

Intraocular Pressure in Pediatric Patients During Prone Surgery

Peter Szmuk, MD,*† Jeffrey W. Steiner, DO,*† Radu B. Pop, MS,*† Jing You, MS,‡§
David R. Weakley, Jr., MD,|| Dale M. Swift, MD,¶ and Daniel I. Sessler, MD§#

BACKGROUND: Intraoperative intraocular pressure (IOP) in the prone position and IOP changes over time have not been evaluated in pediatric surgical patients. We sought to determine time-dependent changes in IOP in children undergoing surgery in prone position.

METHODS: Thirty patients undergoing neurosurgical procedures in prone position were included. Using a pulse-mode pneumatonometer, IOP was measured in supine position after induction and before emergence of anesthesia and in prone position before the start and after the end of surgery. IOP changes over time in the prone position were assessed with a linear mixed model (i.e., random slope and intercept model) to adjust for the within-patient correlation.

RESULTS: IOP in prone position increased by an average of 2.2 mmHg per hour ($P < 0.001$). Sixty-three percent of patients (95% confidence interval [CI], 46%–81%) had at least 1 IOP value exceeding 30 mmHg, and 13% (95% CI, 1%–25%) had at least 1 IOP value exceeding 40 mmHg while prone. Mean IOP increased 7 mmHg (95% CI, 6–9) during the position change from supine to prone ($P < 0.001$) and decreased 10 mmHg (95% CI, 9–12) after changing the position from prone back to supine ($P < 0.001$).

CONCLUSIONS: Changing position from supine to prone significantly increases IOP in anesthetized pediatric patients. Moreover, the IOP continued to increase during surgery and reached potentially harmful values, especially when combined with low mean arterial blood pressures that are common during major surgery. (Anesth Analg 2013;116:1309–13)

Vision loss is a devastating perioperative complication^{1,2} that has been reported as a complication of cranial vault reconstruction,³ spine surgery,⁴ and orbital surgery.⁵ A United States national study estimated the overall incidence of perioperative visual loss to be 2.4 per 10,000 cases (0.02%), but that the risk is 0.03% for spinal fusion and 0.09% for cardiac surgery.⁶

An unexpected finding from analysis of the Nationwide Inpatient Sample was an alarmingly high risk of pediatric patients developing postoperative visual loss after all surgical procedures (odds ratio 6.9 versus adults). The odds ratio for developing visual loss in patients younger than 18 years after spinal fusion surgery was 5.8,⁷ whereas the odds ratio of young patients to develop cortical blindness versus adults across all procedures was 64.⁶ The reason for the

increased visual loss risk in pediatric patients is not clear, but an embolic mechanism seems more likely than stroke (which is uncommon in children).⁶ There are nonetheless only sporadic published reports of postoperative visual loss in pediatric patients.^{3–5,8–10}

The causes of vision loss after spine surgery in prone position remain poorly understood, but appear to be multifactorial and may include impaired perfusion of the eye, occlusion of retinal vessels, or an “eye compartment syndrome” caused by increased orbital pressure and decreased perfusion secondary to use of large amounts of crystalloids.¹¹ Inadequate ocular perfusion pressure can cause retinal ischemia and may contribute to postoperative visual loss.^{12,13} Ocular perfusion pressure is commonly defined as the difference between mean arterial blood pressure and intraocular pressure (IOP).¹⁴ At a given mean arterial pressure, retinal perfusion pressure is determined by IOP. Factors that influence perioperative IOP are thus of considerable interest. IOP can be influenced by general anesthesia, fluid balance, and end-tidal carbon dioxide partial pressure. Aqueous humor flow, choroidal blood volume, central venous pressure, and extraocular muscle tone also contribute.¹⁵ Positioning is yet another factor that influences IOP during surgery.¹⁶ For example, IOP is increased by prone^{17,18} and deep Trendelenburg¹⁹ positions, with the increases being comparable with and without general anesthesia.^{19,20} IOP also continues to increase over time in the prone position,^{16,19–21} an effect that is thought to result from continued production of aqueous fluid by the ciliary body inside the eye¹⁹ or to the accumulation of edema in the orbit.¹¹ With only a single exception,¹⁸ all studies have found a time-dependent increase in IOP in adults.

