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Complications of In-Dwelling Venous Access Devices: a Single Institution Experience

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Abstract

Objective: To determine the complications of venous access devices (VADs) in cancer patients.

Setting: Retrospective study in a tertiary referral center with specialist hematology and oncology services.

Subjects: First one hundred consecutive patients who were implanted a VAD. All patients had an underlying cancer and the devices were inserted by the same surgeon. The duration of use of VADs and causes of their premature removal were noted.

Results: One hundred VADs (55 port-a-caths and 45 Hickman's lines) were inserted in a total of 89 patients over a 7.5 year period. Majority of patients had acute myeloid leukemia (22) gastrointestinal malignancies (20) breast cancer (19) and genito-urinary cancers (15). The mean duration of use was 110 days; 157 days for the port-a-cath and 53 days for the Hickman's line. Nineteen devices (10 port-a-caths and 9 Hickman's lines) had to be removed prematurely. Two Hickman's lines got removed accidentally. The causes of premature removal included device failure (9), exit site infection (4), luminal infection (3) and tunnel infection (3).

Conclusion: The mean duration of use and the complication rates are comparable with studies reported in the literature (JPMA 51:434,2001).

Introduction

Management of cancer has seen major advancements in the last few decades in the form of chemotherapeutics and supportive care. Modern chemotherapeutic management depends upon repeated and safe access to the venous system for the delivery of drugs, fluids and blood products and on the periodic monitoring of the effects of treatment¹. Peripheral veins are rapidly destroyed by repeated venipuncture and by long term chemotherapy. In addition, thrombosis and sclerosis of peripheral veins enhances the risk of drug extravasation and tissue necrosis².

A variety of methods have been used to provide long term venous access. Surgically constructed arteriovenous fistulas have been used in the past with variable success. Although such fistulas are ideal for hemodialysis they have not been found satisfactory for intermittent long-term chemotherapy. Recently indwelling silicone rubber catheters placed in the superior vena cava have enjoyed greater popularity. Their use has been ideal in avoiding multiple needle pricks and local thrombophlebitis due to the sclerosing effect of chemotherapeutic agents^{3,4}. The long-term venous access devices (VADs) have also helped to overcome the need for repeated peripheral or central venous puncture. The Hickman line is an indwelling silicone catheter, a modification of Broviac's catheter^{3,4}. The Port-a-cath is a totally implantable venous access device in which a conventional central venous catheter is attached to a subcutaneous injection port usually on the chest wall⁵. With both these devices, chemotherapy and supportive care requirements can be administered safely, easily and without the need for repeated venipuncture^{6,7}. But despite the benefits of these devices there are certain risks. Lack of expertise in insertion, as well as the potential for thrombosis and catheter infections are hazards that must be anticipated⁸⁻¹⁰. The use of these devices is now beginning to gain widespread popularity in the developing countries. The purpose of this study was to report our initial experience with VADs, assess

the nature and causes of complications and to see whether the complication rate was comparable to the western data.

Patients and Methods

The data were collected retrospectively. The clinical records were retrieved by the medical record section of the hospital. The section uses the International Classification of Diseases, Version 9.0 (ICD 9.0) data base. At the time of insertion of the VAD, the procedure is logged on to the face sheet by the attending surgeon. Data of patients presenting to the hematology and oncology service of the hospital between Jan 1988 and December 2000 and implanted with a VAD was retrieved. First 100 patients who received the device were the subjects of this study. The fate of the devices on 31st December 1996 was noted. The VADs were all inserted at this hospital by a vascular surgeon (ZS) under operating room conditions, either percutaneously or by employing the venous cut down technique. Insertion of the VAD was always followed by a chest x-ray to check the position of the tip of the catheter. Towards the later part of the study, VADs were inserted under fluoroscopic control. The tip of the catheter was aimed to be positioned at the level of the fourth thoracic vertebra. The total duration of use of the VAD as well as reasons for discontinuation of use were studied. The data were analyzed on Epiinfo Version 6 and MS Dos FoxPro II.

Results

Our hundred VADs were inserted in 89 cancer patients over a period of 7.5 years. Of the devices implanted, 55 were Port-a-caths while 45 were Hickman lines. These were used for administration of chemotherapeutic agents, blood products, parenteral nutrition, and for laboratory sampling. Among the patients there were 47 females and 42 males. The mean age of patients at the time of insertion of the VAD was 42.3 years. All patients had underlying malignant conditions, which were both hematological as well as non-hematological. Acute myeloid leukemia (AML) was the most common underlying diagnosis (22%), followed by gastrointestinal tract malignancies and breast cancer. The details of various underlying malignancies are shown in Table 1.

Table 1. Underlying conditions.

Types of Malignancy	Port-A-cath.	Haickman's	Total
Acute Myeloid Leukemia	11	11	22
GI Malignancies	13	7	20
Breast Cancer	14	5	19
Genito-urinary Cancers	7	8	15
Sarcomas	3	4	7
Non Hodgkins Lymphoma	4	3	7
Acute Lymphoblastic Lymphoma	1	5	6
Head and Neck Cancer	1	1	2
Lung Cancer	1	0	1
Chronic Myeloid Leukemia	0	1	1
Total	55	45	100

Laboratory investigations carried out at the time of insertion showed a mean hemoglobin of 9.3 ± 1.6 gm/dl, total leukocyte count $4.2 \pm 4.4 \times 10^9$ /L and platelet count $222 \pm 186 \times 10^9$ /L. The coagulation profile was available in 25 of the patients and was significantly deranged in 10 of them (40%). The most common site of insertion was the subclavian vein (69%), followed by the external jugular vein. The details are shown in Table 2.

