romanian NEUROSURGERY

Vol. XXXVI | No. 4 December 2022

Ventriculoperitoneal shunt surgery in a Nigerian city. A single institutional experience

Ukpong Secundus Udoffa, Stephen Agbomhekhe Oga, Folafoluwa Ayokunle Aduloju, Okezie Obasi Kanu



Ventriculoperitoneal shunt surgery in a Nigerian city. A single institutional experience

Ukpong Secundus Udoffa¹, Stephen Agbomhekhe Oga², Folafoluwa Ayokunle Aduloju¹, Okezie Obasi Kanu³

¹ Department of Surgery Federal Medical Centre Lokoja, Niger State, NIGERIA

² Mount Gerizim Medical Centre, Lokoja NIGERIA

³ Neurosurgery Division, Department of Surgery, College of Medicine, University of Lagos, Idi-Araba, Lagos State, NIGERIA

Keywords

ventriculoperitoneal shunt, hydrocephalus, paediatric hydrocephalus

ABSTRACT

Introduction. Improvements in surgical techniques and advancements in antibiotic management have significantly reduced the complications associated with Ventriculoperitoneal Shunting which is still the most common procedure for the treatment of hydrocephalus. These complications are believed to be highest in Low and middle-income countries due to delayed presentation to care facilities. The authors report our experience of managing paediatric hydrocephalus in an institution with evolving Neurosurgical practice in North Central Nigeria.

Material and methods. We retrospectively reviewed all cases of VP shunting from 2011 to 2018 taking into consideration the demographics, aetiologies of hydrocephalus, length of hospital stay, postoperative morbidity and mortality, and overall outcome. Complications sought included, Surgical Site infections, shunt exposure, obstruction or any other cause of shunt malfunction. The minimum follow-up period was 24 months

Results. There were 27 VPS procedures done in 25 patients of whom 15 were males and 10 females (M:F ratio of 1.5:1). The age range was 8 days to 9 years with a median age of 5 months. Of the 25 cases, 24 (96%) were non-tumoral in origin comprising 17 congenital and 7 acquired hydrocephalus. At 2 years post shunt insertion, 21 (84%) of the 25 initial cases were still functional. The total complication rate was 28%, comprising Surgical Site infection, shunt exposure, shunt obstruction, seizure, and one death, There were 3 (12%) shunt failures from shunt obstruction (2) and shunt exposure (1).

Conclusion

With meticulous control of the surgical environment and improved experience in ventriculoperitoneal shunting, the complication rate can be significantly reduced. There is a need to increase the awareness of the population to the availability of care for seemingly hopeless conditions. The burden of the cost of care on individuals should be lightened through better health insurance coverage.

 \succ

Corresponding author: Ukpong Udoffa

Department of Surgery, Federal Medical Centre, Lokoja Kogi, State, Nigeria

ukpongudoffa1811@gmail.com

Copyright and usage. This is an Open Access article, distributed under the terms of the Creative Commons Attribution Non-Commercial No Derivatives License (https://creativecommons .org/licenses/by-nc-nd/4.0/) which permits noncommercial re-use, distribution, and reproduction in any medium, provided the original work is unaltered and is properly cited. The written permission of the Romanian Society of Neurosurgery must be obtained for commercial re-use or in order to create a derivative work.

> ISSN online 2344-4959 © Romanian Society of Neurosurgery



First published December 2022 by London Academic Publishing www.lapub.co.uk

INTRODUCTION

Hydrocephalus is one of the most common clinical conditions affecting children in Neurological surgery. Ventriculoperitoneal (VP) shunt placement is the mainstay of treatment for hydrocephalus in both adult and paediatric patients.(1, 2) Hydrocephalus accounts for over 69,000 hospital admissions and over 30,000 procedures performed every year in the United States.(3-7) with failure rates of 30-40% at 1 year and approximately 50% at 2 years in pediatric patients.(8) Some recent studies report a relatively lower rate of shunt failure.(1, 9, 10) It is believed that VP shunting complications are highest in Africa and other developing countries.(11) This is not unconnected with delayed presentation to care facilities as a result of poverty, ignorance and other socio-cultural factors.

