

See the corresponding editorial in this issue, pp 307–309.

J Neurosurg Pediatrics 6:310–315, 2010

See the corresponding erratum notice, DOI: 10.3171/2010.12.PEDS103a, for full details.

Predicting who will benefit from endoscopic third ventriculostomy compared with shunt insertion in childhood hydrocephalus using the ETV Success Score

Clinical article

ABHAYA V. KULKARNI, M.D., PH.D.,¹ JAMES M. DRAKE, F.R.C.S.C.,¹
JOHN R. W. KESTLE, M.D.,² CONOR L. MALLUCCI, F.R.C.S.(SN),³
SPYROS SGOUROS, M.D., F.R.C.S.(SN),⁴ SHLOMI CONSTANTINI, M.D., M.Sc.,⁵
AND THE CANADIAN PEDIATRIC NEUROSURGERY STUDY GROUP

¹Hospital for Sick Children, Toronto, Canada; ²Primary Children's Medical Center, Salt Lake City, Utah; ³Royal Liverpool Children's Hospital, Liverpool; ⁴Birmingham Children's Hospital, Birmingham, United Kingdom; and ⁵Dana Children's Hospital, Tel Aviv Medical Center, Tel Aviv, Israel

Object. The authors recently developed and internally validated the ETV Success Score (ETVSS)—a simplified means of predicting the 6-month success rate of endoscopic third ventriculostomy (ETV) for a child with hydrocephalus, based on age, etiology of hydrocephalus, and presence of a previous shunt. A high ETVSS predicts a high chance of early ETV success. In this paper, they assess the clinical utility of the ETVSS by determining whether long-term survival outcomes for ETV versus shunt insertion are different within strata of ETVSS (low, moderate, and high scores).

Methods. A multicenter, international cohort of children (≤ 19 years old) with newly diagnosed hydrocephalus treated with either ETV (489 patients) or shunt insertion (720 patients) was analyzed. The ETVSS was calculated for all patients. Survival analyses with time-dependent modeling of the hazard ratios were performed.

Results. For the High-ETVSS Group (255 ETV-treated patients, 117 shunt-treated patients), ETV appeared to have a lower risk of failure right from the early postoperative phase and became more favorable with time. For the Moderate-ETVSS Group (172 ETV-treated patients, 245 shunt-treated patients), ETV appeared to have a higher initial failure rate, but after about 3 months the instantaneous risk of ETV failure became slightly lower than shunt failure (that is, the hazard ratio became < 1). For the Low-ETVSS Group (62 ETV-treated patients, 358 shunt-treated patients), the early risk of ETV failure was much higher than the risk of shunt failure, but the instantaneous risk of ETV failure became lower than the risk of shunt failure at about 6 months following surgery (the hazard ratio became < 1).

Conclusions. Across all ETVSS strata, the risk of ETV failure becomes progressively lower compared with the risk of shunt failure with increasing time from the surgery. In the best ETV candidates (ETVSS ≥ 80), however, the risk of ETV failure is lower than the risk of shunt failure very soon after surgery, while for less-than-ideal ETV candidates (ETVSS ≤ 70), the risk of ETV failure is initially higher than the risk of shunt failure and only becomes lower after 3–6 months from surgery. These results need to be confirmed by larger, prospective, and preferably randomized studies. (DOI: 10.3171/2010.8.PEDS103)

KEY WORDS • endoscopy • hydrocephalus • pediatric neurosurgery • endoscopic third ventriculostomy • shunt

THE debate over ETV versus CSF shunting for the treatment of childhood hydrocephalus remains largely unresolved.^{1,2,4,6,7,9,11,16,18,20} Recently, we used confounder-adjustment techniques (propensity score modeling) to compare the failure rate of ETV versus shunt placement.¹³ We found that, once adjusted for confounders, ETV

is associated with a higher initial failure rate, but, over time, the failure rate becomes lower than that associated with shunt placement. In a separate publication, we used logistic regression techniques to develop and internally validate a prediction score that would, with good accuracy, predict the 6 month success rate of ETV, taking into account the patient's age and the etiology of their hydrocephalus.¹⁴ The end product of this was the development of the ETV Success Score (ETVSS) (Table 1)—a simple means to predict

Abbreviations used in this paper: ETV = endoscopic third ventriculostomy; ETVSS = ETV Success Score.

