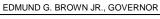


STATE BOARD OF OPTOMETRY

2450 DEL PASO ROAD, SUITE 105, SACRAMENTO, CA 95834 P (916) 575-7170 F (916) 575-7292 www.optometry .ca.gov





Continuing Education Course Approval Checklist

Title:

Provider Name:

□Completed Application Open to all Optometrists? ☑Yes □No Maintain Record Agreement?☑Yes □No

Correct Application Fee

□ Detailed Course Summary

Detailed Course Outline

PowerPoint and/or other Presentation Materials

□ Advertising (optional)

CV for EACH Course Instructor

☑License Verification for Each Course Instructor Disciplinary History? □Yes ☑No

Optometry, P	ested information below and it has b ed below, please attach a copy of th	APPROVALg an RSE APPROVALg an Receipt # Payor ID 1-2323 U 395914 approve continuing education been determined that the court	Beneficiary ID Amo 4395914 51 n (CE) courses after
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course Title	Course P	resentation Date	
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Joseph	Pruitt	Allan	
(First) rovider Mailing Address	(Last)	(Middl	e)
rovider Email Address /ill the proposed course be open t	· · · · · · · · · · · · · · · · · · ·	rists?	YES □NO
Do you agree to maintain and furnis of course content and attendance a rom the date of course presentatio	as the Board requires, for a perio on?	d of at least three years	∎YES □NO
Please provide the information below there are more instructors in the counstructor Name		each instructor or lecturer inv	
Joseph	Pruitt	Allan	
	(Last)	(Mi	iddle)
(First)			
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(First) icense Number <u>13429</u> Phone Number (<u>909</u>) <u>721-7751</u>	License Ty	_{ype} <u>TLG</u> ress pruitt.joseph@gmail	l.com
icense Number <u>13429</u>	License Ty Email Add	ress pruitt.joseph@gmail	

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1 The Pharmacological Management of Glaucoma

Joseph A. Pruitt, O.D., M.B.A., FAAO

Riverside-San Bernardino County Indian Health, Inc.

2 History

- Phase I:
 - Miotics; Pilocarpine, 1877
 - -Adrenergics; Epinephrine, 1920's
 - Systemic Carbonic Anhydrase Inhibitors: acetazolamide, 1954

3 🔲 History

- Phase II
 - Beta-Blockers; timolol, 1979
- Phase III
 - Alpha-Selective Agonists; apraclonidine, 1993
 - Topical Carbonic Anhydrase Inhibitors; dorzolamide, 1996
 - Prostaglandin analogues: latanoprost, 1997
 - Docosanoids: unoprostone, 2000
 - Prostamides; bimatoprost, 2001
 - -

4 History

- Phase IV
 - Neuroprotection

5 Mechanism of Action

Drugs increasing Aqueous Outflow

- Cholinergic agonists (miotics) = increased trabecular outflow
- Adrenergic agonists (epinephrine, dipivefrin) = increased trabecular outflow > increased uveoscleral outflow

6 Mechanism of Action

- Drugs that decrease Aqueous Production
 - Adrenergic blocking agents (beta-blockers)
 - Carbonic Anhydrase inhibitors

7 Hechanism of Action

- Drugs that increase Aqueous Outflow and/or decrease Aqueous Production
 - Alpha-2-agonists
 - Apraclonidine = decrease production + increase uveoscleral outflow
 - Brimonidine = decrease production + increase uveoscleral outflow
 - Prostanoids
 - Prostaglandin analogues = increased uveoscleral outflow

8 Mechanism of Action

Prostanoids continued

2

- Prostamides = ???; increased uveoscleral outflow + trabecular outflow

9 Drugs

- Cholinerigic Agonist (miotics)
 - Oldest class of glaucoma medications
 - Introduced in 1877 by von Weber
 - Acts directly on muscarinic receptors or indirectly by inhibiting Acetylcholine Esterase
 - Lowers IOP by increasing aqueous outflow
 - Works best with normal trabeculum

10 Drugs

- Cholinergic Agonists (continued)
 - Pilocarpine
 - ~20-30% IOP reduction (depending on concentration)
 - IOP becomes recalcitrant over time
 - -In which case an increase in concentration is warranted
 - Additive with most all other drugs
 - -Although, least with prostaglandins
 - Darker pigmented iridies typically require higher concentrations
 - Rarely used as a first-line drug

11 Drugs

- Cholinergic Agonist (continued)
 - Pilocarpine
 - Adverse Ocular effects
 - -Accommodative spasm
 - -Brow or head ache
 - -Pupillary block with shallow anterior chamber
 - -Miosis (decreased vision in low illumination)
 - -Myokymia (with higher concentrations)
 - --Chronic red eyes
 - -*Retinal Detachments
 - »Controversial, but remains a risk for aphakes and myopes with
 - predisposing conditions

12 Drugs

- Cholinergic Agonists (continued)
 - Echothiophate (Phospholine Iodide)
 - Indirect acting (does not mimic ACh)
 - Prolong duration (bid dosing)
 - · Narrow therapeutic index; rarely used
 - Carbachol
 - · Direct acting cholinomimetic
 - · Longer duration of action the pilocarpine
 - · Used by patients allergic or refractory to pilocarpine

- Cholinergic Agonists (continued)
 - Contraindications
 - Cataract (especially on visual axis)
 - < 40 years old

- Neovascular or uveitic glaucoma
- H/O Retinal Detachment

14 Drugs

• Adrenergic Agonists

- Non-selective agents

- Epinephrine
- Dipivalyl epinephrine (dipiverfrin)
- Alpha-2-agonists
 - Apraclonidine
 - brimonidine

15 Drugs

- Adrenergic Agonists (continued)
 - Epinephrine
 - Non-selective (alpha 1 + betas 1 & 2)
 - IOP reduction 15-20% with BID dosing
 - Adverse drugs reactions
 - -Rebound hyperemia
 - -CME in up to 20% of aphakic patients

16 Drugs

- Adrenergic Agonists (continued)
 - Epinephrine (continued)
 - Systemic effects via absorption at nasal mucosa
 - –Anxiety
 - -Palpitations
 - -Headache
 - -Hypertension
 - -Tachycardia
 - -Risk of stroke

17 Drugs

- Adrenergic Agonists (continued)
 - Epinephrine (continued)
 - Contraindications
 - -Concurrent systemic use of MAO inhibitors & tricyclics
 - –Aphakes
 - -Narrow angles
 - -Cerebral/coronary insufficiency
 - -Hypertension

18 Drugs

- Adrenergic Agonist (continued)
 - Dipivefrin (Propine)
 - Prodrug of epinephrine
 - Excellent corneal penetration secondary to being lipophilic; thus requires a weak concentration
 - Much safer than epinephrine
 - IOP reduction, adverse drug effects and contraindications presumably identical to epinephrine

4

19 Drugs

• Adrenergic Agonists (continued)

3

4

– Alpha-2-Receptor Agonists

• Apraclonidine (Iopidine)

