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Analysis of 1027 Adverse Events Reports for Interspinous Process Devices From the US Food and Drug Administration Manufacturer and User Facility Device Experience Database

GREGORY S. KAZARIAN, MD¹; YUSEF J. JORDAN, MD¹; MITCHELL JOHNSON, MD¹; SATYAJ BHARGAVA, BS¹; ROBERT CECERE, BS¹; TAKASHI HIRASE, MD¹; SHEERAZ QURESHI, MD¹; JAMES DOWDELL, MD¹; EVAN SHEHA, MD¹; FRANCIS LOVECCHIO, MD¹; AND SRAVISHT IYER, MD¹

¹Orthopedic Surgery, Hospital for Special Surgery, New York, NY, USA

ABSTRACT

Background: Interspinous process devices (IPDs) introduce a new class of complications to surgical decompression without fusion: hardware-related complications. The purpose of this study was to describe the adverse events associated with IPDs.

Study Design: This was a retrospective review of the Food and Drug Administration Manufacturer and User Facility Device Experience database.

Methods: The database was queried from its inception to November 2022 for reports associated with “Prosthesis, Spinous Process Spacer/Plate.” Entries were categorized by event type, patient impact, and interventions.

Results: A total of 943 surgery-related adverse events were identified. The most common intraoperative events were implant malfunctions (39.7%, $n = 374$) and fractures (2.2%, $n = 21$). The most common postoperative events were persistent pain (26.6%, $n = 251$), implant migration (19.1%, $n = 180$), and fracture (6.8%, $n = 64$). The most common resultant outcome of an adverse event was the need for revision surgery (48.8%, $n = 460$). The need for revision surgery was common in patients who experienced fracture (47.1%), implant migration (84.5%), infection (76.7%), and neurological complications (76.9%). Implant migration, fracture, and implant malfunction, 3 complications that are unique to decompression with an IPD as compared with traditional laminectomy, accounted for 45.9% of revisions (211/460), and revision was required in 33.0% of cases where 1 of these complications was reported (211/640). Implant malfunction made up 21.2% of Coflex complications, 47.3% of Superion complications, and 5.2% of X-Stop complications.

Conclusions: The most common adverse events were implant malfunction, inadequate efficacy, implant migration, and fracture. Concerningly, these complications require revision surgery in one-third of cases when they occur. Implant-specific assessments demonstrate a high prevalence of implant malfunctions for the Coflex and Superion implants.

Clinical Relevance: Interspinous process devices introduce a new class of complications to isolated spinal decompression surgery: implant-related complications. These complications occur both intraoperatively and postoperatively, and they frequently necessitate revision surgery.

Level of Evidence: 4.

Lumbar Spine

Keywords: interspinous process device, adverse events, MAUDE, implant malfunction, migration, fracture

INTRODUCTION

Degenerative lumbar spinal stenosis (LSS) is a common condition that affects roughly 20,000 people in the United States.¹ LSS can cause neurogenic claudication, which is associated with significant impairment and diminished quality of life, making LSS the most common reason for spine surgery in the elderly population.¹ Traditionally, surgical management has focused on direct central decompression at the affected level, most commonly through an open or minimally invasive

(MIS) laminectomy. In an effort to minimize surgical trauma and improve recovery time, however, alternative MIS techniques have been introduced in recent years.²

One alternative MIS technique that has gained recent attention is the interspinous process device (IPD). IPDs are implants that are placed between the spinous processes at the level of pathology to relieve central stenosis.² The reported benefits of an interspinous implant are indirect decompression of central stenosis in a manner that causes less tissue trauma and allows for

shorter operative times and faster recoveries than traditional decompression techniques.³ These potential advantages, however, come with many potential pitfalls, including concerns regarding safety and efficacy.

When comparing complication rates of IPDs to traditional direct decompression techniques, such as open and MIS laminectomies, Zhao et al, Wu et al, and Welton et al have shown higher reoperation rates associated with IPDs.⁴⁻⁶ Multiple complications have been reported with IPDs, such as implant loosening, new or persistent pain, spinous process fractures, and implant migration, many of which require further treatment or surgical intervention.³ These complications are often caused by improper patient selection, inappropriate device size, and incorrect positioning.^{3,7,8} Furthermore, by introducing hardware in a procedure intended for decompression alone, IPDs introduce an entirely new class of complications to the realm of surgical decompression without fusion: implant-related complications. These implant-related complications can be so pervasive that they require a product recall, as was the case with the temporary Class II product recall associated with the X-STOP (Medtronic) device from September 2008 to July 2009.⁹

