



PSDR UNPACKED: WHAT IT IS & WHY IT MATTERS

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Patients First. *Always.*

UNPACKING PSDR: MEET THE SPEAKERS



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LEARNING OBJECTIVES:

- Define Patient Safety Data Reporting (PSDR)
- Explain the specific reporting nuances associated with PSDR
- Explain the importance of the data obtained from PSDR





WHAT IS PSDR?

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PATIENT SAFETY DATA REPORTING (PSDR)

- QUAD A's proprietary data collection system that enhances quality control measures by quantifying safety data as reported by our accredited facilities
- This mandatory data reporting process fosters clinical excellence by demonstrating the benefits of accreditation and revealing trends that impact patient outcomes
- With data from hundreds of thousands of cases, it forms one of the nation's largest healthcare data repositories

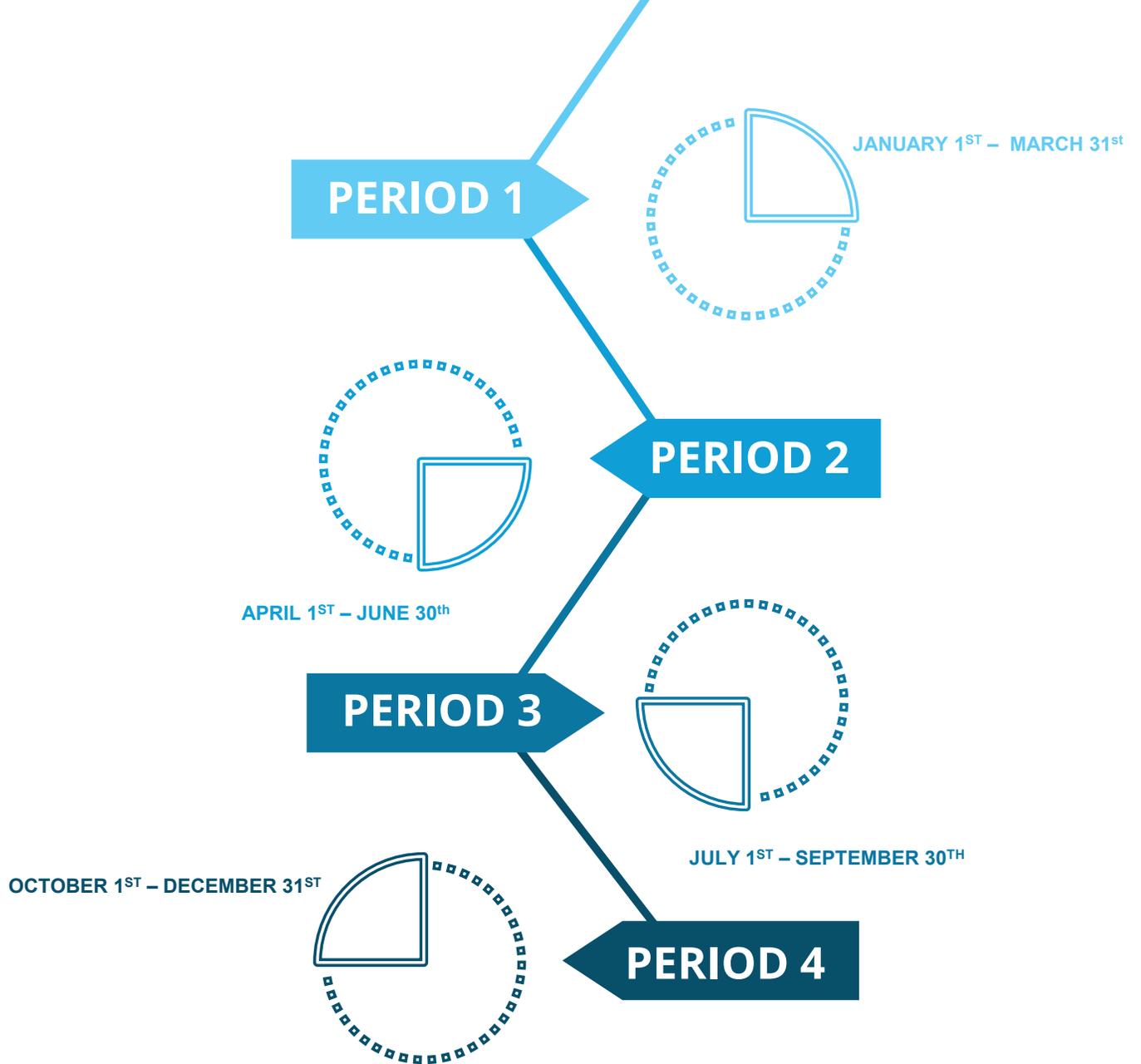




REPORTING SPECIFICS

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PSDR REPORTING PERIODS



REPORTING PERIOD INFORMATION

PERIOD 1:

January 1st – March 31st



Deadline to Report: April 15th

PERIOD 2:

April 1st – June 30th



Deadline to Report: July 15th

PERIOD 3:

July 1st – September 30th



Deadline to Report: October 15th

PERIOD 4:

October 1st – December 31st



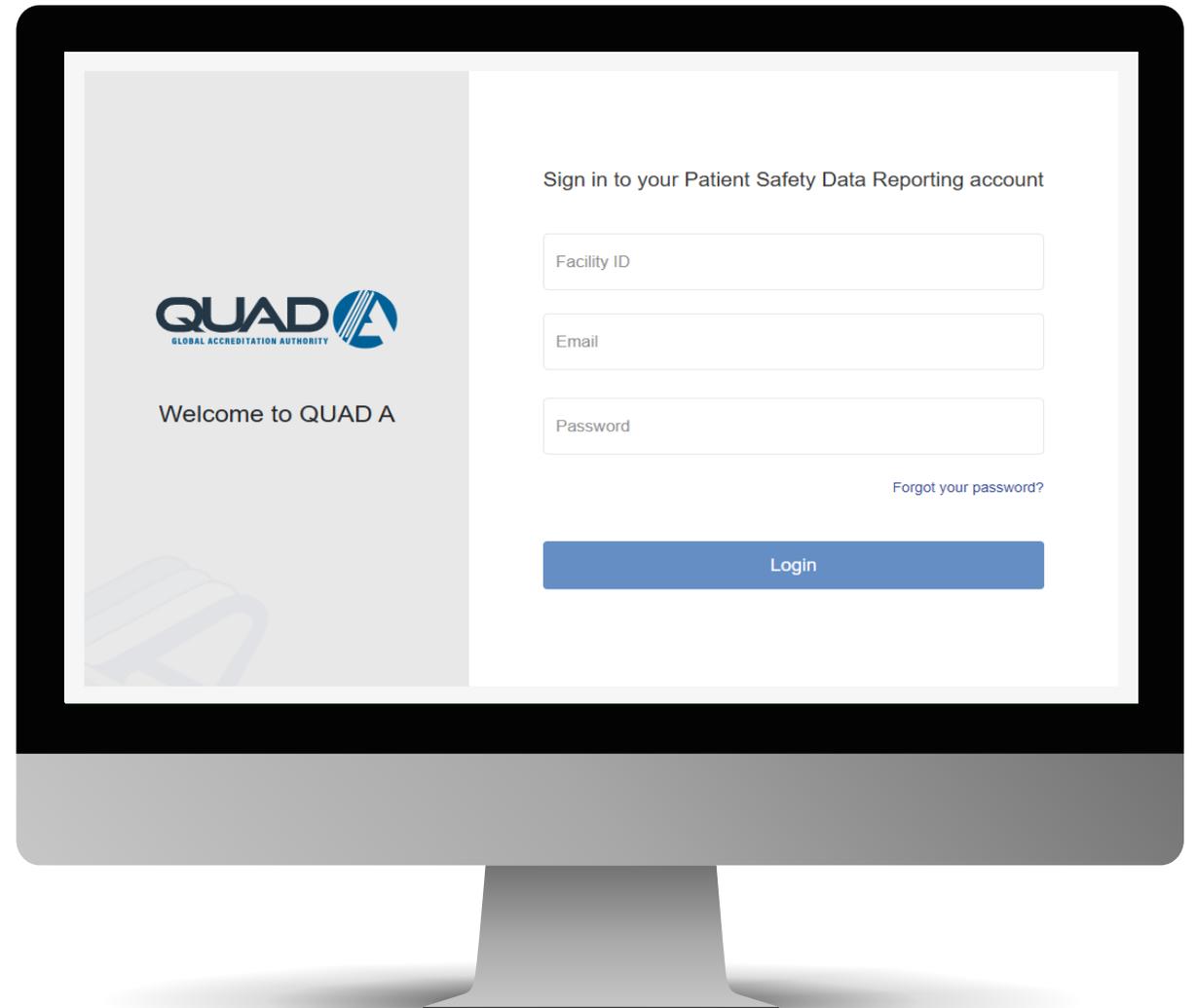
Deadline to Report: January 15th

- Four fixed reporting periods (quarters) per calendar year
- Fifteen-day grace period following the conclusion of each period to submit mandatory data
- Non-compliance results in probation and additional fees

SUBMITTING YOUR CASES

- **Case submissions must be submitted by designated PSDR Administrator**
 - PSDR Administrator Authorization Form must be filled out and submitted by the Medical Director
- **All cases must still be entered directly into the QUAD A online portal**
 - QUAD A offices do not accept paper forms
 - Must fill in all required fields to submit PSDR random cases
- **Failure to comply with reporting deadlines results in:**
 - 60-day probation period
 - \$100 late fee per non-compliant provider

PSDR.QUADA.ORG



WHICH CASES NEED TO BE REPORTED?