- -Selective Alpha 2 agonist; some alpha 1
- -Derived from clonidine, but no CNS penetration
- -Efficacy not influenced by age, race or iris color

-Clinical use:

- »Prevention of post-op spikes
- »Short-term IOP control

20 🔲 Drugs

Adrenergic Agonist

– Alpha 2-Receptor Agonist

• Apraclonidine (continued)

-Adverse Effects:

»Conjunctival blanching

»Eyelid retraction

»Mydriasis

- »Tachyphylaxis (loss of efficacy)
- »Increased risk for hypersensitivity with prolonged use
- »Dry mouth and/or dry nose (20-50%)
- »Fatigue (~15%)
- »Headache
- »Sensation of head cold

21 Drugs

Adrenergic Agonists

– Alpha-2-Receptor Agonist

• Brimonidine (Alphagan P)

- -Stronger selective alpha-2 agonist
- -Lipophilic; thus better CNS and corneal penetration
- -No loss of efficacy
- -IOP reduction ~18-25%
- -Neuroprotective?
 - »Inhibits alutamate-mediated injury?

22 🔄 Drugs

Adrenergic Agonists

– Alpha-2-Receptor Agonist

• Brimonidine (Alphagan P)

-Adverse Effects

- »Conjunctival hyperemia
- »Blurred vision
- »FB sensation
- »Dry Eye
- »CNS adverse reactions (more so than apraclonidine)
 - Dry mouth
 - Fatigue/Drowsiness
 - Headache

23 Drugs

Adrenergic Agonists

- Alpha-2-Receptor Agonist
 - Brimonidine (Alphagan P)
 - -Contraindications:
 - »Patients taking MOA inhibitors
 - »Coronary insufficiency
 - »Cerebral insufficiency
 - »Recent myocardial infarction

24 Drugs

- Beta-Blockers
 - Primary target are beta receptors on ciliary body
 - Some formulations have 2 concentrations
 - 0.25% (blue cap) & 0.5% (yellow cap)
 - IOP reduction 0.5% ≥ 0.25%
 - Proposed to be less effective during sleep
 - Beta-receptors thought to "shut off" during sleep
 - -10-20% of patients unresponsive to drug class

25 🗍 Drugs

- Beta-Blockers (continued)
 - -Timolol (Timoptic)
 - Non-selective beta-blocker
 - "Gold standard" for comparison of newer agents
 - IOP reduction ranges from ~17% 28%
 - Known for "short-term escape"
 - -An exaggerated IOP reduction in the first 2-weeks of therapy
 - Known for "long-term drift"
 - -A gradual loss of IOP reduction @ 3-6 months (or later)

26 🔲 Drugs

- Beta-Blockers
 - Timolol (continued)
 - Hypotensive effect for up to 2 weeks on d/c
 - · Uniocular administration causes contralateral effect in untreated eye
 - · IOP reduction is minimized in patients already taking some form of beta-blocker
 - Various preparations:
 - -Timoptic, Timoptic XE, Istalol

27 Drugs

- Beta-Blockers (continued)
 - Levobunolol (Betagan)
 - Non-selective beta-blocker
 - IOP reduction similar to timolol 0.5%
 - Carteolol
 - Non-selective beta-blocker
 - IOP reduction similar to timolol 0.5%

- Beta-Blockers (continued)
 - Metipranolol (OptiPranolol)
 - Non-selective beta-blocker

6

- Efficacy similar to timolol 0.5% (perhaps slightly less?)
- Some corneal anesthetic effect
- Drug-induced anterior uveitis?
 - -Uncommon with U.S. concentration of 0.3%
 - --More common with European concentration of 0.6%
 - -Sterilization/Packaging? Or truly higher concentration

29 🗐 Drugs

- Beta-Blockers (continued)
 - Betaxolol
 - Relative selectivity for beta-1 receptors
 - Less effect on IOP than non-selective
 - Calcium antagonistic effects
 - -May reduce calcium in neuroprotection model
 - -Tends to be the beta-blocker of choice in normal-tension GLC

30 Drugs

- Beta-Blockers
 - Betaxolol (continued)
 - Better preservation of visual field
 - -mid 90's studies showed better mean sensitivities and less severe mean defects
 - -More recent study suggests blue/yellow field preserved (using SWAP)
 - Considered to be the "safest" beta-blocker for least side effects

31 Drugs

Beta-Blockers (continued)

- Adverse Systemic Effects:
 - –Cardiovascular
 - »In normal patients
 - 3-8 mmHg drop in systolic blood pressure
 - 1-5 mmHg drop in diastolic blood pressure
 - 2-4 beats/minute slower heart rate
 - · Blunted exercise-induced tachycardia
 - »Effects are worse and serious those already susceptible cardiovascular systems
 - -Pulmonary

»Bronchospasm

»Decreased Forced Expiratory Volume

»Effects should not manifest in normal patients

32 Drugs

- Beta-Blockers
 - Adverse Systemic Effects (continued)

-CNS

»Depression? (disputed and not confirmed)

- »Fatigue
- »Lethargy
- »Confusion
- »Memory loss
- »Dizziness
- »Insomnia

»Somnolence

33 Drugs

- Beta-Blockers
 - Adverse Systemic Effects (continued)
 - -Gastrointestinal
 - »Nausea
 - »Diarrhea
 - –Metabolic
 - »Reduction in HDLs
 - »Masks signs of hypoglycemia
 - -Sexual Dysfunction
 - »Impotence
 - »Decreased libido (disputed)

»

34 Drugs

- Beta-Blockers
 - Adverse Ocular Effects
 - Decreased tear production
 - Decreased goblet cell density
 - Corneal anesthesia

35 Drugs

Beta-Blockers

- Contraindications
 - Congestive Heart Failure (CHF)
 - -Possibly not if stable and treated???
 - Cardiac Arrhythmia
 - -Symptomatic bradycardia (e.g. syncope or presyncope)
 - -Bradycardia (< 55 bpm)
 - -Implanted pacemaker
 - Airway Disease
 - –Asthma
 - -COPD

36 Drugs

- Beta-Blockers
 - Contraindications (continued)
 - Hyperthyroidism
 - -"Thyroid storm" symptoms are masked by beta-blockers
 - Diabetes
 - -Hypoglycemia symptoms are masked by beta-blockers
 - Older patients
 - -High risk for undiagnosed and/or subclinical respiratory or cardiovascular disease
 - Depression???
 - –Your call...