With the growing utilization of this technology, there is a need for continued vigilance to assess its safety and efficacy. The Food and Drug Administration (FDA) Manufacturer and User Facility Device Experience (MAUDE) database, a publicly available repository of adverse events reports, is a particularly useful tool for assessing adverse events associated with an implant. This database provides a near real-time summary of adverse events associated with new implants and technologies and allows for monitoring of their safety and efficacy. The objective of this study, therefore, was to use the MAUDE database to summarize the adverse event reports associated with interspinous implants in hopes of uncovering any concerning trends in the safety or efficacy of these devices. We hypothesized that the most common adverse event types associated with these implants would be inadequate efficacy, implant migration, and spinous process fractures.

MATERIALS AND METHODS

Institutional Review Board approval was attained prior to the completion of this study (IRB 2023–1876). The FDA MAUDE database is a publicly available repository of adverse events reports voluntarily submitted by health care professionals, manufacturers, and patients, providing a comprehensive overview of adverse events associated with medical devices. The database contains

voluntary adverse events reports dating from June 1993, user facility reports since 1991, distributor reports since 1993, and manufacturer reports since August 1996 for medical devices, including IPDs. Mandatory reporting to the FDA is required by manufacturers, importers, and device user facilities. Additionally, voluntary reporting is accepted by health care professionals, patients, and consumers.

We retrospectively reviewed adverse events reports in the MAUDE database regarding IPDs for the treatment of LSS. In order to gather adverse events reports, we selected “Prosthesis, Spinous Process Spacer/Plate” as the product class. All available reports from the creation of the database to November 2022 were queried and included in our analysis. For each complication, we collected the following data: type of adverse event, patient impact, and necessary interventions associated with the adverse event. Narrative reports of each adverse event were used to accurately categorize events for analysis. FDA reports relating to the Coflex (XTant Medical, Belgrade, Montana; FDA approval 10/2012), Superior (Boston Scientific, Marlborough, Massachusetts; FDA approval 5/2015), and X-Stop (Medtronic, Minneapolis, Minnesota; FDA approval 11/2005) were included in our report.

Complications were categorized into 2 groups: (1) unrelated anesthesia or medical events and (2) surgery or implant-related events. Unrelated anesthesia or medical events were eliminated from our analysis in order to focus specifically on implant-related surgical adverse events. Adverse events were further categorized as intraoperative or postoperative to allow for differentiation of events that were associated with complications during implantation vs complications that occurred as postoperative sequelae.

Adverse events were analyzed both in aggregate (across implants), as well as for individual implant manufacturers (within implants). This allowed us to determine what proportion of each adverse event type was attributable to each implant, as well as to determine the relative representation of each adverse event for a given implant. No tests for statistical significance were performed, but counts of adverse events were summed, and proportions were calculated using SPSS statistical software (IBM Corp., Armonk, NY).

RESULTS

Overall Results

Our search yielded a total of 1027 adverse events reports. Thirty-nine of these reports were generated