GUIDELINES

- **Each surgeon/proceduralist must submit 3 cases per period**
 - The 1st case performed by surgeon/proceduralist in each month of the reporting period are the cases submitted for PSDR
- **A total of 12 cases must be submitted annually**
 - These cases should correspond to the 1st case performed each month throughout the year
- **Any cases involving unanticipated sequelae should be reported**



RESOURCES AVAILABLE TO YOU: PSDR REPORTING TEMPLATES

The image displays three overlapping forms from QUAD A (Global Accreditation Authority). The top-left form is the 'Unanticipated Sequela Template', the top-right is the 'Unanticipated Sequela Addendum Template', and the largest, central form is the 'Patient Safety Data Reporting Template'. Each form includes the QUAD A logo and contact information: 600 Central Ave. Ste 265 | Highland Park, IL 60035; (direct) 847.775.1970 | (fax) 847.775.1985; info@QuadA.org. The 'Patient Safety Data Reporting Template' includes sections for Facility Information, Patient Information, Surgical/Procedural Information, Anesthesia Information, and Medical Record Review. The 'Medical Record Review' section contains a checklist of items such as Pathology Report, Pre-Op Plan for Treatment, Informed Consent, Medical History, Physical Examination, Laboratory Reports, Post-Op Recovery Record, Anesthesia Record, RX Given to Patient, Discharge Instructions, Operative Report, and Recorded in Log, each with 'Yes', 'No', and 'N/A' options.

- **Templates available on QUAD A website**
 - Unanticipated sequela = adverse advent
 - Verbiage used interchangeably
 - These resources are available to help your facility understand the information that will be captured when submitting in the PSDR portal
- **Use these templates as internal tool at your facility**
 - Assists in record-keeping
 - Outlines all information that you will need to fill in for each submission within the portal

PSDR REPORTING EXAMPLES

Random Review

Reporting Period: Period 1 (January 1, 2025 - March 31, 2025) Physician: Scott A. Bianchi Surgery Facility: AAAASF Test Facility - Facility Name Status: 1/3 Open

Patient Information

Patient Initials: EP Patient Age: 28 Height type: ft Height (in feet): Gender: Ethnicity: Weight Type: lbs

Weight (in pounds):

Procedure Information

Surgery Date: Duration in Hours: Minutes:

Procedure Name:

Anesthesia Information

Type: Provider: Duration in Hours: Minutes:

BAD REPORTING

PSDR REPORTING EXAMPLES



Random Review

Facility ID: 4474

Reporting Period: Period 1 (January 1, 2025 - March 31, 2025)

Physician: Scott Bianchi

Surgery Facility: AAAASF Test Facility - Facility Name

Submission Status: Complete

Patient Information

Patient Initials: RA

Patient Age: 1

Height: 9 m

Height: 99 cm

Gender: Female

Weight: 3.6 kg

Ethnicity: Asian

Procedure Information

Surgery Date: January 1, 2025

Duration in Hours: 0

Minutes: 0

Procedure Name	Notes
1-stage repair of distal hypospadias and chordee with urethroplasty using flip-flap	test

Anesthesia Information

Type	Provider	Duration in Hours Minu	
Local Anesthesia	Registered Nurse	0	0

Patient Care Review

Pathology report adequate Yes No N/A

Pre-op plan for treatment Yes No N/A

Plan informed consent Yes No N/A

Medical history Yes No N/A

Physical examination Yes No N/A

Laboratory reports Yes No N/A

Post-op recovery record Yes No N/A

Anesthesia record Yes No N/A

Rx given to patient Yes No N/A

Discharge instructions Yes No N/A

Operative report Yes No N/A

Recorded in Log Book Yes No N/A

GOOD REPORTING

PSDR REPORTING: REPORTING A DEATH



- Any death occurring in an accredited facility or any death occurring within thirty (30) days of a procedure performed in an accredited facility must be reported within five (5) business days after the facility is notified or otherwise becomes aware of that death.
- Report as an Adverse Event in the online Patient Safety Data Reporting portal.
- QUAD A office will not accept paper forms or email notifications as an official way to report a death of a patient.