37 Drugs

• Carbonic Anhydrase Inhibitors (Oral)

- Decreases bicarbonate entry into posterior chamber, which reduces hypertonic concentration; thus less aqueous production
- Need ~99% inhibition of carbonic anhydrase to achieve an effect on IOP
 Therefore, very high oral doses are required

38 Drugs

- Carbonic Anhydrase Inhibitors (Oral)
 - Acetazolamide (Diamox Sequels)
 - Good GI asorption with peak levels within 2-4 hours and maintained for 4-6
 hours
 - IOP reduction parallels plasma drug levels
 - Adverse Effects:
 - -Decreased libido
 - -Depression
 - -Fatigue
 - –Malaise
 - --Anorexia
 - –Weight loss

39 Drugs

- Carbonic Anhydrase Inhibitors (Oral)
 - Acetazolamide (Diamox Sequels)
 - Adverse Effects (continued)
 - –Numbness
 - -Polyuria
 - -GI upset
 - -Metabolic acidosis
 - -Hypokalemia (loss of potassium)
 - -Renal calculi (kidney stones)
 - –Transient myopia

40 Drugs

- Carbonic Anhydrase Inhibitors (Oral)
 - Acetazolamide (Diamox Sequels)
 - Contraindications
 - –Liver Disease
 - -Severe COPD
 - -Renal disease
 - -Pregnancy
 - -Severe cardiac disease

- Carbonic Anhydrase Inhibitor (Oral)
 - Methazolamide
 - IOP reduction is dose-dependent
 - · Good PO absorption; peaks @ 2-3 hours and maintained for 8 hours
 - Greater ocular penetration than acetazolamide
 - Adverse Effects:
 - -Best tolerated oral CAI
 - -Less acidosis, association w/kidney stones, less paresthesia
 - –Drowsiness
 - -Polyuria

9

-Dermatitis

42 Drugs

- Carbonic Anhydrase Inhibitors (Oral)
 - Methazolamide
 - Contraindications
 - -Essentially same as acetazolamide
 - -Better for patients with tendency toward kidney stones
 - -COPD- may be better tolerated since less metabolic acidosis

43 Drugs

- Carbonic Anhydrase Inhibitors (Topical)
 - -High activity against carbonic anhydrase II and IV enzymes
 - -Balanced lipid/water solubility for corneal penetration
 - -Effect limited to treated eye
 - -Effective during sleep

44 🖂 Drugs

• Carbonic Anhydrase Inhibitors (Topical)

- Dorzolamide (Trusopt)

- Sulfonamide derivative
- TID preferred for monotherapy; BID adjunctive
- IOP reduction ~21.8-24.4% (BID) & 22.2-26.2% (TID)
- Peak effect: 2 hours
- · Should not be used along with oral CAI

45 Drugs

- Carbonic Anhydrase Inhibitors (Topical)
 - Dorzolamide (continued)
 - Adverse Effects
 - -Stinging (secondary to acidic pH)
 - -Burning
 - -Blurred vision
 - –Allergic blepharoconjunctivitis in ~10%
 - -Bitter taste (25-30%)
 - -headaches

46 Drugs

- Carbonic Anhydrase Inhibitor (Topical)
 - Brinzolamide (Azopt)
 - Selective inhibitor of CA II isoenzyme
 - Sulfonamide derivative
 - BID or TID dosage yield similar reductions
 - -~19.1% reduction ranging from -2.7 to -3.9 mmHg

- Carbonic Anhydrase Inhibitor (Topical)
 - Brinzolamide (continued)
 - Adverse Effects
 - -Much less ocular discomfort (<6%)
 - –Itching
 - -FB sensation
 - -Dry eyes (< 2%)

48 🖃 Drugs

• Carbonic Anhydrase Inhibitor (Topical)

- Contraindications

- Severe Kidney Disease
- Liver Disease (not critical)
- COPD (not as critical)
- CHF (not as critical)

49 🔲 Drugs

- Prostanoids
 - Includes prostaglandin analogues, docosanoids and prostamides
 - Pharmacologic and ocular effects are dose-dependent

50 Drugs

- Prostanoids
 - Latanaprost (Xalatan)
 - Effective during sleep
 - IOP reduction independent of race, sex, age, iris color, type of glaucoma (with exception of inflammatory types) and/or previous therapy
 - Shelf life of ~6 weeks unrefrigerated
 - Dosage QD (not critical to be QHS)
 - IOP reduction ~25-35%; thus effective as monotherapy
 - Patent recently expired 3/2011

51 🖾 Drugs

Prostanoids

- Latanoprost (continued)

• Adverse Effects:

-Conjunctival hyperemia

-Stinging, burning and tearing

-Punctate corneal erosions

-Iris pigmentation darkens

»Thought to be permanent

- -Eyelid pigmentation
- –Hypertrichosis

»Reversible once discontinued

- -Anterior uveitis
- --CME
- -Migraines

52 🖾 Drugs

Prostanoids

- Latanoprost (continued)

- Contraindications
 - -History of uveitis

-Prior "incision surgery" or YAG capsulotomy

-Previous episodes of recurrent HSV keratitis

-*Relative* contraindication is <u>unilateral</u> therapy

- Prostanoids
 - Travoprost (Travatan Z)
 - Average IOP reduction between 7 to 8 mmHg

- Mean IOP reduction of up to 1.8-2.4 mmHg GREATER in blacks patients
- Rumored to have "slippage" after ~6-12 months (unsubstantiated)
- Dosage QD "evening" not bedtime

•

54 Drugs

Prostanoids

- Travoprost (continued)
 - Adverse Effects
 - -Conjunctival hyperemia
 - -FB sensation
 - -Tearing
 - -Dry Eyes
 - -Icreased pigmentation in iris and periorbital tissue
 - -Increased pigmentation and growth of eyelashes

55 Drugs

- Prostanoids
 - Brimatoprost (Lumigan)
 - Synthetic analogue of fatty acid prostamides
 - Prostamides are present in ocular tissues
 - Prostamides presumably lower IOP by the same mechanism as prostaglandins
 - Dosage QHS

56 Drugs

- Prostanoids
 - Brimatoprost (continued)
 - Adverse Effects
 - -Conjunctival hyperemia
 - -FB sensation
 - -Growth and darkening of eyelashes
 - »Latisse
 - -Pigmentation of periocular skin

57 🔲 Drugs

- Prostanoids
 - Tafluprost (Zioptan)
 - FDA approved for the treatment of OHTN & POAG Feb. 2012
 - * per Merck
 - -Average IOP reduction at 3 months -6 to -8 mmHg
 - -Average IOP reduction at 6 months -5 to -8 mmHg
 - -Adverse Effects
 - »Conjunctival Hyperemia (~4-20%) *

58 Drugs

- Prostanoids
 - Unoprostone (Rescula)
 - Originally on market 2000
 - Reintroduced 2012-2013
 - Discontinued March 31, 2015

Prostanoids

- VESNEO (latanoprostene bunod ophthalmic solution 0.024%)

 Per B&L: Upon instillation in the eye, latanoprostene bunod is rapidly metabolized to two actives; latanoprost acid, a prostaglandin analog, and nitric oxide. Nitric oxide is an important physiological signaling molecule, which plays a key role in IOP regulation in healthy eyes. *VESNEO* is thought to increase aqueous humor outflow by acting on both the uveoscleral (non-conventional) pathway via latanoprost acid, and trabecular meshwork and Schlemm's canal (conventional pathway) via nitric oxide signaling