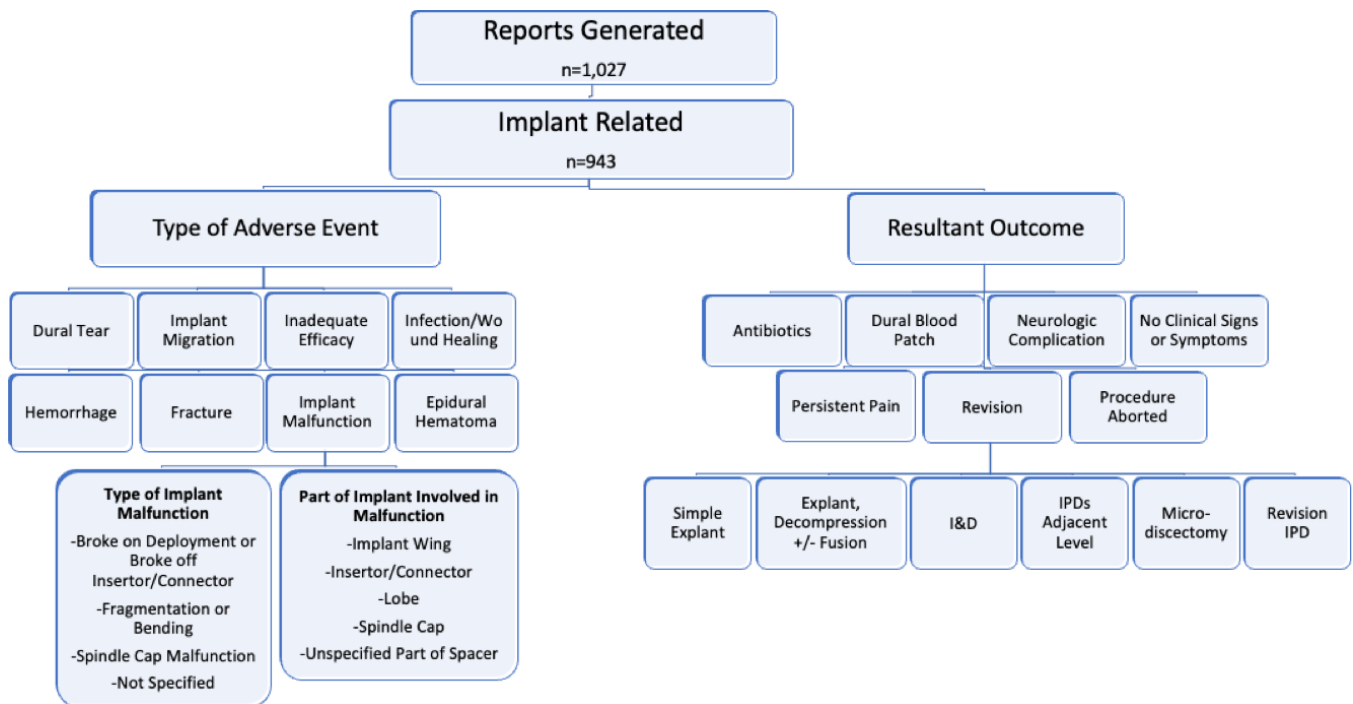


Figure 1. Summary of reports generated from the Manufacturer and User Facility Device Experience database for "Prosthesis, Spinous Process Spacer/Plate."

from published abstracts or manuscripts and were eliminated due to the high likelihood that these could represent duplications of previously submitted reports. Five were technical errors (wrong-level surgery) and were also eliminated. This yielded 983 reports for analysis. Forty of these events were unrelated anesthesia or medical events, while the remaining 943 were surgery related (Figure 1).

Of the 943 surgery-related adverse events, 42.8% (404) were detected intraoperatively and 57.2% (539) were postoperative sequelae. The most common intraoperative events were implant malfunctions (39.7%, $n = 374$) and fractures (2.2%, $n = 21$). The most common

postoperative events were persistent pain (26.6%, $n = 251$), implant migration (19.1%, $n = 180$), and fracture (6.8%, $n = 64$; Table 1). The 2 most common types of implant malfunctions, which were unique to the Supreion implant, were spindle cap malfunction during deployment (52.1%, $n = 195$) and implant breakage off of the insertor/connector (35.8%, $n = 134$; Table 2).

The resultant outcome of each adverse event is summarized in Table 3. While a large number of adverse events had no clinical impact on the patient (aside from unmeasured impacts, such as increased operative time and cost; 34.8%, $n = 328$), the most common resultant outcome of an adverse event was the need for revision

Table 1. Intraoperative and postoperative adverse events related to interspinous process devices.

Complications	Across Implants ^a				Within Implants ^b		
	Coflex	Supreion	X-Stop	Overall, % (n)	Coflex	Supreion	X-Stop
Intraoperative	2.0%	97.0%	1.0%	42.8% (404)	21.1%	47.3%	5.2%
Implant malfunction	2.1%	97.6%	0.3%	39.7% (374)	21.1%	44.1%	1.3%
Fracture	0.0%	85.7%	14.3%	2.2% (21)	0.0%	2.2%	3.9%
Dural tear	0.0%	100.0%	0.0%	0.6% (6)	0.0%	0.7%	0.0%
Hemorrhage	0.0%	100.0%	0.0%	0.2% (2)	0.0%	0.2%	0.0%
Implant migration	0.0%	100.0%	0.0%	0.1% (1)	0.0%	0.1%	0.0%
Postoperative	5.6%	80.9%	13.5%	57.2% (539)	78.9%	52.7%	94.8%
Inadequate efficacy	4.4%	78.9%	16.7%	26.6% (251)	28.9%	23.9%	54.5%
Implant migration	2.8%	89.4%	7.8%	19.1% (180)	13.2%	19.4%	18.2%
Fracture	7.8%	78.1%	14.1%	6.8% (64)	13.2%	6.0%	11.7%
Infection or wound healing	26.7%	60.0%	13.3%	3.2% (30)	21.1%	2.2%	5.2%
Neurological complication	7.7%	61.5%	30.8%	1.4% (13)	2.6%	1.0%	5.2%
Epidural hematoma	0.0%	100.0%	0.0%	0.1% (1)	0.0%	0.1%	0.0%

^aAcross group analysis demonstrates that the proportion of total adverse events for a given adverse event type is attributed to each implant manufacturer.

