# Journal Article Summary Worksheet

## **Article Title**

Implanted vascular access device related deep vein thrombosis in oncology patients; A prospective cohort study. (Aug 2015 – Sept. 2017, Data collected at Ottawa Hospital)

#### Name of Journal /Date

Thrombosis Research / February 2019 Authors: Suleman A, Jarvis V, Hadziomerovic A, Carrier M, McDiarmid S

### **Study Objectives**

Assess the incidence of Implanted Vascular Access Device (IVAD)-related upper extremity deep venin thrombosis (UEDVT) associated with the AngioDynamics BioFlo Port.

Study Design/Methods		
Number of Patients	501 cancer patients were assessed for eligibility – (501 BioFlo ports placed)	
Number of Patients	394 of those patients enrolled in the study	
	389 of those patients were enrolled over 12 months & included in the final analysis	
	*breakdown why certain patients were excluded and removed in Fig 1	
	Prospective single center cohort study – study followed one group of patients after	
Patient Groups	the port was implanted for a timeframe of 12 months, or until a patient received an	
	anticoagulant, the catheter was removed, or death.	
	Incidence of IVAD-related UEDVT associated with BioFlo ports where IVAD-related	
Primary Endpoints	UEDVT was defined as symptomatic ipsilateral upper extremity (axillary vein or	
	proximal) DVT and symptomatic Pulmonary Embolism (PE)	
	Retrospective studies, assessing the incidence of IVAD-related UEDVT associated	
Secondary Endpoints	with other ports – specifically the Bard X-port ISP.	

### **Study Results**

- Of the 389 patients included in the analysis, it was determined that only 5 patients (1.29%) had symptomatic port-related UEDVT (1.29%, 95% CI 0.2 50 2.4%)
- In a previous study at the same institution with similar sample size and patient population, the port-associated DVT rate was 4.5% (X-port ISP, Bard Access SystemsInc, Salt Lake City, US). This represents a 71% reduction of port-associated DVT [1-(1.29/4.5) = 71%]
- The median age of the cohort was 58.2 years; 68% (n=273) were females. Sixty-six percent had gastrointestionalcancer (including pancreatic cancer) and 68% had metastases. Eighty four percent of IVADs were right sided insertions. Ninety eight percent of catheter tip placements were distal superior vena cava (n=237), cavo-atrial junction (n=67) or atrium (n=90)

### **Study Conclusions**

- IVAD-related UEDVT is an infrequent complication in cancer patients with BioFlo ports
- Specifically, using a BioFlo port can reduce port-associated UEDVT by 71%
- The risk of thrombotic catheter complications that are associated with increased morbidity, mortality, healthcare costs and diminished quality of life may be reduced by the use of the BioFlo\*port



### **Points/Key Take-Aways**

- This study suggests that cancer patients who receive an implanted BioFlo port have a reduced chance of getting an upper extremity DVT
  - We can make this determination because the rate of symptomatic IVAD-related UEDVTs reported in this study is lower than other previously reported rates.
  - While this study reported rates at 1.29%, other studies retrospectively reported rates of VTE complications ranging between 3.8% – 5.5%, with 4-10% being the estimated industry standard.
  - And while the difference between 1.29% and 4.5% may not be initially overwhelming, the difference is important to note when considering patient outcomes.

Especially when VTE is the second leading cause of death in cancer patient

- AngioDynamics BioFlo port reported a lower rate of IVAD-related UEDVT when compared to the Bard X-port ISP. The Bard port reported a 4.5% incidence rate of UEDVT in a similar cohort study (article 10 in references).
  - Incidence and risk factors of symptomatic venous thromboembolism related to implanted ports in cancer patients retrospectively assessed the incidence of IVAD-related UEDVTs associated with the Bard X-port ISP (4.5%, 95% CI, 2.5 to 6.3%)

Possible Objections		Response
1.	Because cancer patients are at higher risk of VTE complications, it is challenging to determine if the VTE is related to the port or not. The study did not include a control group comparing ports. Therefore, it is difficult to determine whether the low risk of IVAD-related DTVs is attributed to the BioFlo port itself, or the highly trained specialized team that implanted and cared for the port.	<ul> <li>aways:</li> <li>While that point is valid, I feel we cannot ignore the difference in reported symptomatic IVAD-related UEDVTs when retrospectively compared to previous studies.</li> </ul>

### In What Sales Scenarios Would You Use this Study?

- A) Selling BioFlo ports
- B) Selling BioFlo ports against Bard X-port ISP
- C) Upselling existing business to BioFlo Ex: Xcela Plus to BioFlo port

