



A case of tubular adenoma developing after bladder augmentation: Case report and literature review

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ABSTRACT

INTRODUCTION: We encountered a rare case of tubular adenoma developing after bladder augmentation. We here report our case as well as summarize reports in the literature on adenomas developing after bladder augmentation.

PRESENTATION OF CASE: A 23-year-old man came to our hospital for routine surveillance cystoscopy. He was born with a lipomyelomeningocele and neurogenic bladder with low bladder compliance, and hence his bladder was routinely emptied by clean intermittent catheterization. He was also treated with anticholinergic agents. However, because the patient's neurogenic bladder was unstable, he underwent sigmoidocolocystoplasty when he was 8-years old. After the bladder augmentation, he was examined annually by surveillance cystoscopy.

On cystoscopy, a 5-mm pedunculated polyp was found on the front side of the sigmoid colon cap. Therefore, we performed snare polypectomy together with electrocoagulation under cystoscopy. The patient's final diagnosis was tubular adenoma (mild atypia) with no malignancy, as assessed by histopathology. There has been no evidence of recurrence after the polypectomy on routine surveillance cystoscopy.

DISCUSSION: To the best of our knowledge, there have been 11 cases of adenoma occurring after bladder augmentation reported in the literature, including our present case. There are several carcinogenic pathways associated with colorectal oncogenesis. Adenomas that are larger than 1.0 cm in diameter with a marked villous component have a high risk of oncogenesis.

CONCLUSION: We believe that the early detection of carcinoma or adenoma and their treatment at an early stage is crucial. Therefore, we recommend routine surveillance cystoscopy for patients after bladder augmentation.

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1. Introduction

This work has been reported in line with the CARE criteria [1]. Enterocystoplasty (ECP) using the colon, ileum, or stomach has become an accepted reconstructive option for patients with intractable incontinence and poor bladder compliance owing to various neurogenic and non-neurogenic disorders [2].

However, recently, there have been an increasing number of reports of benign and malignant tumors developing in the neobladder of post-ECP patients, particularly around the line of anastomosis between the colon cap and the native bladder remnant, which are detected on long-term follow-up [2]. We previously reported that 55 cases of malignancy occurring after ECP have been published

in the literature [3]. Although *N*-nitrosamines were reported to be associated with carcinogenesis following bladder augmentation, the mechanisms involved remained unclear [4]. There are several carcinogenic pathways associated with colorectal oncogenesis [5]. Adenomas that are greater than 1.0 cm in diameter, and with a marked villous component have a high risk of developing into cancer [5]. Therefore, the early detection of adenomas is important as it will enable early treatment, and therefore the prevention of malignant changes.

We routinely perform annual surveillance cystoscopy in patients who have undergone bladder augmentation in our departments. One such patient was found to have developed tubular adenoma at the patch site of the sigmoid colon cap. The incidence of adenoma after ECP is rare. To the best of our knowledge, there are 11 cases of adenoma reported in the literature, including our present case [4,6–14]. Here we report our rare case and review the literature on cases of benign tumors that occurred after ECP.

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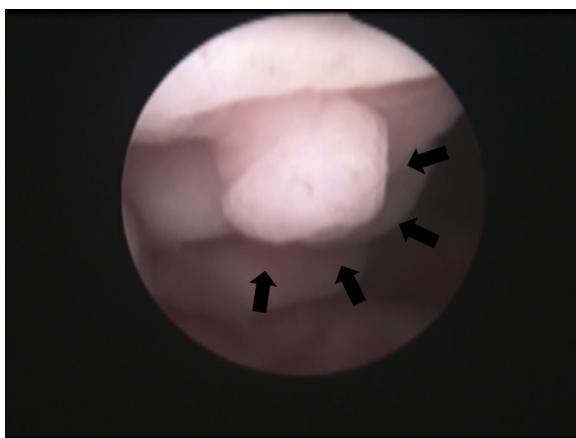


Fig. 1. Cystoscopic findings before transurethral resection.

An image of the patient's bladder. A 5-mm pedunculated polyp was found on the front side of the colon cap. There were no abnormalities in other areas of the bladder, including in the anastomosis between the colon cap and the native bladder (arrows).