^bWithin groups analysis demonstrates the contribution of each adverse event to the total adverse events pool for a given manufacturer/implant.

Table 2. Malfunctions of interspinous process devices.

Implant Malfunctions	Across Implants ^a				Within Implants ^b		
	Coflex	Superion	X-Stop	Overall, % (n)	Coflex	Superion	X-Stop
Spindle cap malfunction during deployment	0.0%	100.0%	0.0%	52.1% (195)	0.0%	53.4%	0.0%
Broke on deployment or broke off of inserter/connector	0.0%	100.0%	0.0%	35.8% (134)	0.0%	36.7%	0.0%
Fragmentation/bending	24.2%	75.8%	0.0%	8.8% (33)	100.0%	6.8%	0.0%
Not specified	0.0%	91.7%	8.3%	3.2% (12)	0.0%	3.0%	100.0%

^aAcross group analysis demonstrates that the proportion of total implant malfunctions for a given malfunction type is attributed to each implant manufacturer.

^bWithin groups analysis demonstrates the contribution of malfunction type to the total pool of implant malfunctions for a given manufacturer/implant.

surgery (48.8%, $n = 460$). The following most common outcomes were persistent pain (13.0%, $n = 123$) and the need to abort the index procedure (34.8%, $n = 328$). The need for revision surgery was common in patients who experienced fracture (47.1%), implant migration (84.5%), infection (76.7%), and neurological complications (76.9%; Table 3). Trends in resultant outcomes

over time are presented graphically in Figure 2 and demonstrate increasing numbers of adverse events, especially for implant malfunction, implant migration, and inadequate efficacy.

Among the patients who required revision surgery, inadequate efficacy (46.7%, $n = 215$), implant migration (33.3%, $n = 153$), and fracture (8.7%, $n = 40$) were the

Table 3. Resultant outcome as a function of adverse event type related to interspinous process devices.