- Awaiting FDA approval

• The FDA has set an action date of July 21, 2016 to complete its review, as per the Prescription Drug User Fee Act (PDUFA)

http://www.bausch.com/our-company/newsroom/2015-archive/fda-acceptanceof-new-drug-application-for-novel-glaucoma-candidate#.VucRDzjn-Uk

60 Drugs

Combination Formulations

– Cosopt

• timolol 0.5% & dorzolaminde 2%

Dosed BID

- As effective as timolol 0.5% BID & dorzolomide 2% TID
- Mean IOP changes similar to Xalatan

--IOP reduction ~25-35%

· Contraindications are the sum of each drug

Combination Formulations

- Combigan
 - brimonidine 0.2% + timolol 0.5%
 - -NOTE: not Alphagan P (either 0.15% or 0.1%)
 - Dosed q12h (*BID*)
 - As effective as both meds given separately
 - Better tolerated than 0.2% brimonidine TID
 - Better tolerated than Cosopt

62 Drugs

- Combination Formulations
 - Simbrinza
 - Brinzoloamide 1.0%/Brimonidine 0.2%



Combination Formulations

- Xalacom and Extravan
 - Latnoprost or travoprost combined with timolol 0.5%
 - Dosing schedules not clarified at this time
 - FDA approval delayed for "Xalacom"
 - -Although has been available in Europe for years...
 - IOP reduction may equal to, or less than, if separate drugs used together
 - Extravan may be closer to FDA approval

66 Conclusion

- Managing IOP remains a staple in glaucoma management
- Consider compliance and quality of life, an the frequently overlooked, cost

14

- "Above all else, do no harm"
 - Manage appropriately utilizing all therapeutics available
 - Do not let ANYONE go blind in your chair
 - Refer whenever appropriate and/or necessary

The Pharmacological Management of Glaucoma

Joseph A. Pruitt, O.D., M.B.A., FAAO Riverside-San Bernardino County Indian Health, Inc.

History

• Phase I: – Miotics; Pilocarpine, 1877 - Adrenergics; Epinephrine, 1920's - Systemic Carbonic Anhydrase Inhibitors: acetazolamide, 1954

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 Drugs that increase Aqueous Outflow and/or decrease Aqueous Production

- Alpha-2-agonists
- Apracionidine = decrease production + Increase uveoscleral outflow
 Brimonidine = decrease production + Increase uveoscleral outflow
- Prostanoids
- Prostaglandin analogues = increased uveoscieral outflow

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- Works best with normal trabeculum

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 Additive with most all other drugs

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 Darker pigmented indies typically require higher
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warranted

Rarely used as a first-line drug

Drugs

Cholinergic Agonist (continued)

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Drugs

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 < 40 years old

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 Apraclonidine brimonidine

Drugs

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 Rebound hyperemia
 CME in up to 20% of aphakic patients

Drugs

Adrenergic Agonists (continued)

- Epinephrine (continued) Systemic effects via absorption at nasal mucosa

- Anxiety Palpitations

- Headache - Hypertension - Tachycardia - Risk of stroke

Drugs

- · Adrenergic Agonists (continued) - Epinephrine (continued)
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Drugs

 Adrenergic Agonists (continued) - Alpha-2-Receptor Agonists
 Apracionaline (oplicine)

 - Selective Alpha 2 agonists ome alpha 1

 - Derived from dondinn, but no CAS penetration

 - Efficacy on Influence du yage, race or inscolor

 - Efficacy on Influence du yage, race or inscolor

 - Efficacy on Universitian of past-op spikes

 > Prevention of past-op spikes

_

Drugs

Adrenergic Agonist - Alpha 2-Receptor Agonisi Alpha 2-Receptor Agonist • Apracionidme (continued) • Adverse Effects: • Equipational blanching • Eyeliaf extraction • Eyeliaf extraction • Excessed risk for soft efficacy) • Excreased risk for hypersensitivity with prolonged use • Drymouth margier of you nose (20-50%) • Fatgue (~15%) • Fatgue (~15%) • Exensation of head cold

Drugs

 Adrenergic Agonists Alpha-2-Receptor Agonist
 Brimonidine (Alphagan P) Brimonidine (Alphagan P) – Stronger selective alpha-2 agonist – Upophilic; thus better CNS and corneal penetr: – No loss of efficacy – IOP reduction ~18-25% – Neuroprotective? > Inhibits glutamate-mediated Injury?

Drugs

 Adrenergic Agonists Adrenergic Agonists — Alpha-2-Receptor Agonist • Brimonidine(Alphagan P) - Adverse Effect > Conjunction hyperemia > Bluered vision > Brassation > Dry Size • CK3-Servise reactions (more so than apractonidine) • Dry mouth • Faigury/Drownless + Headsche

Drugs

 Adrenergic Agonists - Alpha-2-Receptor Agonist Patients taking MOA initiato
 Coronaryinsufficiency
 Cerebralinsufficiency
 Recent myocardial infarction

Drugs

- Beta-Blockers
- Primary target are beta receptors on ciliary body - Some formulations have 2 concentrations

• 0.25% (blue cap) & 0.5% (yellow cap) IOP reduction 0.5% ≥ 0.25%

- Proposed to be less effective during sleep
- Beta-receptors thought to "shut off" during sleep
- 10-20% of patients unresponsive to drug class

Drugs

Beta-Blockers (continued)

- Timolol (Timoptic) Non-selective beta-blocker

- "Gold standard" for comparison of newer agents
- IOP reduction ranges from ~17% 28%

- Known for "short-term escape"
 An esagerated IOP reduction in the first 2-weeks of therapy
 Known for "long-term drift"
 A graduations of IOP reduction @ 3-6 months (or later)

Drugs

- Beta-Blockers
- Timolol (continued)
- Hypotensive effect for up to 2 weeks on d/c
 Uniocular administration causes contralateral effect in
- untreated eve

4

- IOP reduction is minimized in patients already taking some form of beta-blocker
- Various preparations:
 Timoptic, Timoptic XE, Istatol

Drugs

 Beta-Blockers (continued) – Levobunolol (Betagan)

- Non-selective beta-blocker
- IOP reduction similar to timolol 0.5%
- Carteolol
- Non-selective beta-blocker
 IOP reduction similar to timolol 0.5%

Drugs

Beta-Blockers (continued) - Metipranolol (OptiPranolol)

- Non-selective beta-blocker
 Efficacy similar to timolol 0.5% (perhaps slightly less?)
- Some corneal anesthetic effect
- · Drug-induced anterior uveltis?
- Jrug-induced Briterior Uvents r Uncommun with U.S. concentration of 0.3% Morecommon with European concentration of 0.6% Sterilization/Packaging? Or truty bigher concentration

Drugs

- Beta-Blockers (continued) - Betaxolol Relative selectivity for beta-1 receptors
 - Less effect on IOP than non-selective
 - Calcium antigonistic effects
 May reduce calcium in neuroprotection model
 Tends to be the beta-blocker of choice in normal-tension GLC

Drugs .