The normal distribution of IOP is well established in unanesthetized, pediatric subjects.²² However, intraoperative IOP and the extent to which it changes over time have

From the *Department of Anesthesiology and Pain Medicine, University of Texas Southwestern Medical School and Children’s Medical Center at Dallas, Texas; †Outcome Research Consortium and Departments of ‡Quantitative Health Sciences and §Outcomes Research, Cleveland Clinic, Cleveland, Ohio; ||Department of Ophthalmology, University of Texas Southwestern Medical School and Children’s Medical Center at Dallas, Texas; ¶Neurosurgery for Children, Children’s Medical Center at Dallas, Texas; and #Population Health Research Institute, McMaster University, Hamilton, Ontario, Canada.

Accepted for publication January 29, 2013.

Funding: Departmental funding. None of the authors has any personal financial interest in this work.

The authors declare no conflicts of interest.

This report was previously presented, in part, at the Society for Pediatric Anesthesia Meeting, March 31–April 3, 2011, San Diego, CA; American Society of Anesthesia 2009 Annual Meeting, October 17–21, 2009, New Orleans, LA.

Reprints will not be available from the authors.

Address correspondence to Peter Szmuk, MD, Department of Anesthesiology and Pain Medicine, University of Texas Southwestern Medical Center Dallas and Children’s Medical Center at Dallas, 1935 Medical District Drive, B3304, Dallas, TX 75235. Address e-mail to Peter.Szmuk@UTSouthwestern.edu.

Copyright © 2013 International Anesthesia Research Society
DOI: 10.1213/ANE.0b013e31828d3730

yet to be evaluated fully in pediatric patients. Furthermore, the effect of prolonged prone positioning remains unknown in pediatric patients. We thus sought to determine time-dependent changes in IOP in children undergoing surgery in the prone position.

METHODS

With approval of the University of Texas Southwestern IRB and written consent from parents, we enrolled consecutive patients from newborn to 18 years of age who were scheduled for neurosurgery in prone position with an expected duration exceeding 2 hours. Patients with a history of increased IOP or glaucoma, known visual impairment, heart failure, or ASA physical status scores >3 were excluded.

Both induction and maintenance of anesthesia were left to the discretion of the anesthesiologist, but typically included propofol (1.5–3.0 mg/kg), fentanyl (1–2 mcg/kg), vecuronium (0.1 mg/kg), and sevoflurane or isoflurane at approximately 1 minimum alveolar concentration. All patients were given dexamethasone 0.5 mg/kg¹ shortly after induction. Arterial blood pressure was monitored from an arterial catheter. Mechanical ventilation was adjusted to provide an end-tidal PCO₂ near 35 mmHg. Anesthetic administration was adjusted as necessary to maintain mean arterial blood pressure and heart rate about 20% below preinduction values.

As is routine in these cases, the patient’s head was secured with skull pins which allowed free access to the eyes while avoiding any direct mechanical pressure to the globe. The patient’s head was elevated 10° to reduce venous stasis.

Patients were given 5 to 7 mL/kg lactated Ringer’s solution in the immediate postinduction period, which was followed by 5 mL/kg/h maintenance hydration. Additional lactated Ringer’s solution was given as necessary to replace blood loss (usually in a 3:1 ratio) and to maintain mean arterial blood pressure about 20% below the preinduction value, heart rate within 20% of the preoperative value, and urine output ≥0.5 mL/kg/h. Blood was transfused as necessary to maintain a hematocrit ≥30%.

Morphometric and demographic characteristics were recorded, along with mean arterial blood pressure, blood loss, fluid administration, urine output, and the duration of surgery. IOP was measured with a Model 30 Classic Tm pulse-mode pneumatonometer (Reichert Technologies, Depew, NY). The pneumatonometer is self-calibrating and records 40 values per second; we thus made a single measurement for each eye at each time point. All measurements were performed by the same investigator (RBP). This system is well validated in pediatric patients.²³

IOP was recorded first with patients supine 15 minutes after anesthetic induction but before the head was positioned in pins; second, 15 minutes after patients were turned prone; third, at the end of surgery while the patient was still in prone position; and fourth, 10 minutes after patients were turned supine at the end of surgery before tracheal extubation. When possible, IOP was determined in each eye at each measurement interval. Anesthesia was discontinued only after the final IOP measurements in supine position.