Endoscopic third ventriculostomy versus shunt insertion

TABLE 1: Calculation of the ETVSS*

Score	Age	Etiology	Previous Shunt
0	<1 mo	postinfectious	previous shunt
10	1 mo to <6 mos		no previous shunt
20		myelomeningocele, IVH, nontectal brain tumor	
30	6 mos to <1 yr	aqueductal stenosis, tectal tumor, other	
40	1 yr to <10 yrs		
50	≥10 yrs		

* The ETVSS is calculated as Age Score + Etiology Score + Previous Shunt Score. Abbreviation: IVH = intraventricular hemorrhage.

ETV success with scores ranging from 0 (extremely poor chance of ETV success) to 90 (extremely high chance of ETV success). We suggested that the ETVSS could be used to meaningfully select good candidates for ETV.

There were 2 main limitations to our previous work, however. First, it left unanswered the question: how would patients with equivalent ETVSSs fare with a shunt instead of an ETV? The ETVSS predicts success based on age and etiology, both of which likely have prognostic significance for shunt-treated patients also. Therefore, it is possible that the ETVSS may simply select patients who will do very well with either ETV or a shunt, without a clear relative benefit of one over the other. Second, the ETVSS was designed to predict relatively short-term outcome only—that is, the chance of being failure-free at 6 months. How would these patients do after a longer period of follow-up, and would the ETVSS still be predictive of good outcome? The current analysis is our attempt to address these issues. In this report, we tested whether the success rate predicted by the ETVSS would specifically translate into different long-term outcomes for ETV compared with shunt placement.

Methods

Patient Population

The cohort for this study was accrued from 3 main sources and has been described previously.^{5,10,13,14} All patients were 19 years old or younger; had newly diagnosed, previously untreated, high-pressure hydrocephalus; and were treated by pediatric neurosurgeons at specialized centers. The ETV cohort was collected from 12 centers in Canada, Israel, and the United Kingdom, and the patients were treated between 1989 and 2006.¹⁴ The shunt cohort was collected from the Shunt Design Trial⁵ (patient recruitment 1993–1995 from Canada, France, the Netherlands, and the US) and the Endoscopic Shunt Insertion Trial¹⁰ (patient recruitment 1996–1999 from Canada, the Netherlands, the United Kingdom, and the US). Permission was granted from each trial’s principal investigators for analysis of data. Since these trials demonstrated no difference in outcome in the treatment arms, data from all patients were analyzed collectively for this study. All data were anonymized and data collection adhered to local research ethics protocols.

We defined the failure of treatment as any subsequent surgical procedure for definitive CSF diversion (either shunting or ETV) or death related to hydrocephalus management.

Statistical Analysis

For each patient in the cohort we calculated the ETVSS (Table 1), regardless of whether the patient was actually treated by ETV or shunt insertion. The ETVSS is based on the patient’s age, hydrocephalus etiology, and presence of a previous shunt. Since all patients in our cohort had newly diagnosed hydrocephalus, no patient had a previous shunt. The ETVSS ranges from 0 to 90, and the number itself roughly approximates the percentage chance that an ETV will be successful at 6 months. For example, an 8-month-old with aqueductal stenosis and no previous shunt would have an ETVSS of (30 + 30 + 10) = 70, or a roughly 70% chance of having a successful ETV without failure at 6 months postprocedure. We have previously demonstrated the internal validity of the ETVSS.¹⁴

