



Surgical Outcomes System

About Us

The Surgical Outcomes System is an orthopedic and sports medicine global registry, that is cloud-based and tablet-friendly. The intuitive platform enables surgeons to easily collect and analyze patient outcomes and procedures to address the demands of evidence-based medicine. The SOS™ global registry allows surgeons to remotely monitor their patients' outcomes over two years through patient-reported outcome questionnaires and clinician-reported objective measures. The SOS will automatically email surveys to patients at predefined time points. The primary purpose of the SOS is for monitoring and evaluating the outcomes of various standard of care orthopedic and sports medicine medical procedures, leading to the development of evidence-based protocols for treating patients. Secondary purposes of SOS include analysis and publication of results, patient education and cost-effectiveness of treatment procedures and products.

Features and Benefits

- ▶ Cloud-based, HIPAA reviewed and IRB approved registry that allows for data collection, remote monitoring of patient data and comparative analysis of treatment outcome data.
- ▶ The SOS requires only a few steps for the surgeon to enroll a patient into a study.
- ▶ The SOS leverages technology, engaging patients in their own healthcare through the automated surveys, video links, images, reminders and alerts.
- ▶ Surgeons can use the SOS de-identified global average for marketing, providing evidence of successful surgical/treatment outcomes to their peers and future patients.
- ▶ The SOS is fully transportable. Both the patient portal and surgeon portal are accessible on PCs, tablets or smart devices.
- ▶ The SOS allows for real-time clinical review and decision making of global effectiveness of various orthopedic and sports medicine clinical procedures, leading to the development of evidence-based protocols for treating patients.
- ▶ Surgeons are able to create, save and edit "favorites." Favorites can be used to auto populate post-treatment details on surgeries that are performed routinely as a time-saving measure.
- ▶ The SOS is a "global registry" with the patient portal available in multiple languages.

Patient Portal

The patient portal is an intuitive, easy-to-use, web-based application for enrolled patients to submit health outcome data over two years posttreatment using validated patient-reported questionnaires. The patient portal is fully transportable and available on any smart device or tablet.

The format of the patient portal is either multiple choice or scale-rating type, which makes the use of the interface easy and intuitive.

The patient can change his/her selection throughout the submission process. Questions cannot be skipped.

The system indicates survey progress.

Universal Pain Assessment

1 Please choose a number between zero and ten which most closely approximates the pain you are experiencing today

0 1 **2.01** 3 4 5 6 7 8 9 10

No Pain Mild Pain Moderate Pain Severe Pain Worst Pain Possible

Please answer the following question(s)

2 Do you take narcotic pain medication (codeine or stronger)?

Yes
No

3 Do you take pain medication (aspirin, Advil, Tylenol, etc)?

Yes
No

4 Are you taking any sleeping medication due to the discomfort of your recent treatment?

Yes
No

Symptoms

These questions should be answered thinking of your knee symptoms during the last week.

2 Do you have swelling in your knee?

✓ Sometimes

3 Do you feel grinding, hear clicking or any other type of noise when your knee moves?

Never
Rarely
Sometimes
Often
Always

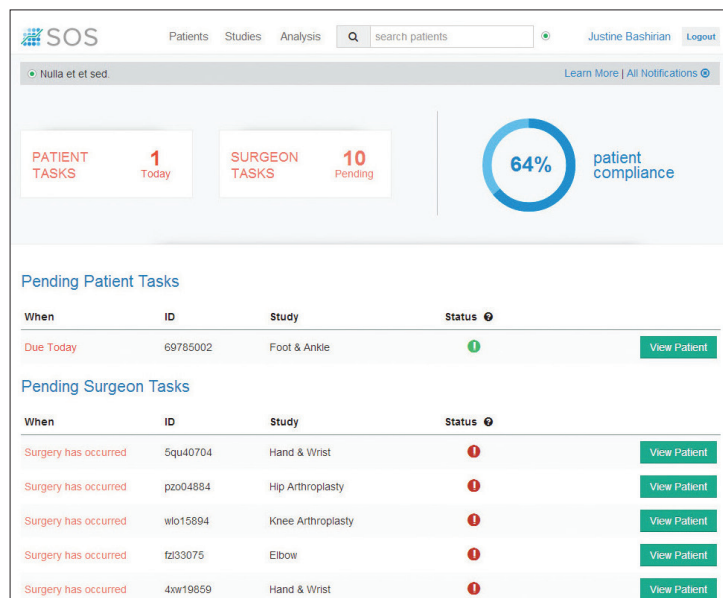
4 Does your knee catch or hang up when moving?

