the time period of June 1st 2015 through August 31st 2016.