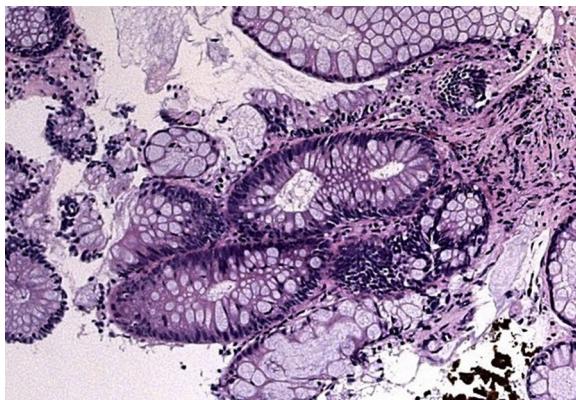


Fig. 2. Histopathological analysis of the tumor on biopsy.

Hematoxylin and eosin staining of the tumor specimen ($\times 40$). The tumor was contained within the glandular component. No malignant changes were detected. The diagnosis was tubular adenoma.

2. Case report

A 23-year-old man came to our hospital for routine surveillance cystoscopy. He had no symptoms, such as upper tract infection, flank pain, and hematuria. He was born with a lipomyelomeningocele and neurogenic bladder with low bladder compliance. He had been undergoing clean intermittent catheterization and taking anti-cholinergic agents since he was 3-years old. However, his neurogenic bladder was unstable, and his bladder capacity decreased and his bladder compliance worsened. He hence underwent sigmoidocolocystoplasty when he was 8-years old. His postoperative course was stable, and his bladder capacity increased and his urinary continence improved. The patient has been undergoing annual routine surveillance cystoscopy from 1 year after the bladder augmentation. The patient has no family history of bladder and colorectal disease.

There were and no abnormalities on laboratory data before cystoscopy. Tumor markers, such as α -fetoprotein, carcinoembryonic antigen, and carbohydrate antigen 19–9 were not increased.

Upon cystoscopy, there were no abnormalities around the anastomosis between the native bladder and sigmoid colon. However, a 5-mm pedunculated polyp was found on the front side of the sigmoid colon cap, and hence we performed a biopsy (Fig. 1). On histopathology, the polyp was identified as tubular adenoma (Fig. 2). Therefore, we performed snare polypectomy together with



Fig. 3. Cystoscopic findings at the site of transurethral resection.

Image of the site of resection after snare polypectomy together with electrocoagulation. No perforations or bleeding was detected.

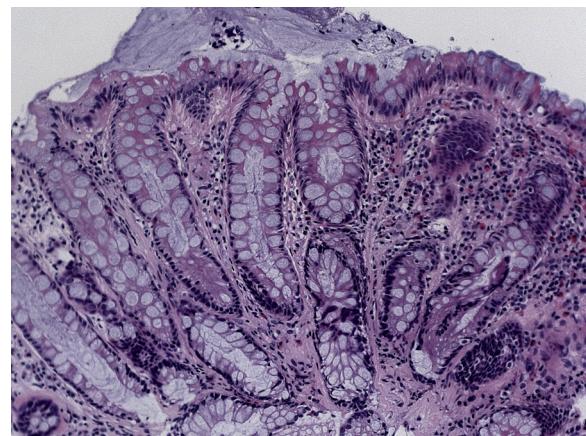


Fig. 4. Histopathological analysis of the tumor on transurethral resection.

A high magnification image of the tumor ($\times 400$) showing the presence of glandular epithelium. No malignant changes were detected.

electrocoagulation under cystoscopy (Fig. 3). The patient's post-operative course was uneventful and his bladder function was maintained. His bladder was drained for 24 h. His final diagnosis was tubular adenoma (mild atypia) and no malignancy was found on histopathology (Fig. 4a,b). We performed routine annual surveillance cystoscopy from 1 year after the polypectomy, because we believe that the early diagnosis of adenoma will decrease the risk of malignancy development. During the subsequent 5 years, there has been no evidence of recurrence.