Never
Rarely

User Dashboard

The surgeon portal is divided into three panels: Patients, Studies, and Analysis. Intuitive navigation and interactive features make the surgeon portal easy to use, while providing all necessary tools to monitor the patient's health progress over time and analyze the outcome data.

- ▶ All features and data are fully transportable and accessible from the SOS™ surgeon portal through a PC, smart device or tablet
- ▶ Patient compliance is highlighted upon login, providing accountability to achieve the goal of at least 80 percent
- ▶ Pending patient tasks and surgeon tasks are organized based on time deadline, keeping sites organized for follow-up
- ▶ Sites are able to view patient details through a direct click from the dashboard
- ▶ Patient search capability is available using custom ID, study or patient email address



Patients

The Current Patients panel displays all enrolled patients by custom ID, email address and corresponding enrolled studies. Patients can be searched and organized by custom ID, email address or study.

Colored icons provide a visual reference as to which case is up-to-date and which case has missing data.

Enrolling a new patient is simple and fast, requiring only 15 seconds to enter demographic information.

SOS

PatientsStudiesAnalysis

search patients

Justine BashirianLogout

Nulla et et sed.

Learn More | All Notifications

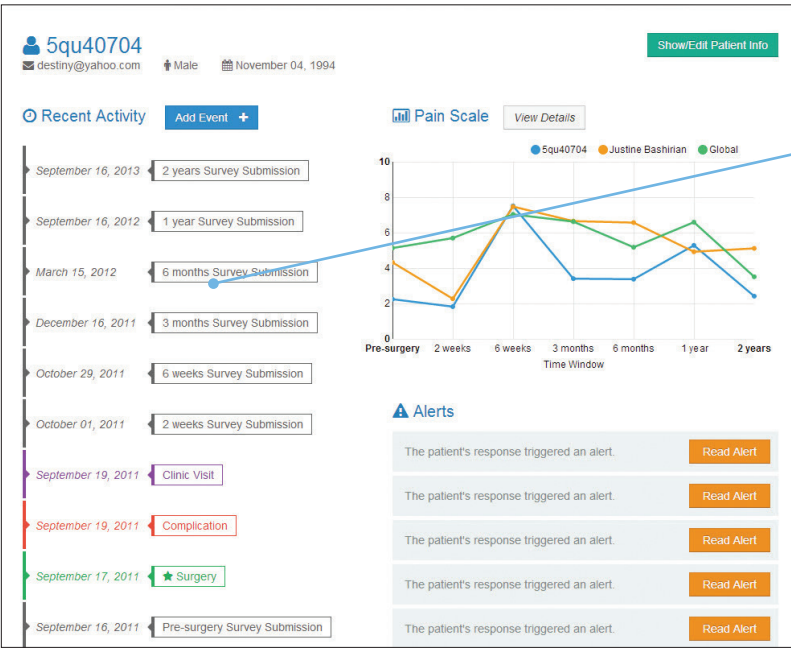
Home > Patients

Patients

Search

ID	Email	Studies	PS	SS	
5qu40704	destiny@yahoo.com	Hand & Wrist			View Patient
pzo04884	hip_arthroplasty@arthrex.com	Hip Arthroplasty			View Patient
ljr24318	hip_arthroscopy@arthrex.com	Hip Arthroscopy			View Patient
wlo15894	knee_arthroplasty@arthrex.com	Knee Arthroplasty			View Patient
91612151	knee_arthroscopy@arthrex.com	Knee Arthroscopy			View Patient
afn27634	shoulder_arthroplasty@arthrex.com	Shoulder Arthro...			View Patient
wl975143	shoulder_arthroscopy@arthrex.com	Shoulder Arthro...			View Patient
fz33075	elbow@arthrex.com	Elbow			View Patient

Enroll Patient +



By clicking on the “View patient” button, sites are able to view the patient’s record in a timeline fashion, highlighting events such as survey submissions, surgeries, clinic visits or complications.

Sites can easily add an event, edit patient information, and view patient outcomes compared to the site outcomes and the global de-identified average within the enrolled study.

Patients (Cont.)

Sites are able to enter treatment detail in 30 seconds or less with the intuitive dropdown menus.

Surgeons are able to create, save and edit "favorite" templates, allowing for autopopulation of treatment detail.