We stratified the sample into patients with a high ETVSS (≥ 80), those with a moderate ETVSS (50–70), and those with a low ETVSS (≤ 40). Within each of these stratified groups we performed comparative survival analyses, using the time to treatment failure as the outcome of interest. We used Kaplan-Meier methods to construct survival curves. We performed survival analyses using Cox proportional hazards models to compare ETV failure relative to shunt failure. This provides a hazard ratio, which is the instantaneous risk of ETV failure compared with shunt failure, at a given moment in time following surgery (values greater than 1 indicate a higher chance of ETV failure than shunt failure). We discovered, however, that the assumption of proportional hazards was violated, based on visual inspection of the treatment survival curves (the curves crossed each other) and significance of the interaction of time and treatment ($p < 0.001$). This meant, therefore, that the risk of ETV failure compared with shunt failure was not constant after surgery. Rather, it changed as a function of time after surgery. Because of this, we used a Cox regression model that would allow the hazard ratio to change over time using a log₁₀(time)-by-treatment interaction. This time-dependent interaction term was, therefore, included in all Cox regression models. All analyses were performed with SPSS Advanced Statistics 17.0 (SPSS Inc.).

Results

The characteristics of the 1209 patients are listed in Table 2 and the distribution of the ETVSS for shunt-treated and ETV-treated patients is shown in Fig. 1. As expected, there was a discrepancy in this distribution, with more ETV patients having higher ETVSS, reflecting the selection bias of the treating surgeons. Nevertheless, there was still reasonable overlap of ETV- and shunt-treated patients across the full range of ETVSS.

The results of the survival analysis are shown in Fig. 2 and Table 3. Because the hazard ratios change as a function of time after surgery, these are represented by the

TABLE 2: Summary of patients' characteristics*

Variable	Treatment		Overall (1209 patients)
	ETV (489 patients)	Shunt (720 patients)	
age at treatment			
<1 mo	36 (7.4)	234 (32.5)	270 (22.3)
1 to <6 mos	78 (16.0)	240 (33.3)	318 (26.3)
6 to <12 mos	35 (7.2)	83 (11.5)	118 (9.8)
1 to <10 yrs	206 (42.1)	122 (16.9)	328 (27.1)
≥10 yrs	134 (27.4)	41 (5.7)	175 (14.5)
etiology of hydrocephalus			
stenosis of cerebral aqueduct	168 (34.4)	53 (7.4)	221 (18.3)
tectal tumor	61 (12.5)	0	61 (5.0)
other brain tumor	114 (23.3)	48 (6.7)	162 (13.4)
post-IVH	53 (10.8)	161 (22.4)	214 (17.7)
myelomeningocele	10 (2.0)	178 (24.7)	188 (15.6)
other	83 (17.0)	280 (38.9)	363 (30.0)
mean ETVSS (± SD)	71 ± 16	51 ± 19	59 ± 20

* Values represent number of patients (%) unless otherwise indicated.

logarithmic curves in Fig. 3. These curves are presented to 48 months after surgery, after which the number of patients left in the analysis was very small.

For the High-ETVSS Group, ETV appeared to have a lower risk of failure right from the early postoperative phase onwards (Fig. 3). There was a reasonable separation of the survival curves at 3 years that appeared to diverge further with time (Fig. 2A), but the number of shunt-treated patients in this analysis was limited.

For the Moderate-ETVSS Group, ETV appeared to have a higher initial failure rate, but it became slowly and progressively more favorable over time. After about 3 months from surgery, the instantaneous risk of ETV failure became slightly lower than the risk of shunt failure (the point at which the hazard ratio curve in Fig. 3 crosses below 1) and the survival curves crossed at about 30 months (Fig. 2B).

For the Low-ETVSS Group, the early risk of ETV failure was very high, as expected, but among those who survived this early phase, their risk of ETV failure became lower than the risk of shunt failure at about 6 months following surgery. Although the survival curves appeared to cross at about 42 months, the number of ETV patients left in the analysis was too small for strong conclusions (Fig. 2C).