Hereby, we report our experience of managing pediatric hydrocephalus in an institution with evolving Neurosurgical practice in North Central Nigeria.

MATERIALS AND METHOD

The authors present a three-year-old female patient who accidentally sustained a PHI with a screwdriver through the right orbit (Figure 1). The patient did not lose consciousness or vomit after the injury. She was previously examined by an ophthalmologist and a pediatric surgeon at the local hospital. During the neurosurgical examination, on admission, the patient was conscious, oriented to time, space and person (Glasgow Coma Scale score of 15), cardiopulmonary stable, without any recorded gross neurological deficits.

RESULTS

Demographics

There were 27 VPS procedures done in 25 patients of whom 15 were males and 10 females with a male-female ratio of 1.5:1. The age range was 8 days to 9 years with a mean age of 1.2 years and a median of 5 months. All patients had clinical features of hydrocephalus and confirmation was made with diagnostic Scan. The Occipito-Frontal circumference ranged betwwn 39cm to 65cm. All patients were treated with CChabra Slit and Spring Shunt. Length of post-operative hospital stay was 3 days to 16 days with a mean duration of 9.2 days.

Aetiology of hydrocephalus

Of the 25 cases of hydrocephalus, 24 (96%) were non-tumoral in origin with only one case (4%) caused by posterior fossa tumour in a 9 year-old child, who coincidentally is the oldest patient in this series.

Seventeen (68%) of the 25 cases were congenital in origin, comprising of Aqueductal stenosis (11), Arnold-Chiari Malformation (4) and Dandy-Walker malformation (2). The major cause of Acquired hydrocephalus was post-meningitic (5 of 8). There were 2 cases of Aqueductal stenosis with no history of infection. (Table 1b)

Table 1. Demographic profile of hydrocephalus

Variables	Frequency (%)	
A)		
Males		15 (60)
Females		10 (40)
B)		
Congenital		17 (68)
a)	Acqueductal stenosis	11 (44)
b)	Arnorld-Chiari Malformation	4 (16)
c)	Dandy-Walker malformation	2 (8)
Acquire	8 (32)	
a)	Post-infective	5 (20)
b)	Acqueductal Stenosis	2 (8)
c)	Post fossa Tumor	1 (4)
C)	Diagnostic Imaging	
MRI		6 (24)
CT Scan		17 (68)
TFUS		2 (8)

Diagnostic Imaging

Diagnosis was made in 17 cases (68%) with Computerized Tomographic (CT) scan. Magnetic Resonance Imaging (MRI) was deployed in 6 (24%) while Trans-fontanel Ultrasonographic Scan (TFUSS) was used in 2 (8%) patients (Table 1c)

Outcome

At 2-year post shunt insertion, 21 (84%) of the 25 initial cases were still functional. There were 3 (12%) shunt failures from shunt obstruction (2) and shunt exposure (1). Two of the shunt failures were revised. There was one post-operative death (4%).

There were 3 (12%) cases of surgical site infection (SSI) at the scalp region noted in the first week postoperative period, all of which grew Staphylococcus aureus that was treated with Amikacin and Rifampicin. One of the cases of SSI led to shunt exposure which was promptly removed but the parents declined revision surgery and requested discharge from the facility. The child was lost to follow-up. One patient (4%) died on the third postoperative day accounting for the shortest period of hospital stay. Permission for autopsy was not granted by the parents. One patient developed seizure disorder post op.

Whereas shunt complication was seen in 28% of all patients, only 12% required shunt revision (Table 2). The other 2 cases of shunt failure were marked by increasing head circumference and reduced activities. These occurred within 6 months of the surgery. Shunt was promptly revised and the cause of failure was determined in both cases to be obstruction of the ventricular catheter by debris. These two patients have remained clinically stable.

