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Cataract (Surgical Management) Chart Abstraction

Print

Chart

March 20, 2018

RECORD IDENTIFIER: CATARACT00001



History

1. Date of birth	December 1950
2. Gender	<input type="radio"/> Male <input type="radio"/> Female
3. Visual complaint	Do symptoms impact activities of daily living? <input type="radio"/> Yes <input type="radio"/> No <input type="radio"/> Not Recorded
4. Medication history	Alpha-1a agonists (i.e. tamsulosin) <input type="radio"/> Yes <input type="radio"/> No <input type="radio"/> Not Recorded
	Anticoagulation <input type="radio"/> Yes <input type="radio"/> No <input type="radio"/> Not Recorded

Preoperative Examination

1. Corrected visual acuity (with glasses or most recent refraction)	20 / []
2. Refraction performed within last 12 months (or results of recent refraction documented?)	<input type="radio"/> Yes <input type="radio"/> No <input type="radio"/> Not Recorded
3. Glare testing	<input type="radio"/> Yes <input type="radio"/> No
4. Potential acuity testing	<input type="radio"/> Yes <input type="radio"/> No
5. Slit lamp exam	<input type="radio"/> Yes <input type="radio"/> No
6. Dilated pupil size	<input type="radio"/> Adequate <input type="radio"/> Small <input type="radio"/> Not Recorded
7. Corneal endothelium	<input type="radio"/> Normal <input type="radio"/> Guttae <input type="radio"/> Other Abnormality <input type="radio"/> Not Recorded
8. Presence of pseudoexfoliation material	<input type="radio"/> Yes <input type="radio"/> No
9. Cataract grading	<input type="radio"/> Documented <input type="radio"/> Not Documented
10. Dilated fundus exam	<input type="radio"/> Done <input type="radio"/> Not Done
11. Macula	<input type="radio"/> Normal <input type="radio"/> Abnormal <input type="radio"/> Not Recorded

Pre-Operative Management

1. Indication for surgery	<input type="checkbox"/> Visually significant cataract <input type="checkbox"/> Clinically significant anisometropia in the presence of cataract <input type="checkbox"/> Lens opacity interfering with diagnosis or management of posterior segment conditions <input type="checkbox"/> Lens causing inflammation <input type="checkbox"/> Lens inducing narrow angle or angle closure <input type="checkbox"/> Other <input type="checkbox"/> Not documented
2. Were risks, benefits and alternatives to surgery discussed?	<input type="radio"/> Yes <input type="radio"/> No <input type="radio"/> Not Recorded
3. Have the refractive goals and/or options for intraocular lenses and options been explained to the patient?	<input type="radio"/> Yes <input type="radio"/> No <input type="radio"/> Not Recorded
4. Was signed informed consent obtained?	<input type="radio"/> Yes <input type="radio"/> No <input type="radio"/> Not Recorded
5. Method of axial length measurement	<input type="checkbox"/> Contact Ultrasound <input type="checkbox"/> Immersion Ultrasound <input type="checkbox"/> Optical Biometry (i.e. IOLMaster, Lenstar) <input type="checkbox"/> Other
5a. If other, please specify:	[]
6. Method of keratometry	<input type="checkbox"/> Manual Keratometry <input type="checkbox"/> Corneal Topography <input type="checkbox"/> Optical Biometry (i.e. IOLMaster, Lenstar) <input type="checkbox"/> Other automated keratometer (i.e. hand held device) <input type="checkbox"/> Other
6a. If other, please specify:	[]
7. What formula was used for IOL power calculation?	<input type="checkbox"/> Haigis <input type="checkbox"/> Hoffer Q <input type="checkbox"/> Holladay <input type="checkbox"/> Holladay II <input type="checkbox"/> SRK-T <input type="checkbox"/> Other
7a. If other, please specify:	[]