Complications	Across Implants ^a				Within Implants ^b			Within Implants, as Percentage of Each Complication ^c			
	Coflex	Superion	X-Stop	Overall	Coflex	Superion	X-Stop	Coflex	Superion	X-Stop	Overall
Implant malfunction	2.1%	97.6%	0.3%	39.7%	21.1%	44.1%	1.3%	-	-	-	-
No clinical signs, symptoms, or conditions	0.0%	100.0%	0.0%	33.8%	0.0%	38.5%	0.0%	0.0%	87.4%	0.0%	85.3%
Persistent pain	2.9%	97.1%	0.0%	3.6%	2.6%	4.0%	0.0%	12.5%	9.0%	0.0%	9.1%
Revision	38.9%	55.6%	5.6%	1.9%	18.4%	1.2%	1.3%	87.5%	2.7%	100.0%	4.8%
Procedure aborted	0.0%	100.0%	0.0%	0.3%	0.0%	0.4%	0.0%	0.0%	0.8%	0.0%	0.8%
Inadequate efficacy	4.4%	78.9%	16.7%	26.6%	28.9%	23.9%	54.5%	-	-	-	-
Revision	4.2%	87.4%	8.4%	22.8%	23.7%	22.7%	23.4%	81.8%	94.9%	42.9%	85.7%
Persistent pain	5.6%	27.8%	66.7%	3.8%	5.3%	1.2%	31.2%	18.2%	5.1%	57.1%	14.3%
Implant migration	2.8%	89.5%	7.7%	19.2%	13.2%	19.6%	18.2%	-	-	-	-
Revision	2.0%	91.5%	6.5%	16.2%	7.9%	16.9%	13.0%	60.0%	86.4%	71.4%	84.5%
Persistent pain	7.7%	84.6%	7.7%	2.8%	5.3%	2.7%	2.6%	40.0%	13.6%	14.3%	14.4%
No clinical signs, symptoms, or conditions	0.0%	0.0%	100.0%	0.2%	0.0%	0.0%	2.6%	0.0%	0.0%	14.3%	1.1%
Fracture	5.9%	80.0%	14.1%	9.0%	13.2%	8.2%	15.6%	-	-	-	-
Revision	10.0%	67.5%	22.5%	4.2%	10.5%	3.3%	11.7%	80.0%	39.7%	75.0%	47.1%
Persistent pain	3.8%	96.2%	0.0%	2.8%	2.6%	3.0%	0.0%	20.0%	36.8%	0.0%	30.6%
Procedure aborted	0.0%	91.7%	8.3%	1.3%	0.0%	1.3%	1.3%	0.0%	16.2%	8.3%	14.1%
No clinical signs, symptoms, or conditions	0.0%	71.4%	28.6%	0.7%	0.0%	0.6%	2.6%	0.0%	7.4%	16.7%	8.2%
Infection or wound healing	26.7%	60.0%	13.3%	3.2%	21.1%	2.2%	5.2%	-	-	-	-
Revision	30.4%	52.2%	17.4%	2.4%	18.4%	1.4%	5.2%	87.5%	66.7%	100.0%	76.7%
Antibiotics	14.3%	85.7%	0.0%	0.7%	2.6%	0.7%	0.0%	12.5%	33.3%	0.0%	23.3%
Neurological complication	7.7%	61.5%	30.8%	1.4%	2.6%	1.0%	5.2%	-	-	-	-
Revision	10.0%	50.0%	40.0%	1.1%	2.6%	0.6%	5.2%	100.0%	62.5%	100.0%	76.9%
Neurological issue	0.0%	100.0%	0.0%	0.3%	0.0%	0.4%	0.0%	0.0%	37.5%	0.0%	23.1%
Dural tear	0.0%	100.0%	0.0%	0.6%	0.0%	0.7%	0.0%	-	-	-	-
Dura blood patch	0.0%	100.0%	0.0%	0.3%	0.0%	0.4%	0.0%	-	50.0%	-	50.0%
Procedure aborted	0.0%	100.0%	0.0%	0.2%	0.0%	0.2%	0.0%	-	33.3%	-	33.3%
Persistent pain	0.0%	100.0%	0.0%	0.1%	0.0%	0.1%	0.0%	-	16.7%	-	16.7%
Hemorrhage	0.0%	100.0%	0.0%	0.2%	0.0%	0.2%	0.0%	-	-	-	-
Procedure aborted	0.0%	100.0%	0.0%	0.2%	0.0%	0.2%	0.0%	-	100.0%	-	100.0%
Epidural hematoma	0.0%	100.0%	0.0%	0.1%	0.0%	0.1%	0.0%	-	-	-	-
Revision	0.0%	100.0%	0.0%	0.1%	0.0%	0.1%	0.0%	-	100.0%	-	100.0%

^aAcross group analysis demonstrates that the proportion of total adverse events for a given adverse event type is attributed to each implant manufacturer.

^bWithin groups analysis demonstrates the contribution of each adverse event to the total adverse events pool for a given manufacturer/implant.

^cWithin groups analysis as a percentage of each complication indicates the percentage of each outcome for each implant within a specific adverse event type.