Table 2. Outcome and complications of VentriculoperitonealShunting

a)	Summary of	Frequency	Comments
Outcome (n=25)		(%)	
Fav	ourable outcome	21 (84)	satisfactory
Poor outcome		4 (16)	3 failed, 1
			died
a.	Complications (7	Frequency	Comments
	of 25)	(%)	
Surgical Site Infection		3 (12)	Antibiotics
(SSI)*		1 (4)	Declined
Shunt Exposure*		2 (8)	Treatment
Shunt Obstruction		1 (4)	Revised
Seizure		1 (4)	Anticonvulsant
Death			Post-op day 3

(*one case of SSI led to shunt exposure)

DISCUSSION

Ventriculoperitoneal shunting remains the mainstay for treatment of hydrocephalus despite recent advances in neurological surgery practice. (1-3)

In this study the age range was between 8 days to 9years with a median of 5 months. Ninety two percent of the patients were below the age of 1 year and it is mainly caused by congenital anomalies. Hydrocephalus is predominantly a disease of infants and this is a common finding in Sub-Saharan Africa. (6, 11-15)

The male to female ratio of 1.5:1 is in keeping with the male preponderance noted in many studies. (6, 12-14, 16) The occiptofrontal circumference (OFC) ranged from 39cm to 65cm. Increase in the OFC is the commonest sign of hydrocephalus seen in infants.(17) Grotesque head enlargement is common in underdeveloped countries due to late presentation and head circumference greater than 60cm is associated with higher rates of shunt failure.(11)

About 70% of the cases of hydrocephalus were congenital with Acqueductal stenosis being the commonest cause. (Table 1b) This is at variance with some of the studies cited (11, 12, 14-16, 18) and the reason may be partly related to volume of the study population. The other reasons may be geographical in nature. Many mothers who had infection during pregnancy in rural areas may not have been adequately looked after, thereby increasing the risk of maternal-to-fetal transmission. Neural tube defects (Arnold-Chiari Malformations with Spina bifida, and congenital posterior fossa anomalies) are still a challenge in low-income communities where adherence to Folic acid supplementation is suboptimal.(15, 19)

The preferred imaging modality is MRI but this was used in only 24% of patients (Table 1c). CT scan was the most commonly used imaging modality in this study because of affordability. Transfontanel Ultrasonography has been used very frequently is our region for the similar reasons.(20, 21)

The overall complication rate in this series was 28%. (Table 2) The commonly reported incidence of complication is between 20 to 40%(10) though there are reported failures as high as 85%.(6) The incidence has reduced in more recent publications. The infection rate was 12% in this study. This is similar to the rate recorded by Yusuf et al(13) in an earlier study though they had a relatively higher volume in their series. The weighted average shunt infection rate across multiple studies is about 5.1% but could be as high as 39% in some studies.(18, 22-28) Staphylococcus aureus is one of the most implicated bacterial organisms in shunt infections.(29) Shunt infection has been reduced with advent of newer techniques including double gloving, prophylactic antibiotics and antibiotic-impregnated shunts (commonly with Rifampicin and Clindamycin)(30) Most of the shunts used in developing countries are antibioticfixed-pressure types of shunt; impregnated shunts are not affordable to most patients in underdeveloped countries where the citizens are not adequately covered by health insurance and health care is paid on out-of-pocket basis.(20, 21) Other established factors that influence the shunt infection rate are the age of the patients, aetiology of the hydrocephalus, operating room settings to prevent infection, total operating time and experience of the surgeon.(26) Sharing the experience of the senior author in center with large volume has helped to cut down shunt failure rates in the country.

There were 3 (12%) failed shunts though 2 (8%) shunt revision surgeries were done in this study. One was caused by infection, while the other two were caused by shunt obstruction. The incidence of shunt failure is commonly seen in children younger than 6 months and often noticed within the first month of shunt placement.(18, 31)

Seizure was seen in one patient in our series, accounting for 4% and this was controlled with anticonvulsant. Seizure is a known complication of VP shunting accounting for 48%. It is believed that seizure is not due to direct placement of the VP shunt but to the underlying neurologic disorder.(32)

One patient (4%) died in the first week following VP shunt placement. The cause of death could not be determined because the parents declined postmortem. The shunt-related mortality has been reported to be 3.4% to 13.7%.(13, 33).