Management and Post-Operative Care

1. Date of surgery	Month Year
2. Which surgical intervention was used?	<input type="radio"/> Large incision extracapsular extraction <input type="radio"/> Manual small incision extracapsular extraction (i.e. SICS) <input type="radio"/> Phacoemulsification
3. What type of anesthesia was utilized?	<input type="checkbox"/> General anesthesia <input type="checkbox"/> Peribulbar injection <input type="checkbox"/> Retrobulbar injection <input type="checkbox"/> Topical <input type="checkbox"/> Topical with intracameral <input type="checkbox"/> Sub tenon <input type="checkbox"/> Other
3a. If other, please specify:	[]
4. What refractive target was selected to the nearest 0.25 diopters?	+ - [] d
5. IOL Placement	<input type="radio"/> Anterior chamber <input type="radio"/> Iris fixated <input type="radio"/> Posterior capsule <input type="radio"/> Sulcus <input type="radio"/> Sulcus with suture fixation
6. Which intraoperative complications were encountered?	<input type="checkbox"/> None <input type="checkbox"/> Iris prolapse or injury <input type="checkbox"/> Loss of lens fragments into vitreous <input type="checkbox"/> Posterior capsular rupture <input type="checkbox"/> Suprachoroidal <input type="checkbox"/> Vitreous loss <input type="checkbox"/> Zonular dehiscence <input type="checkbox"/> Other
6a. If other, please specify:	[]
7. Which adjunctive procedures were necessary?	<input type="checkbox"/> None <input type="checkbox"/> Anterior vitrectomy <input type="checkbox"/> Capsular tension ring <input type="checkbox"/> Iris retractors or expanders <input type="checkbox"/> Other
7a. If other, please specify:	[]
8. Were intracameral or subconjunctival antibiotics given at the time of surgery?	<input type="radio"/> Yes <input type="radio"/> No

Post-Operative Care

1. Was the patient seen within 48 hours of surgery?	<input type="radio"/> Yes <input type="radio"/> No
2. Was topical antibiotic medication prescribed?	<input type="radio"/> Yes <input type="radio"/> No
2a. If Yes, was this medication prescribed to be given postoperatively on the same day as surgery?	<input type="radio"/> Yes <input type="radio"/> No <input type="radio"/> Not Recorded
3. Was topical anti-inflammatory medication prescribed?	<input type="radio"/> Yes <input type="radio"/> No
4. Which immediate (<48 hours) post-operative complications were encountered?	<input type="checkbox"/> None <input type="checkbox"/> Elevated intraocular pressure (>30mm HG) <input type="checkbox"/> Hyphema <input type="checkbox"/> Wound Leak <input type="checkbox"/> Other
4a. If "Other", please specify:	[]
5. Were verbal and/or written instructions given on the symptoms that should lead the patient to contact the ophthalmologist immediately?	<input type="radio"/> Yes <input type="radio"/> No <input type="radio"/> Not Recorded

Outcomes

1. Were any major post-operative complications encountered?	<input type="checkbox"/> None <input type="checkbox"/> Cystoid macular edema <input type="checkbox"/> Dislocated IOL <input type="checkbox"/> Endophthalmitis <input type="checkbox"/> Pseudophakic bullous keratopathy <input type="checkbox"/> Retained lens fragments (anterior chamber) <input type="checkbox"/> Retained lens fragments (vitreous cavity) <input type="checkbox"/> Retinal detachment <input type="checkbox"/> Toxic anterior segment syndrome (TASS) <input type="checkbox"/> Wound dehiscence <input type="checkbox"/> Wrong power IOL <input type="checkbox"/> Other
1a. If other, please specify:	[]
2. Was further surgery necessary within 90 days due to a complication of the cataract surgery?	<input type="radio"/> Yes <input type="radio"/> No
2a. If yes, specify reason and procedure below:	[]
3. Best corrected visual acuity within 90 days	20 / []
4. Refracted spherical equivalent (to the nearest 0.25 diopters)	+ - [] d
5. Did the patient have a pre-existing co-morbid condition that affected the postoperative visual acuity?	<input type="radio"/> Yes <input type="radio"/> No
6. If above acuity is worse than 20/40, what is the reason?	<input type="checkbox"/> Corneal disease <input type="checkbox"/> Optic nerve disease <input type="checkbox"/> Retinal disease <input type="checkbox"/> Not documented <input type="checkbox"/> Unable to determine <input type="checkbox"/> Other
6a. If other, please specify:	[]

Submit Chart

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