- Beta-Blockers - Betaxolol (continued)
- Better preservation of visual field mid 90's studies showed better mean sensitivities and less severe mean defects
 - More recent study suggests blue/yellow field preserved (using SWAP)
- Considered to be the "safest" beta-blocker for least
 side effects

Drugs

atible

 Beta-Blockers (continued) - Adverse Systemic Effects: - Cardiovascular spècler armal parients 3-8 mmHg drop in systolic blood pressure 3-8 mmHg drop in distolic blood pressure 2-4 beats/minute slower heart rate Blunted exercise-induced tachyrardia -the are worse and serious these already sust

 Bronet 255450 Decreased Forced Expiratory Volume
 Effects should not manifest in normal patients Drugs

- Beta-Blockers
 - Adverse Systemic Effects (continued)
 - erse Systemic Effects (continued) Cro > pepression? (disputed and not confirmed) > fatigue > Lethurgy > Contusion > Memery loss > Distribers > Insomnia > Somnolence

Drugs

- Beta-Blockers
- ta-Blockers Adverse Systemic Effects (continued) Gastocintestinal Nausea Diarrhea Metabolic Metabolic

Drugs

Beta-Blockers

- Adverse Ocular Effects
- Decreased tear production
 Decreased gobiet cell density

Corneal anesthesia

Drugs

Beta-Blockers

- Contraindications
- Loncrafindications Congesitive Heart Failure (CHF) Possibly not if stable and treated??? Cardiac Arrhythmia Symptomatic tradycardia (c.g., syncope or presyncope) Budycardia (< 55 bpm) Implanted pacemaker Altway Disease Automatic Stable Sta

- Asthma COPD

- Beta-Blockers
 - Contraindications (continued)
 - Hyperthyroldism
 ---- Thyroid storm" symptoms are masked by beta-blockers
 - Throng storm: symptoms are masked by beta-blockers
 Diabetes
 Hypoglycenia symptoms are masked by beta-blockers
 Older patients
 Highrisk for undiagnosed and/or subclinical respiratory or
 cardioxecular disease

 - Depression??? -- Your call...

Drugs

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Carbonic Anhydrase Inhibitors (Oral) - Decreases bicarbonate entry into posterior chamber, which reduces hypertonic concentration; thus less aqueous production - Need ~99% Inhibition of carbonic anhydrase to achieve an effect on IOP

Therefore, very high oral doses are required

Drugs

- Carbonic Anhydrase Inhibitors (Oral)
- Actazolamide (Diamox Sequels)
 Good Gi asorption with peak levels within 2-4 hours and
 malatalned for 4-6 hours
 iOP reduction parallels plasma drug levels

 - IOP reduction paral
 Adverse Effects:

 Decreased libido
 Depression
 Fatigue
 Malaise
 Anorexia
 Weight loss

Drugs

- Carbonic Anhydrase Inhibitors (Oral) - Acetazolamide (Diamox Sequels) Adverse Effects (continued)

 - Adverse Effects (continued) Numbness Polyuria Giupset Metabolic acidosis Hypokalemia (loss of potassium) Renal calculi (kidney stones) Translent myopia

Drugs

Carbonic Anhydrase Inhibitors (Oral) . - Acetazolamide (Diamox Sequels) Contraindications

Contraindications - Uver Disease - Severe COPD - Renal disease - Pregnancy - Severe cardiac disease

Drugs

- Carbonic Anhydrase Inhibitor (Oral)
- Carbonic Anhydrase Inhibitor (Oral) Methazolamide IOP reduction is dose-dependent Good PO absorption; peske @ 2-3 hours and maintained for 8 hours Greater acular penetration than acetatolamide Adverse Effects: Best beferate oral CAI Loss addosis, association w/Aidney stones, less parestilesia Drowiness Polyuria Dermatilis

.

Drugs

· Carbonic Anhydrase Inhibitors (Oral) – Methazolamide

- Contraindications
 Essentially same as acetazolamide
 Better for patients with tendency towardkidney stones
 COPO-may be better tolerated since less metabolic acidosis

Drugs

 Carbonic Anhydrase Inhibitors (Topical) If ghactivity against carbonic anhydrase II and IV entymes
 Balanced lipid/water solubility for corneal penetration
 Effect limited to treated eye
 Effective during sleep

Drugs

- Carbonic Anhydrase Inhibitors (Topical) - Dorzolamide (Trusopt)

 - Sulfonamide derivative
 TID preferred for monotherapy; BID adjunctive
 - 10P reduction ~21.8-24.4% (BID) & 22.2-26.2% (TID)
 - Peak effect: 2 hours • Should not be used along with oral CA1

Drugs

· Carbonic Anhydrase Inhibitors (Topical) - Dorzolamide (continued)

- Adverse Effects
- Adverse Effects Stinging (secondary to acidic pH) Burning Blurred vision Allergic biepharoconjunctivitis in ~10% Bitter taste (25-30%)

- headaches

Drugs

 Carbonic Anhydrase Inhibitor (Topical) – Brinzolamide (Azopt) Selective inhibitor of CA II Isoenzyme

Sulfonamide derivative
 BID or TID dosage yield similar reductions

 ~ 19.1%: eduction: anging from -2.7 to -3.9 mmHg

Drugs

Carbonic Anhydrase Inhibitor (Topical) - Brinzolamide (continued) Adverse Effects
 – Much less ocular discomfort (<6%) Itching
 FB sensation
 Dry eyes (< 2%)

Drugs

Carbonic Anhydrase Inhibitor (Topical) - Contraindications Severe Kidney Disease

• Liver Disease (not critical) COPD (not as critical)

CHF (not as critical)

Drugs

Prostanoids

- Includes prostaglandin analogues, docosanoids and prostamides - Pharmacologic and ocular effects are dose-
- dependent

Drugs



- Latanaprost (Xalatan)
- Effective during sleep Chective during steep
 IOP reduction independent of race, sex, age, iris color, type of glaucoma (with exception of inflammatory types) and/or previous therapy
 Shelf Ilie of ~6 weeks unrefrigerated

- Dosage QD (not critical to be QHS)
 IOP reduction ~25-35%; thus effective as monotherapy
- Patent recently expired 3/2011

Drugs Prostanoids Prostanoids - Latanoprost (continued) - Adverse Effects - Goginacional Ingreenia - Single, Junnia, and tearing - Purctae correal evaluation - The provide the permanent - Synd pignentation - Hypervisitions - Reversible core discontinued - Anorem version - Anterior w - CME - Migraines

Drugs Prostanoids ~ Latanoprost (continued) Contraindications History of uveilis History of uveilis Prior "incision surgery" or YAG capsulatamy Previous episodes of recurrent HSV ketalitis Relative contraindication is <u>unilateral</u> therapy

Drugs

• Prostanoids

- Travoprost (Travatan Z)
- Average IOP reduction between 7 to 8 mmHg Mean IOP reduction of up to 1.8-2.4 mmHg GREATER in blacks patients
- Rumored to have "slippage" after ~6-12 months
 - (unsubstantiated) Dosage QD "evening" not bedtime

- Prostanoids - Travoprost (continued)

 - Adverse Effects
 Conjunctivalityperemia
 F8 sensation
 Tearing
 Dry Eyes
 Increase diplomentation in iris and periorbital tissue
 Increase diplomentation and growth of evaluates

Drugs

Prostanoids

. .