Discussion

Although the role of ETV appears to be expanding, the major unanswered challenge has been to better define the outcome of ETV, especially in comparison with standard shunting. The development of the ETVSS was our attempt to more strictly predict successful outcome from ETV and thereby formally quantify the effect of the known patient prognostic factors of age and etiology.¹⁴ The current analysis suggests that the ETVSS can be useful in guid-

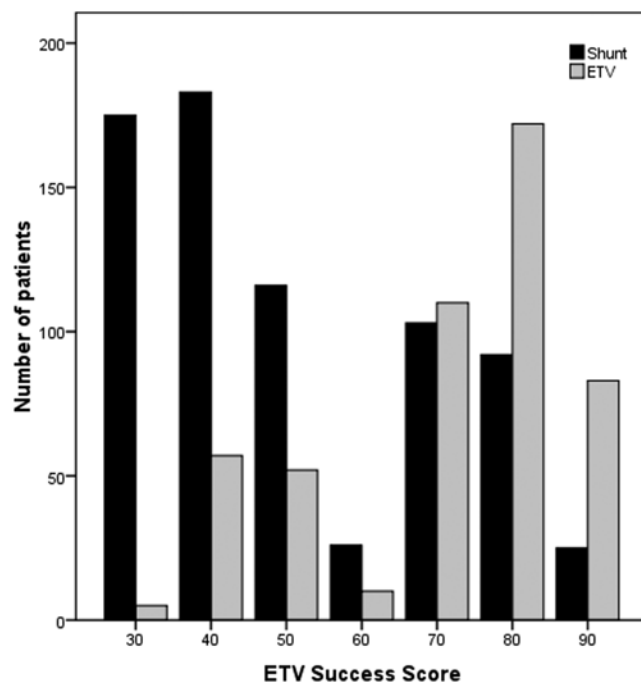


FIG. 1. Frequency histogram showing the distribution of ETVSS among the shunt- and ETV-treated patients.

ing clinical decision making. First, we have shown that the ETVSS predicts not just short-term ETV outcome (as it was designed to do), but also longer-term outcome. The 3-year success rates for the High- and Moderate-ETVSS groups were 72% and 52%, respectively, suggesting that early ETV success leads to a high chance of longer-term success, as well. The corollary of this is that ETV appears to have a low delayed failure rate in comparison with shunt placement, although late ETV failures certainly do occur.³ Interestingly, a lower delayed ETV failure rate compared with shunt placement (that is, a hazard ratio of < 1) was seen across all strata of ETVSS after 6 months following surgery (Fig. 3). Second, we have shown that the success of ETV and CSF shunting are different within strata of ETVSS. In patients with ETVSS ≥ 50 (the High-ETVSS and Moderate-ETVSS groups), there was little difference among the shunt-treated patients, who all had a 3-year survival rate of approximately 50%. In contrast, there is a stark difference in ETV outcome for those with a high ETVSS versus those with a moderate ETVSS: the 3-year success rates were 72% and 52%, respectively. Within the Low-ETVSS Group, shunt insertion appears to have superior success for at least the first 2 years after treatment, although the longer-term failure rate still appears to favor ETV. Therefore, the ETVSS does not simply indiscriminately predict overall hydrocephalus treatment success, but, rather, it specifically differentiates the expected survival outcome of shunt insertion versus ETV.

It is important to recognize the limitations of our analysis. Our data are not randomized nor was outcome blinded. While for the shunt-treated cohort all outcome data were collected prospectively and subject to independent adjudication, this was not the case for the ETV-treated cohort, for whom failure was determined by the treating surgeon. Although our sample is the largest of