Filter Menu

Patient baseline and case details

Diagnosis

Operations

Surgical time and personnel

Rotator Cuff (Complete Tear)

Rotator Cuff (Partial Tear)

Subacromial Decompression

Tendon Transfer

Biceps

Coracoidplasty

Distal Clavicle Excision

AC Joint Stabilization

Os Acromiale

Osteotomy

Cartilage treatment

Osteochondral Treatment

Other intra-articular

Fracture Management (ORIF)

Miscellaneous

Operations - Labrum & Instability

Other type grafts and bone graft details

Implants

Injections/Applications

Favorites

Speedbridge

Add Favorite +

Edit Surgery on November 04, 2013

Update Event

Rotator Cuff (Complete Tear)

Procedures: Double row repair

Anchor repair technique used: SpeedBridge - Medial knots not tied

Primary or Revision

Select

Open

Yes

No

Procedures

Double row repair x

Anchor repair technique used

SpeedBridge - Medial knots not tied x

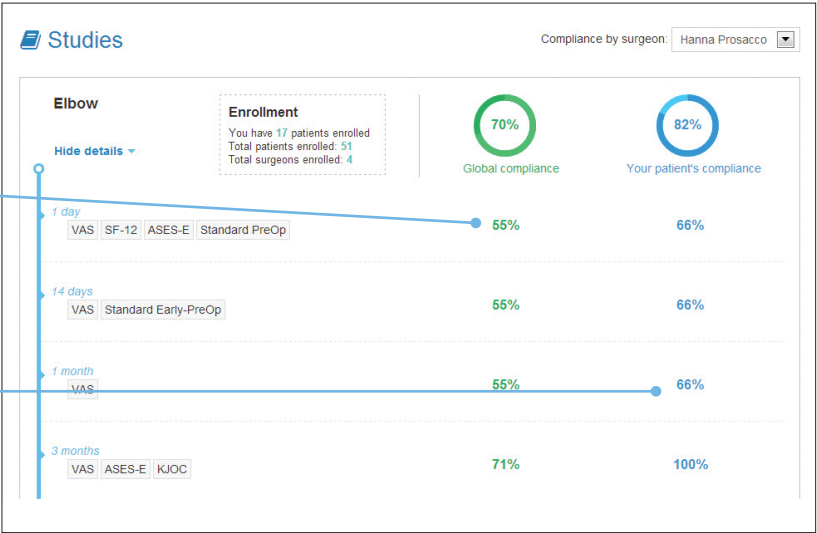
Remove

Studies

Surgeons are able to view a summary of enrolled patients per study, as well as patient compliance per time point.

Number of patients enrolled and patient compliance of surveys are benchmarked with the global information.

Displaying patient compliance per time point allows sites to pinpoint areas of opportunity.

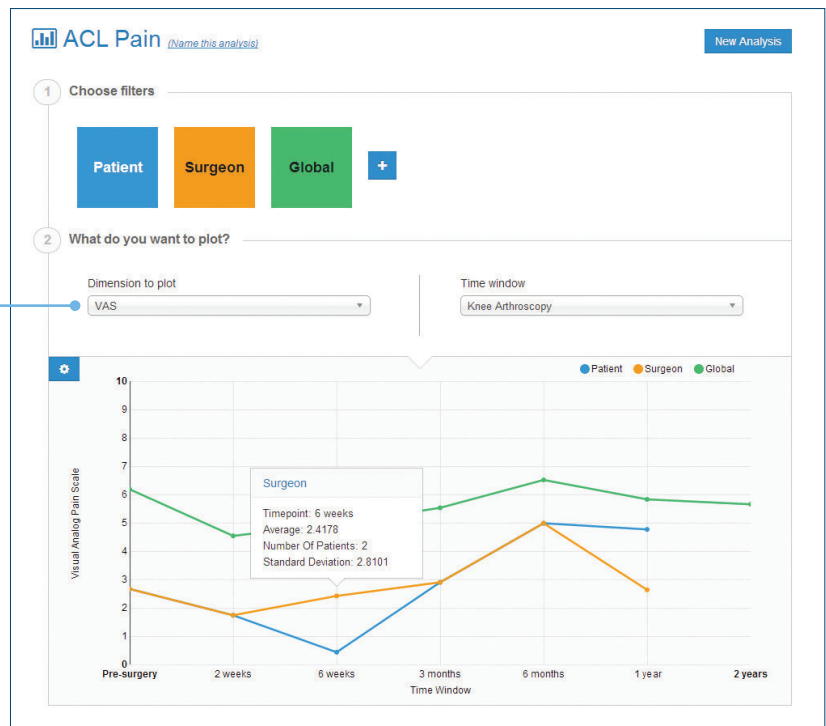


Analysis

The Patient Analysis panel provides a graphical representation of the outcomes data. Sites are able to perform comparative analysis on individual outcomes, site outcomes or the global de-identified average by choosing filters. Data can be exported in raw data format or graphical format, and emailed as an image to patients for educational purposes.