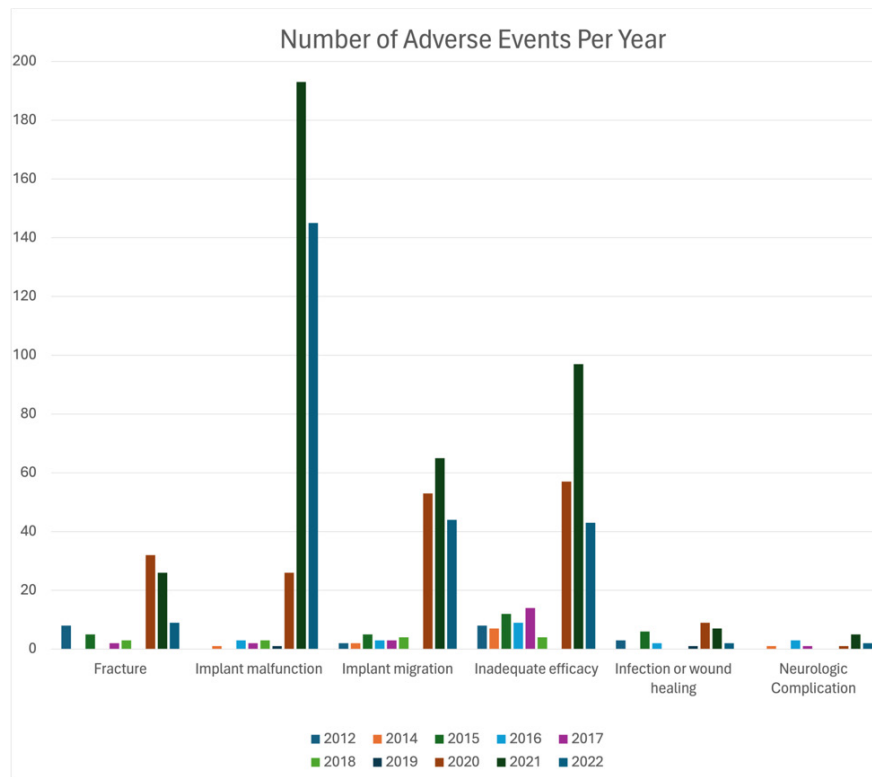


Figure 2. Number of adverse events of each type per year.

most common indications (Table 4). Implant migration, fracture, and implant malfunction, 3 complications that are unique to decompression with an IPD as compared with traditional laminectomy, accounted for 45.9% of revisions (211/460), and revision was required in 33.0% of cases where 1 of these complications was reported (211/640).

Among patients who had the index procedure aborted, fracture (61.1%, $n = 11$), implant malfunction (16.7%, $n = 3$), and dural tears (11.1%, $n = 2$) were the most common reasons. The most common revision procedures were simple explant (59.9%, $n = 226$), revision

IPD (22.3%, $n = 84$), and explant with decompression with or without fusion (14.3%, $n = 54$; Table 5).

Implant-Specific Events

Coflex

There were 38 adverse events reports associated with the Coflex device, accounting for 4.0% of all adverse events reports. A total of 21.1% were intra-operative, while 78.9% were postoperative (Table 1). The most common complications were inadequate efficacy (28.9%, $n = 11$), implant malfunction (21.1%,

Table 4. Reasons for revision after placement of interspinous process devices.

Revision ($n = 460$)	Across Implants ^a				Within Implants ^b		
	Coflex	Superion	X-Stop	Overall, % (n)	Coflex	Superion	X-Stop
Inadequate efficacy	4.2%	87.4%	8.4%	46.7% (215)	29.0%	49.1%	39.1%
Implant migration	2.0%	91.5%	6.5%	33.3% (153)	9.7%	36.6%	21.7%
Fracture	10.0%	67.5%	22.5%	8.7% (40)	12.9%	7.0%	19.6%
Infection or wound healing	30.4%	52.2%	17.4%	5.0% (23)	22.6%	3.1%	8.7%
Implant malfunction	38.9%	55.6%	5.6%	3.9% (18)	22.6%	2.6%	2.2%
Neurological complication	10.0%	50.0%	40.0%	2.2% (10)	3.2%	1.3%	8.7%
Epidural hematoma	0.0%	100.0%	0.0%	0.2% (1)	0.0%	0.3%	0.0%
IPD-specific complication	6.6%	83.9%	9.5%	45.9% (211)	45.2%	46.2%	43.5%

Abbreviation: IPD, interspinous process device.

^aThe percentage of revisions attributed to each implant brand is shown for each revision indication.

^bThe percentage of overall revisions indicates the percentage of overall revisions attributed to each adverse event type. IPD-specific complications include implant migration, fracture, and implant malfunction.

Table 5. Type of revision after interspinous process device placement.

Type of Revision	Across Implants				Within Implants		
	Coflex	Superion	X-Stop	Overall, % (n)	Coflex	Superion	X-Stop
Explant	5.3%	88.5%	6.2%	59.9% (226)	40.0%	67.1%	28.6%
Revision interspinous implant	7.1%	88.1%	4.8%	22.3% (84)	20.0%	24.8%	8.2%
Explant, decompression ± fusion	16.7%	31.5%	51.9%	14.3% (54)	0.0%	0.3%	0.0%
Irrigation and debridement	33.3%	66.7%	0.0%	1.6% (6)	6.7%	1.3%	0.0%
Unknown	20.0%	20.0%	60.0%	1.3% (5)	3.3%	0.3%	6.1%
Implanted level above	0.0%	100.0%	0.0%	0.3% (1)	0.0%	0.3%	0.0%
Microdiscectomy	0.0%	100.0%	0.0%	0.3% (1)	30.0%	5.7%	57.1%

$n = 8$), and infection or wound healing complications (21.1%, $n = 8$; Table 3). There were 8 reports of implant malfunction, all of which were due to fragmentation/bending (Table 2).

