Periocular Triamcinolone vs. Intravitreal Triamcinolone vs. Intravitreal Dexamethasone Implant for the Treatment of **Uveitic Macular Edema**

doi:10.1016/j.ophtha.2018.08.021

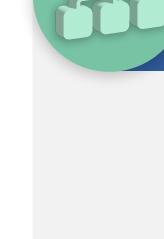
comparative trial.

Thorne JE, Sugar EA, Holbrook JT, et al. Ophthalmology. 2019;126:283-295.

Macular edema (ME) is a common structural ocular complication of uveitis responsible for a substantial amount of visual impairment among patients with uveitis. Despite a wider availability of newer classes of medications used to treat uveitis, the frequency of ME has been relatively stable. Uveitic ME that persists despite control of the uveitis is typically treated with adjunctive regional corticosteroid injections, which may be delivered via a periocular or intravitreal route. There have been, however, limited comparative trials of the periocular and intravitreal corticosteroid therapies. The PeriOcular versus INTravitreal corticosteroids for the treatment of uveitic macular edema (POINT) Trial

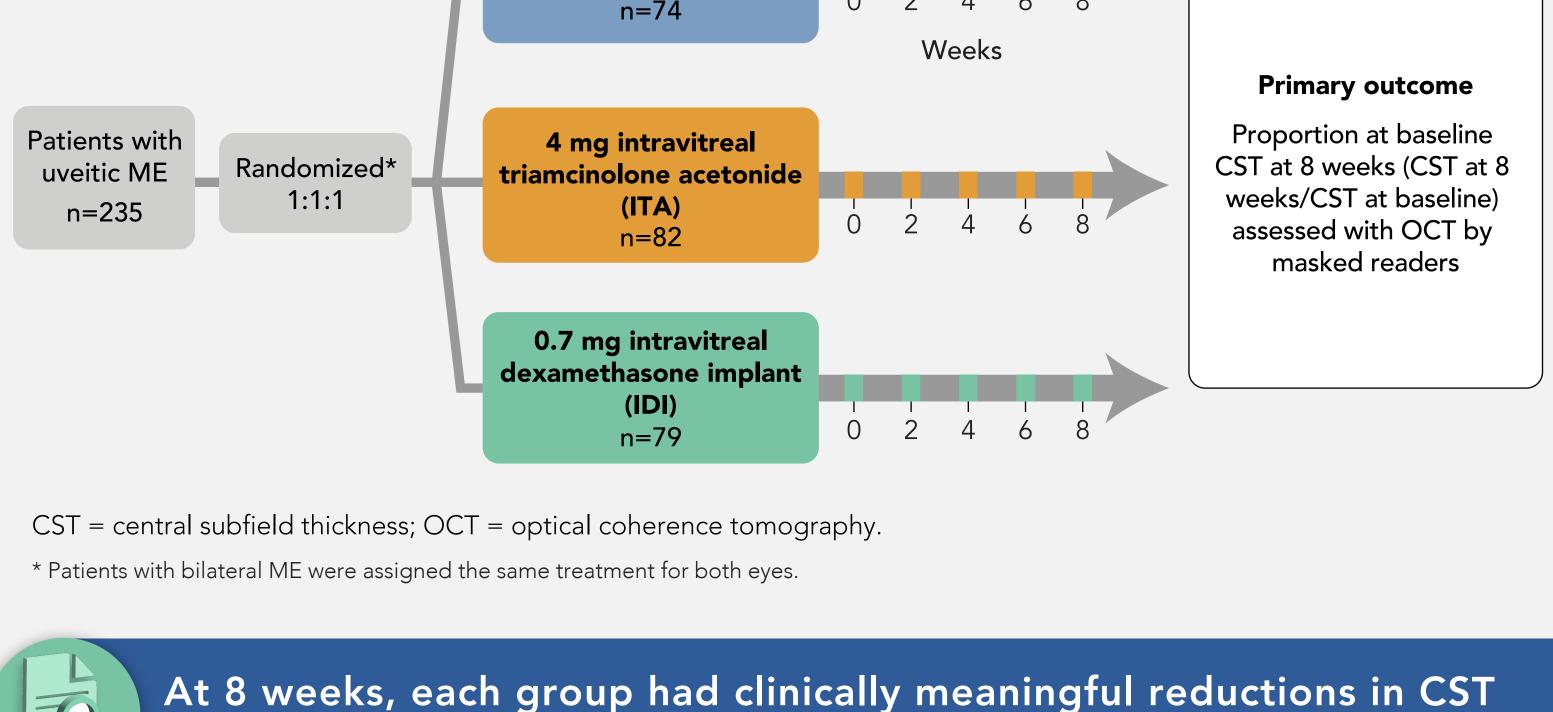
dexamethasone implant (IDI) in the initial treatment of uveitic ME, to evaluate their comparative effectiveness.

compared periocular triamcinolone acetonide (PTA), intravitreal triamcinolone acetonide (ITA), and intravitreal

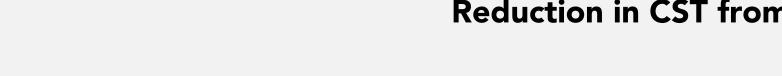


40 mg periocular triamcinolone acetonide (PTA)

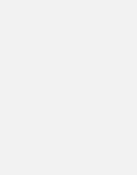
The POINT Trial was a multicenter, randomized, parallel-treatment,



Reduction in CST from baseline at 8 weeks



relative to baseline.