IOP changes over time in the prone position were assessed using a linear mixed model (i.e., a random slope

Table 1. Demographics Baseline and Intraoperative Characteristics (N = 30)

Variables	Statistics ^a
Age, mo	104 (58)
Weight, kg	38 (25)
Length, cm	131 (33)
Gender (male), %	40
ASA physical status class	1.7 (0.6)
Type of surgery, %	
Lumbar laminectomy for tethered cord	43
Craniotomy	13
Chiari decompression	43
Volatile anesthetic used, %	
Isoflurane	38
Desflurane	4
Sevoflurane	58
Length of surgery, h	4.3 (1.3)

^aSummary statistics presented as mean (SD) or % of patients.

and intercept model) with an unstructured covariance matrix to adjust for the within-patient correlation. This model assumes that patient effects (intercepts) and time effects (slopes—IOP changes over time) are random (i.e., differ among patients). The average IOP change per hour in the prone position was estimated with 95% confidence interval (CI). In addition, percentages of patients who had at least 1 IOP in the prone position exceeding 30 and 40 mmHg were reported along with Wald confidence limits.

We assessed the IOP change from supine to prone position by comparing the initial measurement in the supine position and the first measurement in the prone position. Similarly, the IOP change from prone back to supine position was also assessed by comparing the final measurement in the prone position and the measurement in the supine position after changing back from the prone position. Pressures were compared with paired Student *t* tests. The corresponding mean (95% CI) of the IOP changes were estimated.

A total sample size of 26 was required to be able to detect a change of 2 mmHg or more per hour in IOP at the 0.05 significance level and 90% power, assuming an SD of 3 mmHg and a correlation of 0.5 based on previous experience. A total sample size of 30 patients was thus selected. SAS statistical software 9.2 for Windows (SAS Institute, Cary, NC) was used for all analyses.

RESULTS

Thirty pediatric patients were included in the study. Table 1 provides the summary of the demographics baseline and intraoperative characteristics. Blood loss was minimal in all patients, and none required blood replacement.

The change of IOP over time during the prone position did not vary by eye side (*P* = 0.19, assessment of interaction). We thus averaged IOP for the left and right sides when both were available at a given time point, or used the nonmissing IOP measurement when only 1 was available.

IOP changed approximately linearly over time in patients with >2 prone measurements (Figs. 1 and 2); a random slope and intercept model was therefore used to assess IOP change over time during the prone position. The estimated average slope was 2.2 (95% CI, 1.5–2.9) mmHg per hour, indicating an average of 2.2 mmHg increase in

Figure 1. Left and right eye intraocular pressure (IOP) for 30 pediatric patients undergoing surgery in prone position. Zero on x-axis refers to the first IOP measurement in prone position. Each line represents IOP measurements for each patient.