The most common resultant outcomes were revision (81.6%, $n = 31$) and persistent pain (15.8%, $n = 6$). The need for revision surgery was common in patients who experienced implant malfunction (87.5%), fracture (80.0%), implant migration (60.0%), infection (87.5%), and neurological complications (100.0%; Table 3). Implant migration, fracture, and implant malfunction accounted for 45.2% of Coflex revisions (Table 4). Among the patients who required revision, the most common revision types were simple explant (40%, $n = 12$), explant with decompression with or without fusion (30%, $n = 9$), and revision interspinous implants (20%, $n = 6$; Table 5).

Superion

There were 828 adverse events reports associated with the Superion device, accounting for 87.8% of all adverse events reports. A total of 47.3% were intraoperative, while 52.7% were postoperative (Table 1). The most common complications were implant malfunction (44.1%, $n = 365$), inadequate efficacy (23.9%, $n = 198$), and implant migration (19.6%, $n = 162$; Table 3). Of the 365 reports of implant malfunction, the most common were spindle cap malfunction during deployment (53.4%, $n = 195$), implant breaking on deployment or breaking off of the connector/insertor (36.7%, $n = 134$), and implant fragmentation/bending (6.8%, $n = 25$; Table 2). The most common parts of the implant involved in malfunction were the spindle cap (53.7%, $n = 196$), an unspecified part of the spacer (40.0%, $n = 146$), and the insertor/connector (4.9%, $n = 18$).

The most common resultant outcomes were revision (46.3%, $n = 383$), no clinical signs or symptoms (39.1%, $n = 324$), and persistent pain (11.0%, $n = 91$). The need for revision surgery was common in patients who experienced implant migration (86.4%), fracture (39.7%), infection (66.7%), and neurological complications

(62.5%; Table 3). Implant migration, fracture, and implant malfunction accounted for 46.2% of Superion revisions. Unlike with the Coflex, the risk of revision associated with implant malfunction was low (2.6%), but migration, fracture, and implant malfunction accounted for 46.2% of Superion revisions (Table 4). Among the patients who required revision, the most common revision types were simple explant (67.1%, $n = 200$), revision interspinous implants (24.8%, $n = 74$), and explant with decompression with or without fusion (5.7%, $n = 17$; Table 5).

X-Stop

There were 77 adverse events reports associated with the X-Stop device, accounting for 8.2% of all adverse events reports. A total of 5.2% were intraoperative, while 94.8% were postoperative (Table 1). The most common complications were inadequate efficacy (54.5%, $n = 42$), implant migration (18.2%, $n = 14$), and fracture (15.6%, $n = 12$; Table 3). There was 1 report of implant malfunction, the cause of which was not specified.

The most common resultant outcomes were revision (59.7%, $n = 46$), persistent pain (33.8%, $n = 26$), and no clinical signs or symptoms (5.2%, $n = 4$). The need for revision surgery was common in patients who experienced fracture (75.0%), implant malfunction (100.0%), implant migration (71.4%), infection (100.0%), and neurological complications (100.0%; Table 3). Unlike with the Coflex, the risk of revision associated with implant malfunction was low (2.2%), but migration, fracture, and implant malfunction accounted for 43.5% of X-Stop revisions (Table 4). Among the patients who required revision, the most common revision types were explant with decompression ± fusion (57.1%, $n = 54$), explant (28.6%, $n = 14$), and revision interspinous implants (8.2%, $n = 4$; Table 5).

DISCUSSION

IPDs have emerged as a popular treatment option for patients with degenerative LSS, providing a less invasive alternative to traditional surgical interventions.² However, the safety and efficacy of these implants have been a topic of debate within the orthopedic community. In the current study, we assessed 943 adverse events that were reported to the FDA and detailed in the MAUDE database. Implant malfunctions (39.7%), inadequate efficacy (26.6%), implant migration (19.2%), and fracture (9.0%) were the most common postoperative adverse events. Among the most common adverse events, revision surgery was often required for definitive treatment (fracture 47.1%, implant migration 84.5%, and inadequate efficacy 85.7%).

There were implant-specific differences in the representation of various adverse events. While fracture, implant migration, and inadequate efficacy represented a large proportion of the adverse events in each group, there were large differences in the proportion of certain adverse events. Implant malfunction, for example, represented a large proportion of the adverse events for the Superion (44.1%) and Coflex (21.1%) but only a small proportion for the X-Stop (1.3%). The high proportion of implant-related malfunctions in the Superion group may reflect a systematic issue with the intraoperative placement or deployment of these devices. Given the high proportion of reports indicating malfunctions related to the spindle cap and the spacer implant itself, it is likely that 1 or both of these components have mechanical or technical shortcomings that have contributed to the high proportion of mechanical failures for this implant. Interestingly, while the X-Stop implant underwent a temporary Class II recall in 2009⁹ due to “potential to cause damage to and/or breakage of the... device’s universal wing assembly,” there have been no such recalls for the Superion despite the high number of the implant malfunctions identified in the current study. While implant malfunctions only required revision in 2.7% of the Superion cases, they required revision in 87.5% of Coflex reports.