Though Endoscopic third Ventriculostomy is available in the country as shown in many studies, (13, 14, 16) this facility is not available in our center at the time of this study. It is expected that the face of hydrocephalus treatment will improve as soon as this is done.

CONCLUSION

With meticulous control of the surgical environment and improved experience in ventriculoperitoneal shunting, the complication rate can be significantly reduced. There is need to increase the awareness of the population to the availability of care for seeming hopeless conditions. The burden of the cost of care on individuals should be lightened through better health insurance coverage.

REFERENCES

 Khan F, Shamim MS, Rehman A, Bari ME. Analysis of factors affecting ventriculoperitoneal shunt survival in pediatric patients. Childs Nerv Syst. 29(5):791-802,2013.

- 2. Khan F, Rehman A, Shamim MS, Bari ME. Factors affecting ventriculoperitoneal shunt survival in adult patients. Surg Neurol Int. 6:25,2015.
- Bondurant CP, Jimenez DF. Epidemiology of cerebrospinal fluid shunting. Pediatr Neurosurg. 23(5):254-8; discussion 9,1995.
- 4. Merkler AE, Ch'ang J, Parker WE, Murthy SB, Kamel H. The Rate of Complications after Ventriculoperitoneal Shunt Surgery. World Neurosurg. 98:654-8,2017.
- Borgbjerg BM, Gjerris F, Albeck MJ, Borgesen SE. Risk of infection after cerebrospinal fluid shunt: an analysis of 884 first-time shunts. Acta Neurochir (Wien). 136(1-2):1-7,1995.
- Stone JJ, Walker CT, Jacobson M, Phillips V, Silberstein HJ. Revision rate of pediatric ventriculoperitoneal shunts after 15 years. J Neurosurg Pediatr. 11(1):15-9,2013.
- Simon TD, Riva-Cambrin J, Srivastava R, Bratton SL, Dean JM, Kestle JR, et al. Hospital care for children with hydrocephalus in the United States: utilization, charges, comorbidities, and deaths. J Neurosurg Pediatr. 1(2):131-7,2008.
- Hanak BW, Bonow RH, Harris CA, Browd SR. Cerebrospinal Fluid Shunting Complications in Children. Pediatr Neurosurg. 52(6):381-400,2017.
- Khan F, Rehman A, Shamim MS, Bari ME. Ventriculoperitoneal (VP) Shunt Survival in Patients Developing Hydrocephalus After Cranial Surgery. Turk Neurosurg. 26(3):369-77,2016.
- Reddy GK, Bollam P, Caldito G. Long-term outcomes of ventriculoperitoneal shunt surgery in patients with hydrocephalus. World Neurosurg. 81(2):404-10,2014.
- 11. Gathura E, Poenaru D, Bransford R, Albright AL. Outcomes of ventriculoperitoneal shunt insertion in Sub-Saharan Africa. J Neurosurg Pediatr. 6(4):329-35,2010.
- Komolafe EO, Adeolu AA, Komolafe MA. Treatment of cerebrospinal fluid shunting complications in a Nigerian neurosurgery programme. Case illustrations and review. Pediatr Neurosurg. 44(1):36-42,2008.
- Yusuf AS, Omokanye HK, Adeleke NA, Akanbi RO, Ajiboye SO, Ibrahim HG. Management and Outcome of Infantile Hydrocephalus in a Tertiary Health Institution in Nigeria. J Neurosci Rural Pract. 8(2):249-53,2017.
- Bankole OB, Ojo OA, Nnadi MN, Kanu OO, Olatosi JO. Early outcome of combined endoscopic third ventriculostomy and choroid plexus cauterization in childhood hydrocephalus. J Neurosurg Pediatr. 15(5):524-8,2015.
- Adebayo BO, Kanu OO, Bankole OB, Ojo OA, Adetunmbi B, Morgan E. Early Outcome of Endoscopic Third Ventriculostomy With Choroid Plexus Cauterization Versus Ventriculoperitoneal Shunt as Primary Treatment of Hydrocephalus in Children With Myelomeningocele: A Prospective Cohort Study. Oper Neurosurg (Hagerstown). 21(6):461-6,2021.
- Ojo OA, Bankole OB, Kanu OO, Okubadejo NU. Efficacy of endoscopic third ventriculostomy in the management of hydrocephalus in children under 2 years of age:

experience from a tertiary institution in Nigeria. Niger J Clin Pract. 18(3):318-22,2015.