Endoscopic third ventriculostomy versus shunt insertion

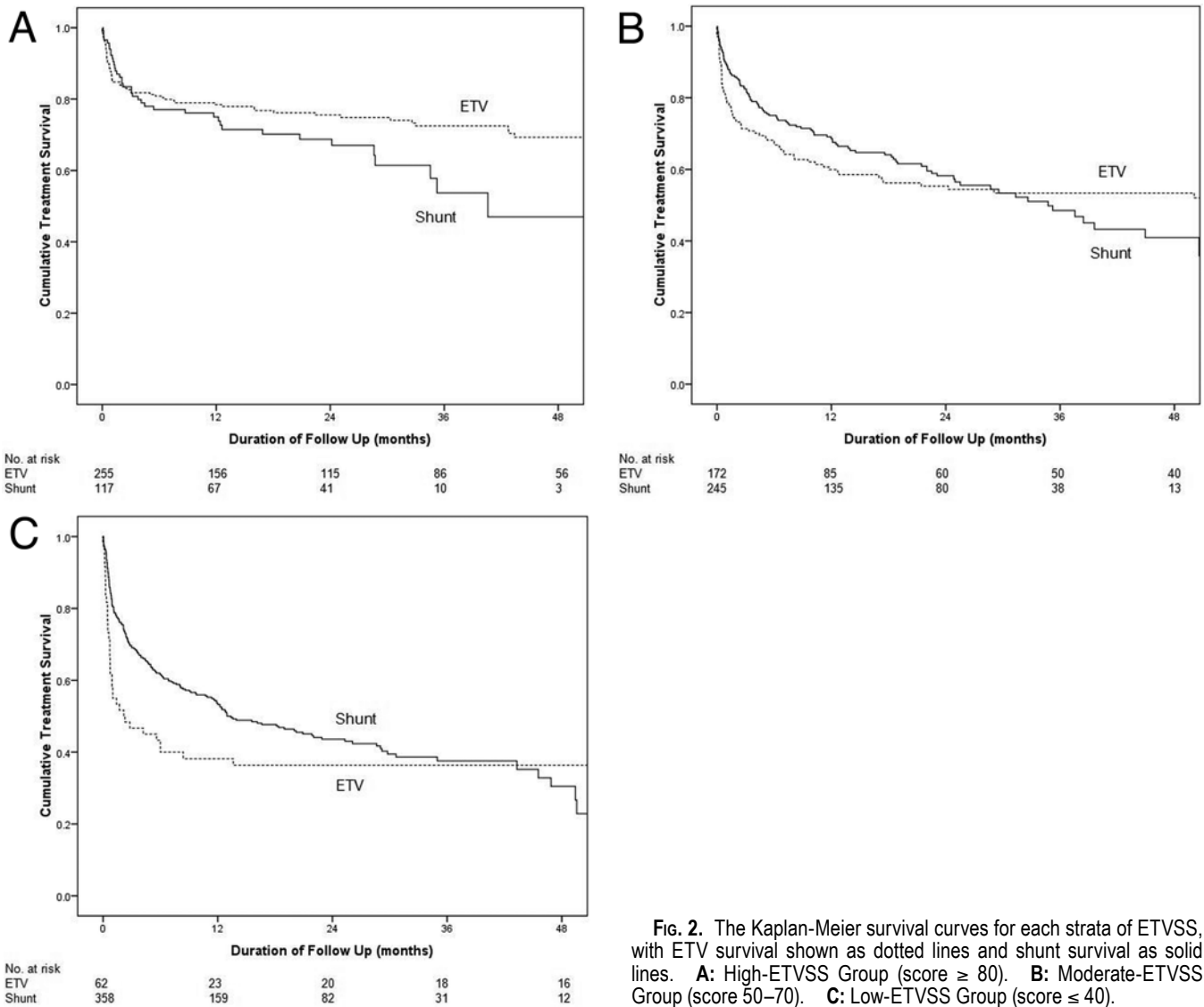


Fig. 2. The Kaplan-Meier survival curves for each strata of ETVSS, with ETV survival shown as dotted lines and shunt survival as solid lines. **A:** High-ETVSS Group (score ≥ 80). **B:** Moderate-ETVSS Group (score 50–70). **C:** Low-ETVSS Group (score ≤ 40).