3. Discussion

Recent reports have suggested an increased incidence of malignancy after ECP [2,15–18]. However, the incidence of adenoma is rare. To the best of our knowledge, there have been 11 cases of adenoma occurring after bladder augmentation reported in the literature, including our present case [4,6–14] (Table 1). The sex distribution of the patients was 7 men (64%) and 4 women (36%). The age at ECP ranged from 5 to 50 years (mean: 27.1 years). The original underlying disease was urinary tuberculosis in 5 cases (45%), neurogenic bladder in 3 cases (27%), rhabdomyosarcoma in 1 case (9%), post bladder surgery complications in 1 case (9%), and detrusor instability in 1 case (9%). Symptoms included hematuria in 6 (55%), lumbar pain and oliguria in 1 (9%), frequent urination in 1 (9%), recurrent pyelonephritis in 1 (9%), and no symptoms (detected

Table 1
Summary of our case and a literature review of cases of adenoma occurring after bladder augmentation.

Authors	Sex	Age at ECP (yrs)	Original disease	Segment for ECP	Duration from ECP (yrs)	Symptoms	Site of tumor	Pathology	Size(mm)	Treatment	Recurrence
King et al. [6]	M	20	Tb	Ileum	30	Dysuria	Ileum	TVA	75	TUR	(–)
Gepi-Attee et al. [7]	M	28	Tb	Colon	25	Frequent urination	Colon	VA	60	Ileocystoplasty	(–)
Gousse et al. [8]	F	77	DI	Colon	5	Hematuria	Colon	TVA	35	Ileocystoplasty	(–)
Yip et al. [9]	F	14	Tb	Colon	24	Hematuria	Colon	VA	ND	TUR	(+)
Yamada et al. [10]	F	18	Tb	Ileum	44	Hematuria	Anastomosis	TVA	ND	TUR	(–)
Armah et al. [11]	M	5	RMS	Ileum	34	Hematuria	Anastomosis	TA	50	TUR	(–)
Elphick et al. [4]	M	50	NB	Colon	12	Hematuria	Colon	TA	25	TUR	(–)
Husillos Alonso et al. [12]	M	42	Tb	Colon	24	Lumbar pain/Oliguria	Anastomosis	VA	ND	TUR	(–)
Rubino et al. [13]	M	ND	MBS	Colon	ND	Pyelonephritis	Anastomosis	TVA	40	Resection	(+)
Lin et al. [14]	F	9	NB	Stomach	9	Hematuria	Native bladder	Adenocarcinoma in adenoma	18	Resection	(–)
Our case	M	8	NB	Colon	15	Routine surveillance	Colon	TA	5	TUR	(–)

ECP: enterocystoplasty, M: male patient, F: female patient, ND: not described, Tb: tuberculosis, DI: detrusor instability, RMS: rhabdomyosarcoma, NB: neurogenic bladder, MBS: multiple bladder surgery, TVA: tubulovillous adenoma, VA: villous adenoma, TA: tubular adenoma, TUR: transurethral resection.

on a routine examination) in 1 (9%). Bladder augmentation involved the colon in 7 cases (64%), small bowel in 3 cases (27%), and stomach in 1 case (9%). Histopathology confirmed tubulovillous adenoma in 4 cases (36%), villous adenoma in 3 cases (27%), tubular adenoma in 3 cases (27%), and adenocarcinoma in the tubulovillous adenoma in 1 case (9%). Adenoma arose from the anastomosis in 4 cases (36%), intestinal cap in 6 cases (55%), and native bladder in 1 case (9%). Treatment was transurethral resection in 7 cases (64%), cystectomy and ECP using other segments of the intestine in 2 cases (18%), and partial resection in 2 cases (18%). There were 2 recurrent cases in the literature [9,13].

The mechanism of colorectal cancer occurrence has been analyzed in detail. There are at least 4 types of carcinogenic pathways that are associated with colorectal oncogenesis, namely, the adenoma–carcinoma sequence type, the *de novo* type, the hereditary non-polyposis colorectal cancer type, and the colitic cancer type [5]. Among these carcinogenic pathways, the *de novo* type and the adenoma–carcinoma sequence type are thought to be particularly important.