- Brimatoprost (Lumigan) Synthetic analogue of fatty acid prostamides
 Prostamides are present in ocular tissues

Prostamides presumably lower IOP by the same mechanism as prostaglandins

Dosage QHS

Drugs

- Prostanoids - Brimatoprost (continued) Adverse Effects Adverse Effects – Conjunctival hyperemia – FB sensation – Growth and darkening of eyelashes » Latisse – Pigmentation of periocular skin

Drugs

Prostanoids

.

- Tafluprost (Zioptan) FDA approved for the treatment of OHTN & POAG Feb.
 2012

• *per Merck

Average IOP reduction at 3 months -6 to -8 mmHg
 Average IOP reduction at 6 months -5 to -8 mmHg
 Adverse Effects
 Conjunctival Hyperemia (~4-20%) *

Drugs

Prostanoids

- Unoprostone (Rescula) Originally on market 2000

Reintroduced 2012-2013
 Discontinued March 31, 2015

Drugs

 Prostanoids VESNEO (latanoprostene bunod ophthalmic solution 0.024%)

Per B&L-Upon instillation in the eye, is tan proteine burnot is rapidly made, third do byte at other is the second strategies of the second strategies the second strategies and the second strategies and the second strategies acy role in (for equilation in head year), second strategies to burnot strategies acy second humor outflow by secing so both the two scheral (non-conventions) adverses humor outflow by secing so both the two scheral (non-conventions) adverses in humor scheral burnot scheral the scheral scheral scheral scheral scheral scheral scheral scheral scheral burnot scheral scheral

Awaiting FDA approval

The FDA has set an action date of July 21, 2016 to complete its review, as per the Prescription Drug User Fee Act (PDUFA)

Drugs

 Combination Formulations -- Cosopt

timolol 0.5% & dorzolaminde 2%
Dosed BID

Osee biD
 As effective as timolol 0.5% BID & dorzolomide 2% TID
 Mean IOP changes similar to Xalatan
 – IOP reduction "25-35%
 Contraindications are the sum of each drug

Drugs

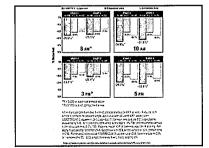
Combination Formulations

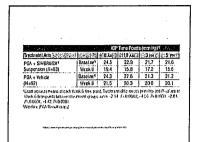
– Combigan

brimonidine 0.2% + timolol 0.5%
NOTE:not Alphagan P (either 0.15% or 0.1%)
Dosed q12h (*BID*)
As effective as both meds given separately

- Better tolerated than 0.2% brimonidine TID
 Better tolerated than Cosopt

- Combination Formulations
- Simbrinza Brinzoloamide 1.0%/Brimonidine 0.2%





Drugs

Combination Formulations

- Xalacom and Extravan Xalacom and Extravan • Latnoprost or travoprost combined with timolol 0.5% • Dosing schedules not clarified at this time • FDA approval delayed for "Xalacom" – Althoughtas bera available in Europe for years... • 10P reduction may equal to, or less than, if separate drugs used together • Extravan may be closer to FDA approval

Conclusion

- Managing IOP remains a staple in glaucoma management
- Consider compliance and quality of life, an the frequently overlooked, cost
- · "Above all else, do no harm" Manage appropriately utilizing all therapeutics available
 Do not let ANYONE go blind in your chair
 * Refer whenever appropriate and/or necessary

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Work: (951) 654-0803 x 2280 jpruitt@rsbcihl.org Cell: (909) 721-7751 pruitt.joseph@gmail.com

Joseph A. Pruitt, O.D., M.B.A., FAAO

Objective:

Objective:		• •
		· · · · · ·
Education:	Nova Southeastern University, Fort Lauderdale-Davie, Florid	a 2008-2011
	Master of Business Administration, 2011	. 2008-2011
	West Los Angeles Veteran Affairs Healthcare Center, Los Ang Residency Certificate, Geriatric/Primary Care, 2008	celes, California 2007-2008
	Illinois College of Optometry, Chicago, Illinois	2003-2007
•	Doctor of Optometry, 2007	
• • • •	California State Polytechnic University, Pomona, California Bachelor of Science, Biology, 2003	2000-2003
	University of Memphis, Memphis, Tennessee	1999-2000
•	Major in Biology	
Licenses:	Tennessee #2753	Date of Issue: July 10, 2007
	Active	Date of issue. July 10, 2007
	Injectible Certification	
	Therapeutic Certification	•
•	California #13429T	Date of Issue: Sept. 28, 2007
	• Active	Date of issue. Sept. 28, 2007
	Therapeutic and Pharmaceutical Agent + Lacrima	l Irrigation
	and Dilation + Glaucoma (TLG) Certified	
	Georgia #OPT002454	Data officerson from 10,0000
	• Active	Date of Issue: June 12, 2008
	 Diagnostic and Therapeutic Pharmaceutical Agen 	t Certified
		· · ·
	Minnesota #3130	Date of Issue: June 17, 2008
•	Active Diamontin Diamontine 1 Amont (DDA) Conting 1	
· .	 Diagnostic Pharmaceutical Agent (DPA) Certified Therapeutic Pharmaceutical Agent (TPA) Certified 	
Board Certi		•
	American Board of Certification in Medical Optometry	Date of recertification: Feb 2018
	Board certified	
Certificatio	ns:	
	Drug Enforcement Agency (DEA) Certified	Date of Expiration: Mar 2020
• • •		· · ·
	Cardiopulmonary Resuscitation (CPR) & Automated External Defibrillator (AED)	Recommended Renewal: Mar 2017
		Recommended Renewal, mai 2017
		ate of Issue/Completion: April 6, 2006
	Certification Number: 20060406002	