this nature ever analyzed, our individual stratified analyses suffered from limited numbers of patients, particularly within the High- and Low-ETVSS groups (lacking shunt-treated and ETV-treated patients, respectively). Our use of a logarithmic time-dependent hazard function is complex and does not lend itself to intuitive interpretation. It was important, however, to use this to faithfully

model the true interaction of treatment failure for these 2 interventions. Our data only allowed for meaningful analysis up to 48 months posttreatment. Given the nature of the survival curves, it seems that the true benefit from ETV would be incurred with longer follow-up. Once beyond the early steep failures seen within the first 3–6 months, all indications are that the failure rate for ETV

TABLE 3: Treatment success stratified by ETVSS*

Variable	High ETVSS (≥ 80)	Moderate ETVSS (50–70)	Low ETVSS (≤ 40)
no. of patients			
ETV	255	172	62
shunt	117	245	358
cumulative treatment survival at 36 mos (95% CI)			
ETV	0.72 (0.65–0.78)	0.52 (0.44–0.60)	0.37 (0.24–0.49)
shunt	0.54 (0.39–0.68)	0.49 (0.41–0.57)	0.38 (0.31–0.44)

* Survival figures obtained via Kaplan-Meier survival methods.

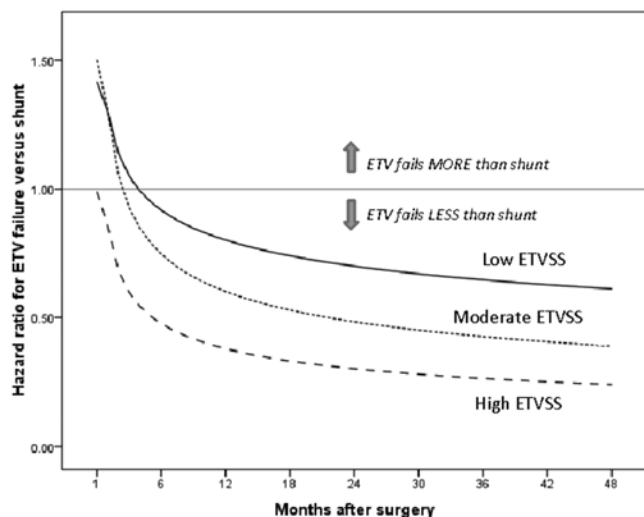


Fig. 3. Line graph demonstrating the treatment failure hazard ratios for ETV compared with shunt insertion as a function of time following surgery. Vertical axis values > 1 indicate that the risk of ETV failure is greater than the risk of shunt failure. Vertical axis values < 1 indicate that the risk of ETV failure is less than the risk of shunt failure. See text for further explanation.

is ultimately lower than that for shunt placement. Therefore, longer-term follow-up might reveal further benefit from ETV that is not evident in our report. Our analysis only examined time to first treatment failure and ignored the many other important aspects of treatment success, including the morbidity associated with the treatment and complications of shunt or ETV failure. As well, we did not examine the effects of repeated failures and other long-term complications, such as slit ventricles. Previous work has suggested that these factors can impact the most important outcome, quality of life.^{12,15} Future studies will need to examine this aspect in further detail. We also recognize that the current version of the ETVSS has not been externally validated and its predictive accuracy might be improved in the future with the inclusion of other predictive features, including preoperative imaging features and intraoperative findings.^{8,19} It is our goal to continually refine the ETVSS as our understanding of ETV success evolves over the coming years.

Given these inherent limitations, our results have to be interpreted cautiously. Specifically, we do not present these as a replacement for randomized data or even contemporaneously collected prospective data. Our analysis is, by no means, definitive. Rather, we feel that this represents, for the time being, one of the most methodologically and statistically rigorous comparisons of ETV versus shunt placement, but much further work is required.