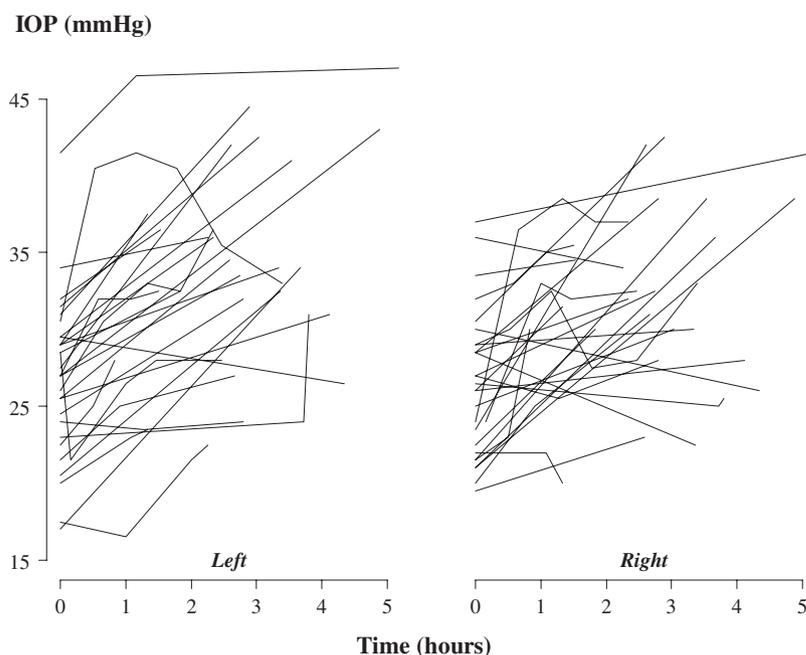
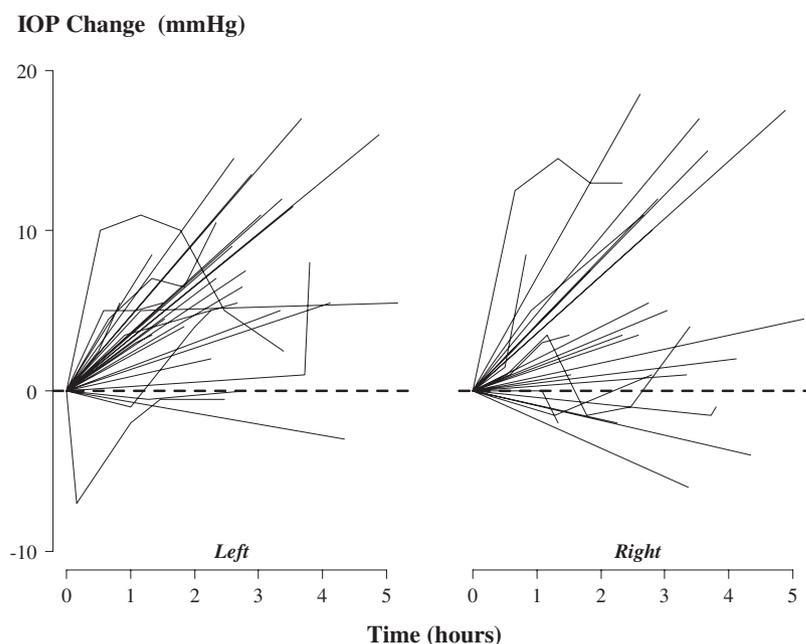


Figure 2. Changes in intraocular pressure (IOP) over time from first prone position (left and right eye) for 30 pediatric patients undergoing surgery in prone position. Zero on x-axis refers to the first IOP measurement in prone position; zero on y-axis (dashed line) refers to no change in IOP. Each line represents changes in IOP for each patient.



IOP per hour in prone position ($P < 0.001$; Fig. 3, middle panel).

Sixty-three percent of patients (95% CI, 46%–81%) had at least 1 IOP value exceeding 30 mmHg, and 13% (95% CI, 1%–25%) had at least 1 IOP value exceeding 40 mmHg while prone.

The observed mean (SD) of IOP was 19 (3) mmHg for the initial measurement in the supine position and 27 (5) mmHg for the first measurement in the prone position (Fig. 3, left panel). Mean IOP thus increased 7 (95% CI, 6–9) mmHg during the position change from supine to prone ($P < 0.001$).

The observed mean (SD) of IOP was 32 (6) mmHg at the last IOP measurement in the prone position and 22 (4)

mmHg in the supine position after changing back from the prone position (Fig. 3, right panel). Mean IOP thus decreased 10 (95% CI, 9–12) mmHg after changing the position from prone back to supine ($P < 0.001$). One patient did not have IOP measured in supine position after changing back from prone position; thus 29 patients were included in this analysis.

DISCUSSION

Our results indicate that in pediatric patients mean IOP increased 7 (95% CI, 6–9) mmHg during the position change from supine to prone ($P < 0.001$) and decreased 10 (95% CI, 9–12) mmHg after changing the position from prone back