EXCEPTIONS TO THE RULES

1. ZERO CASES PER MONTH

- If a surgeon/proceduralist has not performed at least one case per month making up the reporting period in question, they can select a case from other months within the period to report

2. LESS THAN 3 CASES PER REPORTING PERIOD

- Must fill out Exemption Form on Quad A's website or PSDR portal.
- All surgeries and/or procedures performed by surgeon/proceduralist with less than 3 cases should be reported online in the PSDR portal for that period

PSDR REPORTING: EXEMPTION FORM

- Send QUAD A team an Exemption Form before the quarterly deadline to helpdesk@quada.org
- All PSDR forms are found on the main website www.quada.org or in PSDR Portal
 - Website: Homepage > Resources & Education > PSDR > PSDR Documents
 - PSDR Portal: Login Page > Dashboard > Forms (on lefthand side).
- This includes physicians that have been removed from 'Active Status' during this reporting period.



QUAD A
GLOBAL ACCREDITATION AUTHORITY
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600 Central Ave. Ste 265 | Highland Park, IL 60035
(direct) 847.775.1970 | (fax) 847.775.1985
helpdesk@QuadA.org

Patient Safety Data Reporting Exemption Form

This form should be used when surgeon/proceduralist has performed fewer than three (3) cases during the reporting period. Please make sure to submit all cases online (psdr.quada.org) prior to submitting this document.

Period I – Jan. 1 to March 31 Period II – April 1 to June 30 Period III – July 1 to Sept. 30 Period IV – Oct. 1 to Dec. 31

Period: Year:

Facility Name: Facility ID #:

Surgeon/Proceduralist Name	# Cases Completed	Reason for Exemption
<i>Example: A. Surgeon MD</i>	<i>0</i>	<i>No cases performed in period (due to COVID)</i>

(Note: Please use more than one form if needed.)

Medical Director Name:

Medical Director Signature: Date:

Please submit this form via fax or email (preferred methods) or mail prior to the Patient Safety Data Reporting deadline to remain in compliance.

Email: helpdesk@QuadA.org

Fax: (847) 775-1985

Mail: QUAD A
600 Central Ave, Suite 265
Highland Park, IL 60035



PSDR VS PEER REVIEW: WHAT'S THE DIFFERENCE?

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PSDR VS PEER REVIEW

PSDR

- Quarterly online data submission to Quad A
- Must include 3 random cases per quarter per surgeon
- Must include submission of all unanticipated sequela

PEER REVIEW

- **Facility-based meeting to improve quality and safety of care**
- Quarterly or Semi-Annual peer review meeting conducted based on case-load
 - Semi-Annual Peer Review Meeting: <50 cases per month
 - Quarterly Peer Review Meeting: >50 cases per month
- **At a minimum, the quarterly sample size for Peer Review is 10% of the average monthly case volume for the quarter, which includes the 1st case of each month for each surgeon/proceduralist.**
- Performed by a surgeon other than the operating surgeon, unless otherwise specified by state regulations



PSDR: WHY IS IT IMPORTANT?

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THE IMPORTANCE OF PSDR

Collects Statistics For Quality Assurance:

- Improves Patient Care
- Establishes quantifiable safety data
 - 12 cases per surgeon per year
- Drives standards revisions
- Contributes to scholarship and safety

All Adverse Events

- Adverse events 1/207 procedures .34%
- Hematoma .11%
- Infection .05%
- VTE .01%
- Death 1/55,032 procedures or .0018%
- VTE: 37% of deaths

PSDR IN THE LITERATURE

SAVING LIVES:

The Key to Preventing a Common Postoperative Complication



OTHER RESEARCH & OUTCOMES

Harvard researchers have maximized the available PSDR data:

- “National Mortality Rates after Outpatient Cosmetic Surgery and Low Rates of Perioperative Deep Vein Thrombosis Screening and Prophylaxis.” *Plastic and Reconstructive Surgery*, 142(1), 90-98.
- “Quantifying the Crisis: Opioid-Related Adverse Events in Outpatient Ambulatory Plastic Surgery” *Plastic and Reconstructive Surgery* 145(3):687-695
- “Complications from Fat Grafting and Gluteal Augmentation in Outpatient Plastic Surgery: An Analysis of American Association for Accreditation of Ambulatory Surgical Facilities”
- “Liposuction Complications in the Outpatient Setting: A National Analysis of 246,119 Cases in Accredited Ambulatory Surgery Facilities”
- “How Altra Medic Hospital in Guatemala Overcame”



YOUR QUESTIONS, OUR ANSWERS

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QUESTION:

How do you use the data submitted in PSDR portal? Can my facility's data be identified based on submission information?

QUESTION:

Since PSDR is technically “research” can we opt out of submitting on a quarterly basis?

QUESTION:

Can we request a copy of the data submitted for PSDR?

CONTACT US!

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Standards Questions?

Email Our Clinical Team!

standards@quada.org

THANK YOU!