Another implant-specific difference is that a far higher proportion of the adverse events reports for the X-Stop relate to inadequate efficacy (54.5%) in comparison to the Coflex (28.9%) and Superion (23.9%). Such disparities may highlight important shortcomings related to the degree of relief provided by these implants.

One of the most important findings in the current study is the high likelihood of revision surgery for complications that specifically can be attributed to the IPD.

Because spinous process fractures, implant migration, and implant malfunction are implant specific, these represent a fairly unique adverse event class compared with traditional uninstrumented laminectomy for decompression. Implant migration, implant malfunction, and fracture accounted for roughly 45% of revisions, and one-third of cases where these adverse events were present required a revision surgery.

The results of the current study corroborate the findings of multiple clinical trials reporting adverse events associated with IPDs. In a 2-year study assessing clinical outcomes of the X-Stop device, Hartjen et al¹⁰ identified a 3.6% risk of persistent pain, 1.8% risk of device migration, and a 1.8% risk of spinous process fracture. Nearly 10% of these patients required explant with or without further decompression within the study period due to adverse events or inadequate efficacy. Similarly, Verhoof et al¹¹ found a 58% revision rate within 2 years following insertion of the X-Stop spacer. In a clinical study assessing the Superion device, a total of 44.4% of patients experienced device-related complications, including device migration (11.1%) and spinous process fractures (21.1%). A total of 20.1% of patients required device explantation, and 24.3% required same-site revision surgery.⁶ Taken together, the results of this analysis and prior studies demonstrate a concerning number of adverse events associated specifically with the efficacy and safety of IPDs that should lead treating physicians to question the use of these devices over standard uninstrumented decompression techniques.

We believe that while the current study cannot make definitive conclusions regarding the safety and efficacy of these implants, it does alert surgeons to the potential adverse events associated with IPDs. This is especially useful in the setting of implant-related complications, which were common. Surgeons should be aware that implants often malfunction on deployment or cause spinous process fractures on deployment. Specific technical pearls can also be derived from these data. For example, surgeons should be aware that Coflex implants are subject to intraoperative fragmentation and bending, while Superion implants are more prone to spine cap malfunction, implant breakage on deployment, and implant breakage off the insertor/connector. These findings may help surgeons determine whether they would like to perform IPDs, which implants they might prefer, and which technical aspects of the surgical procedure require additional caution or vigilance.

This study had many limitations. First, because the MAUDE database lacks data on the number of implants placed over the period under analysis, it

is difficult to ascertain a prevalence or incidence. Therefore, it is difficult to compare the number of adverse events reports between implants in any meaningful way, as a high number of adverse events reports may simply be a reflection of a higher utilization of a given implant. This makes it impossible to comment directly on the comparative safety and efficacy of these implants. Second, the MAUDE database relies on voluntary reporting, which may lead to underrepresentation of certain adverse events. Third, this database lacks patient- and surgery-specific details that may contribute to the occurrence of an adverse event, such as bone quality, anatomical abnormalities, or other conditions. Finally, because these implants are often placed by nonsurgeon physicians, it is possible that technical errors surrounding proper implant placement techniques may be partially responsible for many of the adverse events seen in this report, including implant malfunction, fractures, and implant migration.

CONCLUSIONS

Per review of the FDA MAUDE database for adverse events, there are a large number of adverse events reports associated with IPDs. The most common reports are for implant malfunction, inadequate efficacy, implant migration, and fracture. While inadequate efficacy and continued pain are possible with traditional decompression techniques, the use of IPDs introduces hardware-related complications to the realm of surgical decompression without fusion (implant malfunction, implant migration, and fracture). Concerningly, these complications require revision surgery in one-third of cases. Furthermore, implant-specific assessments demonstrate a high prevalence of implant malfunctions for the Coflex and Superior implants. While these rarely led to revision surgery in the Superior, nearly 90% of Coflex implant malfunctions led to revision surgery.

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Corresponding Author: Gregory S. Kazarian, Hospital for Special Surgery, 535 E 70th St, New York, NY 10021, USA; kazariang@hss.edu

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