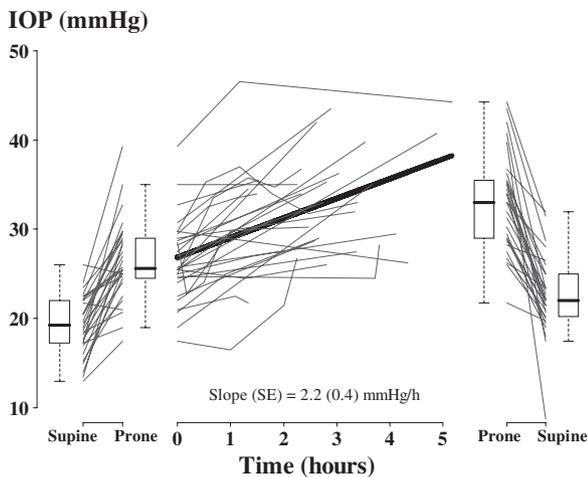


Figure 3. Average (left and right eye) intraocular pressure (IOP) for 30 pediatric patients undergoing surgery in prone position. Left panel: Box plots of IOP in the initial supine position and the first measurement in the prone position, and a plot of individual changes from supine to prone position. Middle panel: Plot of the IOP over time from the first IOP measurement in the prone position. Right panel: Box plots of the final IOP measurement in the prone position and IOP in the final supine position, and a plot of individual changes from prone back to supine position. Each line represents changes in IOP for each patient. The middle, upper, and lower edges of the box indicate the 50th, 75th, and 25th percentile of the data. The ends of the vertical lines indicate 1.5 times the interquartile range.

to supine ($P < 0.001$). IOP in prone position increased by an average of 2.2 mmHg per hour ($P < 0.001$). These results are comparable with those in adults.^{4,5}

IOP in children is generally low, reaching adult pressures at about 12 years of age. For example, IOP in a normal pediatric population ranges from a mean (SD) of 8 (3) mmHg in infants younger than 1 year to 15 (3) mmHg in children aged 11 to 12 years.^{5,22} These values were slightly lower than the 19 (3) mmHg we observed in our patients while supine after induction of anesthesia. The discrepancy might result from differing approaches to IOP determination because previous measurements were made with conventional tonometry rather than applanation pneumotometry as in our study. The distinction is important as Eisenberg et al.²³ found that conventional tonometry significantly underestimates IOP in pediatric patients; in contrast, applanation pneumotometry remained accurate.

Sugata et al.²⁴ showed that the choice of anesthetic (sevoflurane or propofol) does not have significant effect on IOP changes during short periods of prone surgery; however, changes in body position have a noticeable effect. IOP increased from 19 (3) mmHg while supine to 27 (5) mmHg after patients were initially turned prone. IOP then decreased from 32 (6) mmHg while prone at the end of surgery to 22 (4) mmHg in the supine position. The effect of changing position was thus similar at the beginning and end of surgery.

As in adults, IOP increased over time in the prone position. However, the slope was only 2.2 (0.4) mmHg/h. Consequently, mean pressure was 32 (6) mmHg at the end of procedures lasting 4.3 (1.3) hours. We found that 63% of our pediatric patients (95% CI, 46%–81%) had IOP exceeding 30 mmHg, and 13% (95% CI, 1%–25%) had IOP

exceeding 40 mmHg while prone. Sustained increases in IOP over time have been reported to have negative effects both in animal and humans studies.^{12,13} It is thus plausible that IOPs exceeding 40 mmHg in our patients could have put them at risk of visual loss.

Blood flow to the optic nerve head is regulated and thus remains relatively constant despite changes in IOP.²⁰ The IOP at which autoregulation fails in pediatric patients is unknown, but in adult volunteers IOP remained nearly constant until ocular pressures reached 40 mmHg.²⁵ Even if 40 mmHg were the safe threshold in pediatric patients, 13% of our patients exceed this pressure. However, it is conceivable that blood flow in the optic nerve is lower in infants and approaches adult values in older children. It is thus concerning that IOP exceeded 30 mmHg in more than half of the patients we evaluated during prone surgery.

Grant et al.²¹ evaluated the anatomy of the posterior optic nerve in volunteers laying supine or prone for 5 hours by using ultrasound imaging. In the prone position only, there was a thickening of the choroid layer which progressed over time, along with an increase in optic nerve diameter. These results support the hypothesis that time-dependent increases in IOP result at least partially from orbital venous congestion and its effect on episcleral venous congestion.

That being said, the clinical implications of increased IOP remain poorly understood. Thus, while pressures exceeding 40 mmHg are certainly concerning, it is unknown whether relatively brief periods (i.e., hours) at such pressures actually provoke visual loss. Ocular perfusion pressure, by definition, depends on mean arterial blood pressure, but blood pressure is often low during surgery which presumably aggravates risk. We also note that it is difficult to accurately assess visual ability in infants and children and that much postoperative visual loss may never be detected clinically or even in studies. Finally, IOP, and changes in IOP during surgery, varied considerably from patient to patient. A consequence is that the average values we report poorly predict individual pressures; a corollary is that without individual IOP measurements, it will be difficult to predict a given patient's pressure at any particular time.