The High-ETVSS Group, with scores ≥ 80 , represents a select group of patients: all are over 1 year of age and the vast majority will have aqueductal stenosis or tectal tumors. These represent the classic indications for ETV, for which there is likely widespread consensus. Ours is one of very few analyses, however, to provide a true direct comparison of this group of patients to similar patients with shunts.¹⁷ An early and lasting survival benefit with ETV is suggested by examination of the survival curves (Fig. 2A) and the hazard ratio curves (Fig. 3). In less-

favorable ETV candidates, however, the survival benefit is either equivocal (for the Moderate-ETVSS Group) or virtually nil (for the Low-ETVSS Group), at least until 4 years after surgery. An optimistic interpretation of the hazard ratio curves in Fig. 3 is that with longer follow-up, ETV might eventually be shown to be beneficial. This would require that a patient survive with the ETV for at least 4 years, after which the risk of long-term failure would be lower than would be seen with a shunt placement. In a sense, this then becomes a question of whether to “front-load” the risk with an ETV (which has a higher early failure rate) for possible long-term benefit or “back-load” the risk with a shunt (which has better early success, but perhaps worse long-term risks). Since individual tolerance for risk varies, sharing this information with families can provide extra guidance for treatment decisions.

Conclusions

We have shown that the ETVSS can be used to differentiate expected survival outcomes for patients treated with ETV compared with CSF shunting. In all ETVSS strata the risk of ETV failure becomes progressively lower compared with the risk of shunt failure with increasing time from the surgery. In the best ETV candidates (those with ETVSS ≥ 80), however, the risk of ETV failure is lower than that of shunt insertion right after surgery, while for less than ideal ETV candidates (ETVSS ≤ 70), the risk of ETV failure is initially higher than that of shunt insertion and only becomes lower after 3–6 months from surgery. Our results will need to be confirmed with further prospective and, preferably, randomized studies.

Disclosure

The authors report no conflict of interest concerning the materials or methods used in this study or the findings specified in this paper. No external funds were received for this study.

Author contributions to the study and manuscript preparation include the following. Conception and design: all authors. Acquisition of data: all authors. Analysis and interpretation of data: Kulkarni, Kestle. Drafting the article: Kulkarni. Critically revising the article: all authors. Reviewed final version of the manuscript and approved it for submission: all authors. Statistical analysis: Kulkarni. Study supervision: Kulkarni.

Appendix 1: Contributing members of the Canadian Pediatric Neurosurgery Study Group

Alberta Children’s Hospital, Calgary, Canada: W Hader; M Hamilton
 Children’s & Women’s Health Centre of BC, Vancouver, Canada: DD Cochrane, P Steinbok
 Children’s Hospital of Eastern Ontario, Ottawa, Canada: M Vassilyadi, E Ventureyra
 Hospital for Sick Children, Toronto, Canada: PB Dirks, JM Drake, AV Kulkarni, JT Rutka, A Van der Stoel, I Veltman
 IWK Health Centre, Halifax, Canada: W Howes, PD McNeely, SA Walling
 London Health Sciences Centre, London, Canada: A Ranger
 Montreal Children’s Hospital, Montreal, Canada: J Atkinson, JP Farmer, J Montes
 Stollery Children’s Hospital, Edmonton, Canada: K Aronyk, V Mehta
 Winnipeg Children’s Hospital, Winnipeg, Canada: PJ McDonald