		-
¢	Paragon Corneal Refractive Therapy (CRT) Date of Issue/Co • Certification Number: 161000	ompletion: Dec. 28, 2007
	Advance Competence in Medical Optometry (ACMO)	te Taken: June 13, 2008
	Administered by the National Board of Examiners in Optometry (NBEO)	
	 Examination only made available to candidates meeting specific clinical experience requirements/pre-requisites 	. ·
	Passed examination	
Employmen		
•	Riverside San Bernardino County Indian Health, Inc (RSBCIHI)	Oct. 2014- present
	Director of Eye CareStaff Optometrist	• •
	Riverside San Bernardino County Indian Health, Inc (RSBCIHI)	July 2014- Oct. 2014
· .	Staff Optometrist	5 uly 2014- Oct. 2014
	Minneapolis Veteran Affairs Health Care System	Nov 2008- June 2014
	Low Vision/Staff Optometrist	
	Optometric Residency Coordinator	
· •	• Spearheaded and implemented program	
. •	 Student Externship Coordinator Spearheaded and implemented program 	
	o spearmaned and implemented program	•
•	Wal-Mart Vision Center (Red Wing & Rochester, MN)	Jul 2008- Nov 2008
	Associate Optometrist	• • • • • • • • • • • • • • • • • • •
	EyExam of California	Oct 2007- June 2008
•	On-call/Fill-in Optometrist	000 2007 - 0 0110 2008
Faculty App		Inn 0015 measure
	Western University of Health Science / College of Optometry, Pomona, California	Jan 2015 - present
	Clinical Assistant Professor of Optometry	· · ·
	• RSBCIHI Externship Site Program Director	
	o As part of being RSBCIHI Eye Care Director	, ,
	University of the Incarnate Word-Rosenberg School of Optometry,	
	San Antonio, Texas	May 2012- June 2014
	Clinical Assistant Professor	
	Minneapolis VA HCS Externship Site Program Director	•
	Midwestern University-Arizona College of Optometry, Glendale, Arizona	May 2012- June 2014
• •	Adjunct Clinical Assistant Professor	May 2012- 0 UIIC 2014
	Minneapolis VA HCS Externship Site Program Director	
	Southern College of Optometry, Memphis, Tennessee	Dec 2010- June 2014
	Adjunct Faculty	
	Minneapolis VA HCS Externship Site Program Director	
	University of Missouri, St. Louis College of Optometry, St. Louis, Missouri	Jul 2009- June 2014
.*	Adjunct Assistant Professor	
•	Minneapolis VA HCS Externship Site Program Director	
Experience:		
mihorronee.	Riverside-San Bernardino Indian Health, Inc	Oct 2014 - present
	Director of Eye Care	
	o Oversee all organizational Eye Care activities	
•		

Staff Optometrist

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Riverside-	-San Bernardir	io Indian Health, Inc		Jul 2014 – Oct 2014
•	Staff Optome	trist		······································
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Minneapo	lis Veteran Aff	airs Medical Center		Nov 2008- June 2014
•	Staff Optome			
		ry Eye Care		
	o Low V			
•	1 1104 0	Sole low vision eye care	provider	
	o Polytr	auma/Traumatic Brain I		h & Vision
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•		ssociated Health Educati		
		Missouri, St. Louis Colleg		
	University Mi	chigan College of Optome	try, & Southern College	of
•	Optometry for	r the optometric externsh	ip program	•
•		iship program director		•
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	Established A	ssociated Health Educati	on Affiliation Agreement	, t with
-		ollege of Optometry for the		
•		ency in Primary Care/Bra		
		ency program director	in injury and vision iter	haomitauon
	o Reside			
•	: .	Designed the program's		
	· · ·	Secured all necessary ap		
	· •	After the initial site visit	, program received full A	ACOE accreditation
•				
			•	
Wal-Mart	Vision Center (Red Wing & Rochester, M	IN)	Jul 2008- Nov 2008
•	Associate Opt	ometrist	· · · · · · · · · · · · · · · · · · ·	
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Residency	•			
		Veteran Affairs Healthca	re Center	Jul 2007- June 2008
	-	rics/Primary Care		
	0		Diabetic exams	•
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		 Given during Ch 		· · · · ·
• •	· · · · O	Precept Southern Califor	mia College of Optometr	'y's
•		interns		•
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Ontometri	c Externships:			
		tute, Jacksonville Beach,	FT	Feb-May 2007
<i></i>				1.00-Way 2007
		D private practice with a		· · ·
		ct Lenses and Primary Ca		
	 Obser 	ved multiple surgical proc	edures:	
	· 0	Cataract Extraction		
	. 0	Blepharoplasty		•
	0	Strabismus recession ar	1d resection	
	Ū			
<i>እለ</i> ራ	mnhis Veteran	s Affairs Medical Center (VAMC), Memphis TN	Nov 2006-Feb 2007
ί Α ΤC			······································	1107 2000-1.60 2007
		asis on Primary Care		
	 Assist 	ed in direct care in a high	. patient volume	
•				•
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	•	25		

	c eye clinic

- Assisted in optometric injections and fluorescence
 - angiographies procedures

Illinois Eye Institute (IEI), Chicago, IL

- Emphasis on Pediatrics/Binocular Vision, Advance Care, and Low Vision
 - Performed comprehensive eye exams on pediatric patients (infants-11yrs of age)
- Performed comprehensive eye exams on "at risk/2nd chance" children one day a week at Maryville Academy
- Constructed, tailored and performed successful binocular vision/vision therapy treatments to 4 children over a 10 week period
- Assisted in the treatment of advance glaucoma with attending University of Chicago ophthalmologist
- Performed problem specific examinations one day per week in IEI's Emergency/Urgent Care/Walk-in clinic
- Performed full Low Vision examinations including Low Vision device selection and training

Body of Christ Optometry Clinic, Tegucigalpa, Honduras

May-Aug 2006

Aug-Nov 2006

- Emphasis on Primary and Advance Care Performed full-scope optometric care in a high
- patient volume medical clinic geared towards the underprivileged
- Also worked closely with a local ophthalmologist
 - o Observed and assisted in Cataract Extraction
 - and Incision and Curettage procedures
 - Provided pre and post-surgical care

Primary Care Clinical Education Illinois Eye Institute, Chicago, IL

Aug 2005-May 2006

Jun-Aug 2004

Volunteer Optometric Assistant

Body of Christ Optometry Clinic, Tegucigalpa, Honduras

• Assisted staff optometrist in direct patient care in the clinic and multiple remote satellite outreach locations

Professional	
Affiliations/Member	ships:

- Accreditation Council on Optometric Education o Consultant, 2014-present
- American Academy of Optometry (AAO) o Fellow; Class of 2009
- American Optometric Association (AOA)
- Armed Forces Optometric Society (AFOS)
- European Academy of Optometry and Optics (EAOO) o Candidate for Fellowship
- Fellowship of Christian Optometrists (FCO)
- Minneapolis VAMC Medical Staff Association
 - o Steering Committee, member 2010-2014
- National Association of Veteran Affairs Optometrists (NAVAO)
 Newsletter Committee, member 2010-2014
- National Optometric Association (NOA)
 - o Minnesota's NOA State Representative 2010-2012
 - o National Optometric Student Association (NOSA)
 - NOSA National Vice-President: 2006-2007
 - NOSA-ICO President: 2005-2006
 - NOSA-ICO Vice-President: 2004-2005

• Volunteer Optometric Service to Humanity (VOSH)

Journal of Rehabilitation Research and Development

o Peer Reviewer, 2013-2014

Activities:

- VOSH Medical Mission Trip, Bamenda, Cameroon (May 2010)
- Mayo Medical School/Brighter Tomorrow's Winter Warmth Festival (Jan 2009 & Jan 2010)
 - Fun day of activities for children battling cancer and their families
 Volunteer
- Veteran Affairs Disaster Emergency Medical Personnel System (DEMPS) o Volunteer (Aug 2009-present)
- FCO Optometry Mission Trip, Port Au Prince, Haiti (Feb 2007)
- SVOSH Medical Mission Trip, Addis Addaba, Ethiopia (Mar-Apr 2006)
- FCO Optometry Mission Trip, Tegucigalpa, Honduras (Apr 2003 & Nov 2004)

Honors/Rewards:

- Recognition of Excellence in Teaching as Clinical Assistant Professor, Western University Health Sciences/College of Optometry (2015-2016 Academic Year)
- Nomination for Medical Staff Clinical Excellence Award, (2012 & 2013)
- Recognition for Outstanding Dedication and Service as Adjunct Assistant Professor, University of Missouri – St. Louis (2010-2011 Academic Year)
 - Journal of the American Optometric Association: Optometry's Eagle Award (Nov 2010)
- Certificate of Appreciation (July 2009)
 - Department of Veterans Affairs VISN 23
 - Awarded for participation in VISN 23 Blind and Low Vision Continuum of Care Conference
 - Recognition for Clinical Excellence (May 2007)
- Derald Taylor Low Vision Award (May 2007)
- Clinical Dean's List (summer 2005; summer & fall 2006, winter & spring 2007)
- Academic Dean's List (fall 2004)
- Wildermuth Leadership Award/Scholarship (Aug 2006)
- Vistakon Acuvue Eye Health Advisor Citizenship Scholarship (Jan 2006)
- NOSA Service Award/Scholarship (Aug 2004)

Publications:

Pruitt JA. The Management of Homonymous Hemianopsia Secondary to Hemispheric Ischemic Cerebral Vascular Accident. Accepted for publication by Review Optometry (July 2010)

Rittenbach TL, Pruitt JA. A Roundup of Recently Approved Ophthalmic Drugs (and their Use in Practice.) Rev Optom. 2014. 151(2):22-28.

Pruitt JA. Management strategies for patients with AION. Rev Optom. 2011. 148(6):57-65.

Pruitt JA. Neuro-Optometric Rehabilitation Association Program Summary. Optimum VA: The Official Newsletter of the National Association of VA Optometrists Summer 2010.

Pruitt JA, Ilsen P. On the frontline: What an optometrist needs to know about myasthenia gravis. Optometry 81(9): 454-460.

Pruitt JA, Sokol T, Maino D. Fragile X Syndrome and the Fragile X-associated Tremor/Ataxia Syndrome. Eye Care Review: Ophthalmology, Optometry, Opticianry 4(2): 17-23

Posters/Presentations

Pruitt JA. The Curious Case of the Functionally Legally Blind Patient with 20/25 (6/7.5) Visual Acuity. Accepted into American Optometric Association Annual Meeting: Optometry's Meeting (2012) Poster Session.

Pruitt JA, Prussing N. Successfully Treated Horizontal Diplopia Returns with Subsequent Traumatic Brain Injury. Accepted into American Optometric Association Annual Meeting: Optometry's Meeting (2012) Poster Session.

Pruitt JA, Prussing N. The Curious Case of the Functionally Legally Blind Patient with 20/25 (6/7.5) Visual Acuity. European Academy of Optometry and Optics Annual Meeting (2012) Poster Session.

Pruitt JA, Prussing N. Successfully Treated Horizontal Diplopia Returns with Subsequent Traumatic Brain Injury. European Academy of Optometry and Optics Annual Meeting (2012) Case Presentation Session.

Pruitt JA, Prussing N. Traumatic Brain Injury Resulting in Horizontal Diplopia Resolved 5 Years Later with 12 Weeks of Vision Therapy. Minnesota Optometric Association Annual Meeting (2012) Poster Session.

Pruitt JA, Wiley LM. Overcoming Mental Barriers in Visual Rehabilitation. American Optometric Association Annual Meeting: Optometry's Meeting (2011) Poster Session.

Pruitt JA, Prussing N. Traumatic Brain Injury Resulting in Horizontal Diplopia Resolved 5 Years Later with 12 Weeks of Vision Therapy. European Academy of Optometry and Optics Annual Meeting (2011) Poster Session.

Pruitt JA. Overcoming Mental Barriers in Visual Rehabilitation. European Academy of Optometry and Optics Annual Meeting (2011) Case Presentation Session.

Pruitt JA, Wiley LM. Overcoming Mental Barriers in Visual Rehabilitation. Minnesota Optometric Association Annual Meeting's (2011) Poster Session

Pruitt JA, Ilsen P, Yeung C. Ptosis Crutch: Success Treating Myogenic Ptosis Secondary to Myasthenia Gravis. American Optometric Association (AOA) 2008 Optometry Meeting Poster Session

Pruitt JA, Ilsen P. Ptosis Crutch: Success Treating Myogenic Ptosis Secondary To Myasthenia Gravis. Southeastern Congress of Optometry (SECO) 2008 Multimedia Poster Session

Lectures and Other:

Riverside-San Bernardino County Indian Health, Inc.: Eye Care Rounds (Nov 2016)

Ptosis Crutch: Success Treating Myogenic Ptosis Secndary to Myasthenia Gravis

• CA Board of Optometry-approved CE

Riverside-San Bernardino County Indian Health, Inc.: Eye Care Rounds (Sept 2016)

- Visual Fields
 - CA Board of Optometry-approved CE

Riverside-San Bernardino County Indian Health, Inc.: Eye Care Rounds (July 2016)

• Ethical Concerns with Short-term Mission Trips

• CA Board of Optometry-approved CE

Riverside-San Bernardino County Indian Health, Inc.: Eye Care Rounds (July 2016)

- Systemic Urgencies and Emergencies
- CA Board of Optometry-approved CE

Riverside-San Bernardino County Indian Health, Inc.: Eye Care Rounds (Mar 2016)

- Episcleritis, Scleritis, and Iritis
- CA Board of Optometry-approved CE

Illinois College of Optometry: Practice Opportunities Symposium (Mar 2011)

- Represented and presented on VA Optometry
- Participated in panel discussion on "Residency-trained Optometrists"

University of Minnesota: Pre-Optometry Club (Oct. 2010)

- Presentation on the profession of Optometry
- Presented and represented VA Optometry and NOA

Illinois College of Optometry: Capstone Ceremony (May 2010)

 Represented and presented on VA Optometry

Illinois College of Optometry: Practice Opportunities Symposium (Mar 2010)

- Participant in Residency-trained Speaker's Panel
- Represented and presented on VA Optometry

Illinois College of Optometry: White Coat Ceremony/Smart Business Program (Sept 2009)

Participant on Recent Graduate Speaker's Panel