Reported differences in IOP among studies may result from various methods used to position patients' heads and from various methods for measuring IOP. The head was supported by scalp pins in all our patients; consequently, there was no direct pressure on the eyes at any time.

Our study was far too small to establish a cause and effect relation between IOP changes and visual loss, and that was never among our goals. Instead, we sought to determine time-dependent changes in IOP in children undergoing surgery in prone position. Due to technical difficulties with IOP assessment during prone position, we measured IOP only before and after surgery in most patients. We were thus unable to fully characterize the shape of the IOP curve over time and have assumed based on limited data that it is approximately linear. The mean duration of surgery was 4.3 (1.3) hours. We do not know whether IOP would continue to increase during longer operations, or if it would reach a plateau.

In summary, changing from supine to the prone position significantly increases IOP in anesthetized pediatric patients. Moreover, IOP progressively increased during surgery and often reached potential harmful values. ■■

DISCLOSURES

Name: Peter Szmuk, MD.

Contribution: This author helped design and conduct the study and write the manuscript.

Attestation: Peter Szmuk has seen the original study data, reviewed the analysis of the data, approved the final manuscript, and is the author responsible for archiving the study files.

Name: Jeffrey W. Steiner, DO.

Contribution: This author helped conduct the study and write the manuscript.

Attestation: Jeffrey W. Steiner has seen the original study data, reviewed the analysis of the data, and approved the final manuscript.

Name: Radu B. Pop, MS.

Contribution: This author helped conduct the study.

Attestation: Radu B. Pop has seen the original study data and approved the final manuscript.

Name: Jing You, MS.

Contribution: This author helped analyze the data.

Attestation: Jing You has seen the original study data, reviewed the analysis of the data, and approved the final manuscript.

Name: David R. Weakley, Jr., MD.

Contribution: This author helped design and conduct the study.

Attestation: David R. Weakley has seen the original study data and approved the final manuscript.

Name: Dale M. Swift, MD.

Contribution: This author helped design and conduct the study.

Attestation: Dale M. Swift has seen the original study data and approved the final manuscript.

Name: Daniel I. Sessler, MD.

Contribution: This author helped design the study, analyze the data, and write the manuscript.

Attestation: Daniel I. Sessler has seen the original study data, reviewed the analysis of the data, and approved the final manuscript.

This manuscript was handled by: Peter J. Davis, MD.

ACKNOWLEDGMENTS

The authors would like to thank Frederick H. Sklar, MD, Bradley E. Weprin, MD, David J. Sacco, MD, and Angela V. Price, MD from the Neurosurgery For Children Group for their support for this project and Maria F. Ortega, MD, Mary F. Harris, MD, and Amy J. Hogge, MD from the Children's Anesthesia Resources and Anesthesia for Children Groups for their extensive help with this study.