- Pan P. Outcome Analysis of Ventriculoperitoneal Shunt Surgery in Pediatric Hydrocephalus. J Pediatr Neurosci. 13(2):176-81,2018.
- Kulkarni A, Drake J, Lamberti-Pasculli M. Cerebrospinal fluid shunt infection: a prospective study of risk factors. J Neurosurg. 94(2):195-201,2001.
- 19. Bankole OB, Arigbabu SO, Kanu OO. Spinal neural tube defects in Lagos University Teaching Hospital, Nigeria. Nig Q J Hosp Med. 22(1):22-4,2012.
- Kanu OO, Nnoli C, Olowoyeye O, Ojo O, Esezobor C, Adeyomoye A, et al. Infantile subdural empyema: The role of brain sonography and percutaneous subdural tapping in a resource-challenged region. J Neurosci Rural Pract. 5(4):355-9,2014.
- Kanu OO, Esezobor CI, Ojo OA, Asoegwu CN, Nnoli C, Dawang Y, et al. Infantile supratentorial subdural empyema managed by percutaneous aspiration: an outcome study in a Nigerian city. Sudan J Paediatr. 19(1):37-43,2019.
- Raygor KP, Oh T, Hwang JY, Phelps RRL, Ghoussaini K, Wong P, et al. Ventriculoperitoneal shunt infection rates using a standard surgical technique, including topical and intraventricular vancomycin: the Children's Hospital Oakland experience. J Neurosurg Pediatr.1-9,2020.
- Choux M, Genitori L, Lang D, Lena G. Shunt implantation: reducing the incidence of shunt infection. J Neurosurg. 77(6):875-80,1992.
- 24. Drake J, Kestle J, Milner R. Randomized trial of cerebrospinal fluid shunt valve design in pediatric hydrocephalus. Neurosurgery. 43(2):294-305,1998.

- 25. Enger P, Svendsen F, Wester K. CSF shunt infections in children: experiences from a population-based study. Acta Neurochir (Wien). 145(4):243-8,2003.
- 26. Reddy G, Bollam P, Caldito G. Ventriculoperitoneal shunt surgery and the risk of shunt infection in patients with hydrocephalus: long-term single institution experience. World Neurosurg. 78(1-2):155-63,2012.
- Rehman A-U, Rehman T-U, Bashir H, Gupta V. A simple method to reduce infection of ventriculoperitoneal shunts. J Neurosurg Pediatr. 5(6):569-72,2010.
- Edwards N, Engelhart L, Casamento E, McGirt M. Costconsequence analysis of antibiotic-impregnated shunts and external ventricular drains in hydrocephalus. J Neurosurg. 122(1):139-47,2015.
- Wu X, Liu Q, Jiang X, Zhang T. Prevention options for ventriculoperitoneal shunt infections: a retrospective analysis during a five-year period. Int J Clin Exp Med. 8(10):19775-80,2015.
- Brown EM, Edwards RJ, Pople IK. Conservative management of patients with cerebrospinal fluid shunt infections. Neurosurgery. 58(4):657-65; discussion -65,2006.
- 31. Faillace WJ. Shunt infection. J Neurosurg. 94(6):1019-20,2001.
- Klepper J, Busse M, Strassburg HM, Sorensen N. Epilepsy in shunt-treated hydrocephalus. Dev Med Child Neurol. 40(11):731-6,1998.
- Tuli S, Tuli J, Drake J, Spears J. Predictors of death in pediatric patients requiring cerebrospinal fluid shunts. J Neurosurg. 100(5 Suppl Pediatrics):442-6,2004.