In the *de novo* type pathway, colorectal cancer develops directly in the normal colorectal mucosa without adenoma. In the 1980s, several Japanese researchers reported that they detected a flat type of carcinoma with a diameter of less than 10 mm arising *de novo*, which tended to reach the deeper layers at an earlier stage than polypoid-type carcinoma in adenoma [5].

In the adenoma–carcinoma sequence pathway, an adenoma initially forms and increases in size, and then a carcinoma forms in the adenoma. This type of carcinogenesis is associated with several genes, such as the adenomatous polyposis coli gene, *K-ras* gene, *p53* gene, and deleted in colorectal carcinoma gene. Adenomas are well-demarcated lumps of epithelial tumor cells, which can be classified into the following 3 major histological types: tubular, villous, and tubulovillous. The mechanism of the adenoma–carcinoma sequence in colonic segments transposed to the urinary tract has not been established, but may be associated with chronic infection and inflammation, possibly via the reduction of urinary nitrates to nitrates by bacteria, accompanied with the formation of *N*-nitrosamines, which are potent carcinogens [4]. Adenomas that are larger than 1.0 cm in diameter with a marked villous component have the risk of developing into cancer [5]. Therefore, we believe that the early diagnosis of adenoma should decrease the risk of malignancy development.

The use of routine surveillance cystoscopy and cytology is controversial [19]. Higuchi et al. [20] reported that no tumors developed in their patients during a median surveillance time of 15 years after ECP. Furthermore, they reported that they discontinued performing annual surveillance endoscopy and cytology because of the low incidence of malignancy, the lack of proven benefit, and for cost containment. Hamid et al. [21] also reported that it is not necessary to perform annual check cystoscopies in patients with augmented bladders at least for the first 15 years after surgery. However, they recommended that if the patient develops hematuria or other symptoms of concern, including suprapubic pain or recurrent unexplained urinary tract infections, a full evaluation including cystoscopy should be performed. Soergel et al. and Biers et al. [16,17] recommended that routine cystoscopic surveillance of all patients with a history of bladder augmentation should be performed from 10 years after the initial surgery. On the other hand, Lin et al. [14] recommended that long-term active surveillance is necessary for all patients who undergo gastrocystoplasty, because they encountered an extremely rare case of a tubulovillous adenoma that transformed into adenocarcinoma in the native bladder segment of an augmented bladder 9 years after surgery. We also perform annual cystoscopic evaluations and cytological examinations in patients after bladder augmentation in our departments. We encountered 1 case of sarcoma occurring 13 years after bladder

augmentation [3], and 1 case of adenoma occurring 15 years after bladder augmentation. Most cases of malignancy occur more than 10 years after bladder augmentation. However, some cases occur within 10 years after bladder augmentation [3]. We cannot predict the time of occurrence of malignancy. Therefore, we believe that the early detection of carcinoma or adenoma should lead to their successful treatment. We also recommend routine annual routine cystoscopy from 1 year after bladder augmentation, because the bladder mucosa can be directly examined by this technique.

4. Conclusions

We reported a rare case of adenoma developing after sigmoidocolocystoplasty. We recommend performing routine annual surveillance in patients with bladder augmentation for the early detection of tumors.

Conflict of interest

None.

Funding

No funding was received for this study.

Ethical approval

This is a case report and hence ethical approval has not been received.

Consent

Written informed consent was obtained from the patient for publication of this case report and accompanying images.

Authors contribution

Yutaka Hayashi was the attending physician and wrote the paper.

Satoko Shiyani treated the patient.

Itsuro Nagae treated the patient.

Tetsuo Ishizaki treated the patient.

Kazuhiko Kasuya treated the patient.

Kenji Katsumata treated the patient.

Atsuyuki Yamataka supervised the patient's treatment.

Akihiko Tsuchida supervised writing of this paper and treated the patient.

Guarantor

Yutaka Hayashi accepts full responsibility for this case report, and can be contacted via e-mail at rimpoo@tokyo-med.ac.jp.

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