Table 2. Site of insertion.

Site of insertion	Total
Subclavian Vein	69
External Jugular Vein	16
Internal Jugular Vein	7
Femoral Vein	6
Saphenous Vein	1
Cephalic Vein	1
Total	100

The fate of VADs was also studied, and the data are shown in Table 3.

Table 3. Fate of catheters.

Fate of catheters	Port-A- cath.		Hickman		Total	Mean duration
	No.	%	No.	%		
Patient died	19	42.2	21	46.7	40	65 (1-112)
Premature removal	10	18.1	9	20.0	19	97 (7-171)
Lost to follow up	8	14.5	6	13.3	14	
End of use	6	10.9	7	15.6	13	224 (56-1155)
In use	12	21.8	0	-	12	175 (33-480)
Accidental removal	0		2	4.4	2	16 (2-30)
Total	55		45		100	110 (1-1155)

Most commonly the patient died with an intact functioning VAD (40%). This was followed by premature removal (19%). A total of 14 patients were either referred or were lost to follow up and the fate of their catheters was not known. Thirteen VADs were removed after completion of the chemotherapy program. Twelve catheters were still in use with a mean duration of use of 175 days. Two Hickman lines got pulled out accidentally 2 and 30 days after insertion.

A total of 19 patients (19%) had premature removal of their VAD's, out of which 10 were Port-a-caths and 9 were Hickman lines. Causes of premature removal included luminal infections, tunnel infections, exit site infections, and device failure. This is shown in Table 4.

Table 4. Causes of premature removal..

Causes of premature removal	Portacath	Hickmans	Total
Device Failure	8	1	9
Exit Site Infection	1	3	4
Luminal Infection	0	3	3
Tunnel Infection	0	3	3
Total	9	10	19

The most common cause for premature removal was device failure (8/19), while 4 patients had multiple causes of premature removal.

The VADs remained in use for a total of 10964 catheter days. The mean duration of use of all VADs was 110 days, with a mean of 157 days for the Port-a-cath, and 53 days for the Hickman line. The mean duration for which the VADs remained implanted till they were removed was 224 days. A total of 6 Port-a-caths were removed at the end of use after a mean duration of 362 days, whereas 7 Hickman lines were removed after a mean duration of 106 days.

The mean duration of use when the VAD had to be removed prematurely, was 97 days (139 days for 10 Port-a-caths and 51 days for 9 Hickman lines), while the mean duration of use in the patients who died with the VAD in place was 65 days (92 days for 19 Port-a-cath and 41 days for 21 Hickman lines).

Discussion

Insertion of VADs is now being attempted on an outpatient basis. Reported success has been comparable to surgical insertion performed as an inpatient procedure¹¹. Moreover devices have now been designed to function as implanted arm ports¹².

The choice of catheter to be inserted is largely dependent upon individual preferences, however, sometimes the type of malignancy, life expectancy of the patient and financial issues make the decision. For example, Hickman lines were generally preferred if the catheter was being implanted to provide terminal supportive care or the underlying malignancy was acute myeloid leukemia requiring prolonged hospital stay for induction chemotherapy and supportive care during the neutropenic phase. In our country the cost of insertion of a Hickman line is about 35% less than the cost of insertion of a Port-a-cath and hence the Hickman lines were preferred where financial constraints were an issue on

the part of the patient. Reports are emerging that despite its higher initial cost a Port-a-cath used for 4 months is in the long-term less expensive than a Hickman line which has to be flushed every other day with heparin saline for this period⁵. Long-term cost considerations may influence the ultimate choice of device in the future.

Catheter-related infections as well as device failure are significant problems seen with the use of these VADs, occurring in upto 20% of all patients². In our study 19% (n= 19) catheters were removed prematurely because of complications. This is consistent with several studies, which have shown rates of premature removal varying from 10% to 30.2%^{13,15}. Catheter related infections were the most common cause of premature removal seen in our study, which occurred in 10% of VADs. This is also in conformity with other studies which have shown catheter-related infection as the cause of premature removal in 7.1% to 13.4% of cases¹³⁻¹⁵.

Although several studies have reported premature removal due to device failure to occur in 2.9 to 3.4% of VADs¹³⁻¹⁵, a much higher incidence was seen in our patients. Nine percent of the VADs inserted at our hospital had to be removed prematurely because of device failure. This could be attributed to the fact that earlier we did not use anticoagulant therapy in our patients during the time the device was in place, a practice we have now changed. There are some reports that the use of continuous prophylactic anticoagulant therapy reduces the incidence of VAD failure¹⁶⁻¹⁸. However, most reports that we have used for comparison with our data, do not indicate whether anticoagulant therapy was given to patients or not at that time.

In conclusion, our results are largely comparable to the published data. However, there is a need to reduce the catheter related complications. Overall, the use of VADs is safe in developing countries.

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