REFERENCES

- Roth S, Barach P. Postoperative visual loss: still no answers—yet. *Anesthesiology* 2001;95:575–7
- Roth S, Thisted RA, Erickson JP, Black S, Schreider BD. Eye injuries after nonocular surgery. A study of 60,965 anesthetics from 1988 to 1992. *Anesthesiology* 1996;85:1020–7
- Lee J, Crawford MW, Drake J, Buncic JR, Forrest C. Anterior ischemic optic neuropathy complicating cranial vault reconstruction for sagittal synostosis in a child. *J Craniofac Surg* 2005;16:559–62
- Ho VT, Newman NJ, Song S, Ksiazek S, Roth S. Ischemic optic neuropathy following spine surgery. *J Neurosurg Anesthesiol* 2005;17:38–44
- Jamous M, Satoh K, Kageji T, Satomi J, Matsubara S, Nagahiro S, Hayashi M, Nakagawa S. Anterior ischemic optic neuropathy after combined ophthalmic artery embolization and craniofacial surgery—case report. *Neurol Med Chir (Tokyo)* 2001;41:419–22
- Shen Y, Drum M, Roth S. The prevalence of perioperative visual loss in the United States: a 10-year study from 1996 to 2005 of spinal, orthopedic, cardiac, and general surgery. *Anesth Analg* 2009;109:1534–45
- Patil CG, Lad EM, Lad SP, Ho C, Boakye M. Visual loss after spine surgery: a population-based study. *Spine (Phila Pa 1976)* 2008;33:1491–6
- Lapeyraque AL, Haddad E, André JL, Brémond-Gignac D, Taylor CM, Rianthavorn P, Salusky IB, Loirat C. Sudden blindness caused by anterior ischemic optic neuropathy in 5 children on continuous peritoneal dialysis. *Am J Kidney Dis* 2003;42:E3–9
- Chutorian AM, Winterkorn JM, Geffner M. Anterior ischemic optic neuropathy in children: case reports and review of the literature. *Pediatr Neurol* 2002;26:358–64
- Giordano M, Seminara G, Infantone L, de Pascale E, Giordano C. A case of transient blindness in a postoperative hyponatremic child. *Clin Nephrol* 2000;53:222–5
- Farag E, Doyle DJ. Vision loss after spine surgery: a new hypothesis. *Can J Anaesth* 2006;53:420
- Setogawa A, Kawai. Measurement of intraocular pressure by both invasive and noninvasive techniques in rabbits exposed to head-down tilt. *Jpn J Physiol* 1998;48:25–31
- Mader TH, Taylor GR, Hunter N, Caputo M, Meehan RT. Intraocular pressure, retinal vascular, and visual acuity changes during 48 hours of 10 degrees head-down tilt. *Aviat Space Environ Med* 1990;61:810–3
- Hayreh SS. Anterior ischemic optic neuropathy. *Clin Neurosci* 1997;4:251–63
- Feldman MA, Patel A. Anesthesia for eye, ear, nose, and throat surgery. In: Miller RD, ed. *Miller's Anesthesia*. 7th ed. Philadelphia, PA: Churchill Livingstone, Elsevier, 2010:2379
- Linder BJ, Trick GL, Wolf ML. Altering body position affects intraocular pressure and visual function. *Invest Ophthalmol Vis Sci* 1988;29:1492–7
- Cheng MA, Todorov A, Tempelhoff R, McHugh T, Crowder CM, Laurysen C. The effect of prone positioning on intraocular pressure in anesthetized patients. *Anesthesiology* 2001;95:1351–5
- Hunt K, Bajekal R, Calder I, Meacher R, Eliahoo J, Acheson JF. Changes in intraocular pressure in anesthetized prone patients. *J Neurosurg Anesthesiol* 2004;16:287–90
- Awad H, Santilli S, Ohr M, Roth A, Yan W, Fernandez S, Roth S, Patel V. The effects of steep trendelenburg positioning on intraocular pressure during robotic radical prostatectomy. *Anesth Analg* 2009;109:473–8
- Walick KS, Kragh JE, Jr, Ward JA, Crawford JJ: Changes in intraocular pressure due to surgical positioning: studying potential risk for postoperative vision loss. *Spine (Phila Pa 1976)* 2007;32:2591–5
- Grant GP, Szirth BC, Bennett HL, Huang SS, Thaker RS, Heary RF, Turbin RE. Effects of prone and reverse trendelenburg positioning on ocular parameters. *Anesthesiology* 2010;112:57–65
- Sihota R, Tuli D, Dada T, Gupta V, Sachdeva MM. Distribution and determinants of intraocular pressure in a normal pediatric population. *J Pediatr Ophthalmol Strabismus* 2006;43:14–8; quiz 36–7
- Eisenberg DL, Sherman BG, McKeown CA, Schuman JS. Tonometry in adults and children. A manometric evaluation of pneumatonometry, applanation, and TonoPen in vitro and in vivo. *Ophthalmology* 1998;105:1173–81
- Sugata A, Hayashi H, Kawaguchi M, Hasuwa K, Nomura Y, Furuya H. Changes in intraocular pressure during prone spine surgery under propofol and sevoflurane anesthesia. *J Neurosurg Anesthesiol* 2012;24:152–6
- Pillunat LE, Anderson DR, Knighton RW, Joos KM, Feuer WJ. Autoregulation of human optic nerve head circulation in response to increased intraocular pressure. *Exp Eye Res* 1997;64:737–44