Multiple patient events can be filtered within the same study or across studies, such as surgery details, complications, clinic visits, or survey submissions. Unlimited analyses can be performed.

Sites are able to analyze “groupings,” including or excluding criteria such as patients, study group, accounts, countries or demographics.



Uni Knee Outcomes [\(edit\)](#)

Groupings +

Groupings

- Account
- Country
- Ethnicity
- Event
- Patient
- Race
- Study
- Time Zone
- User

Uni Knee Outcomes [\(edit\)](#) Plot Line Color Cancel Save Filter

Groupings **Knee Arthroplasty - Su** +

Filter Menu

Select catalog items from the menu on the left to populate the event for this patient.

Patient baseline and case details
Diagnosis
Operations/Implants
Injections/Applications

[Hide Advanced Options](#)

Event is included

Filter By Age Range

Filter By Date Range

Select Age Range

Min Max

Ok

Sites are able to filter by age and date of surgery.

Surgeons interested in utilizing the Surgical Outcomes System™ should contact SOS@arthrex.com, or their local Arthrex sales representative for more information.

Workflow

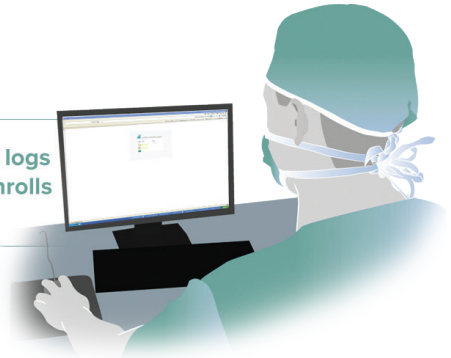


BEGIN

Patient agrees to participate

1.

Surgeon or designated staff **logs on** to outcome portal and **enrolls patient** into the study



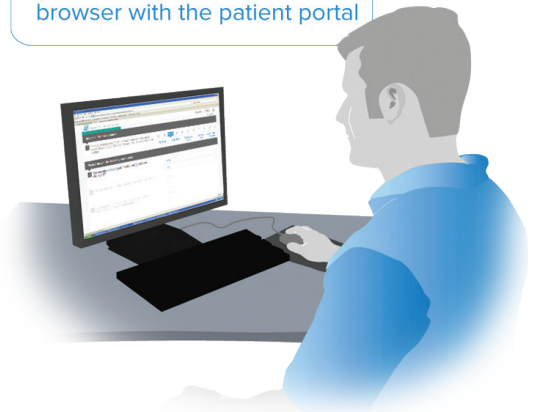
2.

System **sends e-mail** notifications to patient at predefined time milestones



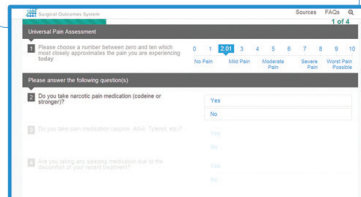
3.

Patient clicks on a **link in the e-mail** which launches a web browser with the patient portal



4.

Patient **answers questions online**



5.

Surgeon or staff enters surgery related **post-op information** immediately following surgery

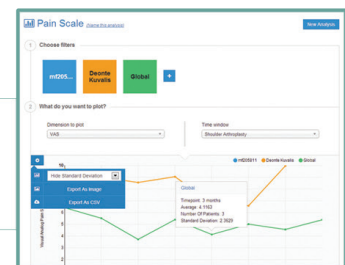


6.

Surgeon **can review** patient's subjective and objective outcomes

7.

Surgeon has the option to **compare patient outcomes** to the global average





Surgical Outcomes System

Advancing Evidence-Based Medicine

This description of technique is provided as an educational tool and clinical aid to assist properly licensed medical professionals in the usage of specific Arthrex products. As part of this professional usage, the medical professional must use their professional judgment in making any final determinations in product usage and technique.

In doing so, the medical professional should rely on their own training and experience and should conduct a thorough review of pertinent medical literature and the product's Directions For Use.