References

1. Balthasar AJ, Kort H, Cornips EM, Beuls EA, Weber JW, Vles JS: Analysis of the success and failure of endoscopic third ventriculostomy in infants less than 1 year of age. **Childs Nerv Syst** **23**:151–155, 2007
2. Di Rocco C, Massimi L, Tamburrini G: Shunts vs endoscopic third ventriculostomy in infants: are there different types and/or rates of complications? A review. **Childs Nerv Syst** **22**: 1573–1589, 2006
3. Drake J, Chumas P, Kestle J, Pierre-Kahn A, Vinchon M, Brown J, et al: Late rapid deterioration after endoscopic third ventriculostomy: additional cases and review of the literature. **J Neurosurg** **105** (2 Suppl):118–126, 2006
4. Drake JM: Endoscopic third ventriculostomy in pediatric patients: the Canadian experience. **Neurosurgery** **60**:881–886, 2007
5. Drake JM, Kestle JR, Milner R, Cinalli G, Boop F, Piatt J Jr, et al: Randomized trial of cerebrospinal fluid shunt valve design in pediatric hydrocephalus. **Neurosurgery** **43**:294–305, 1998
6. Etus V, Ceylan S: Success of endoscopic third ventriculostomy in children less than 2 years of age. **Neurosurg Rev** **28**: 284–288, 2005
7. Fritsch MJ, Kienke S, Ankermann T, Padoin M, Mehdorn HM: Endoscopic third ventriculostomy in infants. **J Neurosurg** **103** (1 Suppl):50–53, 2005
8. Greenfield JP, Hoffman C, Kuo E, Christos PJ, Souweidane MM: Intraoperative assessment of endoscopic third ventriculostomy success. Clinical article. **J Neurosurg Pediatr** **2**: 298–303, 2008
9. Kadrian D, van Gelder J, Florida D, Jones R, Vonau M, Teo C, et al: Long-term reliability of endoscopic third ventriculostomy. **Neurosurgery** **56**:1271–1278, 2005
10. Kestle JR, Drake JM, Cochrane DD, Milner R, Walker ML, Abbott R III, et al: Lack of benefit of endoscopic ventriculoperitoneal shunt insertion: a multicenter randomized trial. **J Neurosurg** **98**:284–290, 2003
11. Koch D, Wagner W: Endoscopic third ventriculostomy in infants of less than 1 year of age: which factors influence the outcome? **Childs Nerv Syst** **20**:405–411, 2004
12. Kulkarni AV, Cochrane DD, McNeely PD, Shams I: Medical, social, and economic factors associated with health-related quality of life in Canadian children with hydrocephalus. **J Pediatr** **153**:689–695, 2008
13. Kulkarni AV, Drake JM, Kestle JR, Mallucci CL, Sgouros S, Constantini S: Endoscopic third ventriculostomy vs cerebrospinal fluid shunt in the treatment of hydrocephalus in children: a propensity score-adjusted analysis. **Neurosurgery** [epub ahead of print], 2010
14. Kulkarni AV, Drake JM, Mallucci CL, Sgouros S, Roth J, Constantini S: Endoscopic third ventriculostomy in the treatment of childhood hydrocephalus. **J Pediatr** **155**:254–259, e1, 2009
15. Kulkarni AV, Shams I: Quality of life in children with hydrocephalus: results from the Hospital for Sick Children, Toronto. **J Neurosurg** **107** (5 Suppl):358–364, 2007
16. O'Brien DF, Javadpour M, Collins DR, Spennato P, Mallucci CL: Endoscopic third ventriculostomy: an outcome analysis of primary cases and procedures performed after ventriculoperitoneal shunt malfunction. **J Neurosurg** **103** (5 Suppl): 393–400, 2005
17. Tuli S, Alshail E, Drake J: Third ventriculostomy versus cerebrospinal fluid shunt as a first procedure in pediatric hydrocephalus. **Pediatr Neurosurg** **30**:11–15, 1999
18. Wagner W, Koch D: Mechanisms of failure after endoscopic third ventriculostomy in young infants. **J Neurosurg** **103** (1 Suppl):43–49, 2005
19. Warf BC, Kulkarni AV: Intraoperative assessment of cerebral aqueduct patency and cisternal scarring: impact on success of endoscopic third ventriculostomy in 403 African children. Clinical article. **J Neurosurg Pediatr** **5**:204–209, 2010
20. Yadav YR, Jaiswal S, Adam N, Basoor A, Jain G: Endoscopic third ventriculostomy in infants. **Neurol India** **54**:161–163, 2006

Manuscript submitted January 15, 2010.

Accepted August 4, 2010.

Current address for Dr. Sgouros: “Attikon” University Hospital, University of Athens, Greece.

Address correspondence to: Abhaya V. Kulkarni, M.D., Ph.D., Hospital for Sick Children, Room 1503, 555 University Avenue, Toronto, Ontario M5G 1X8, Canada. email: abhaya.kulkarni@sickkids.ca.