PTA

ITA

0.61 39%

23%

0.65

0.69

8.0

1.0

The IDI group also had a BCVA improvement of

5 letters greater than the PTA group at week 8.

0.56

HR (99.87% CI)

0.6

HR (99.87% CI)

10-

8

6-

2-

Although all treatment groups demonstrated improvement in best-corrected

had improvements in BCVA that was greater than the PTA group at 8 weeks.

visual acuity (BCVA) compared to baseline BCVA, the ITA and IDI groups

0.79

0.86

0.8

0.96

P< .0001

0.9

P<.0001

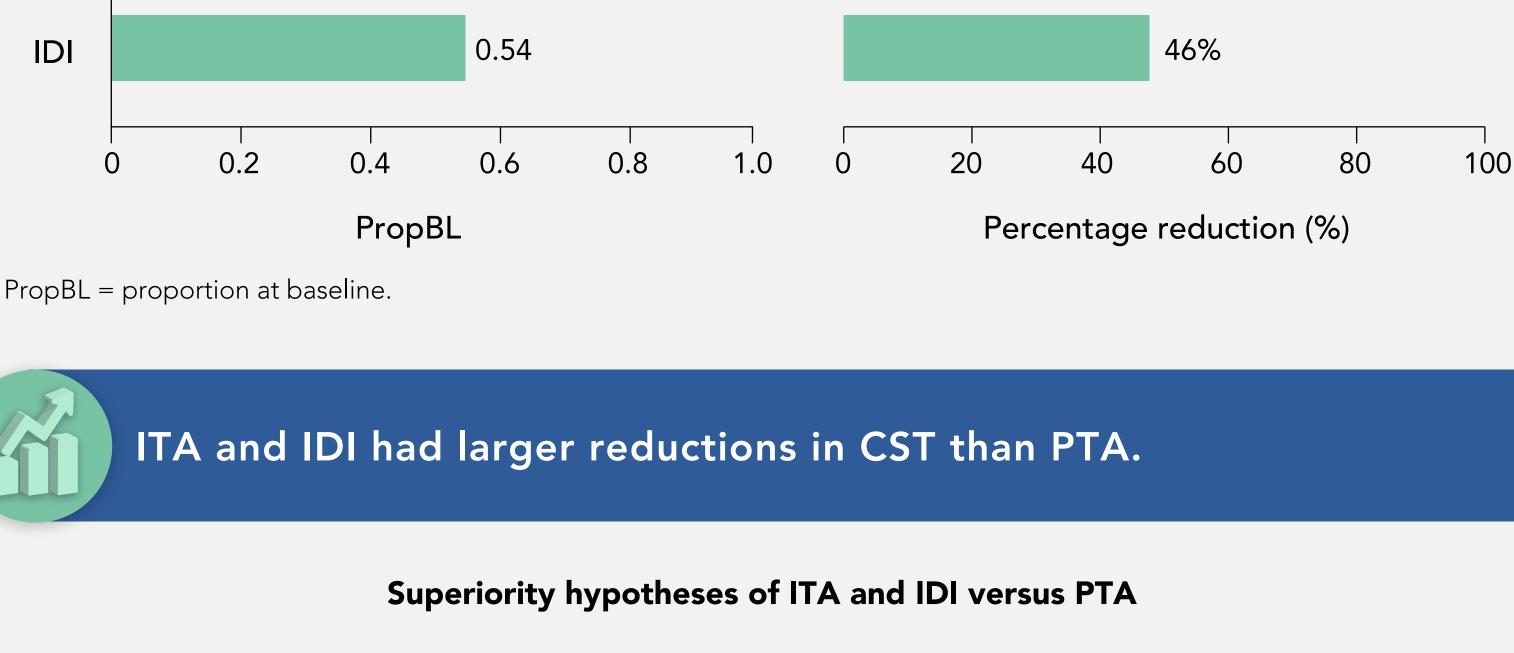
1.0

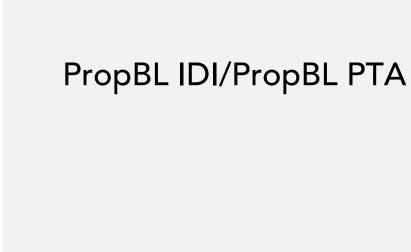
1.2

-5 letters greater

P< .004

0.77

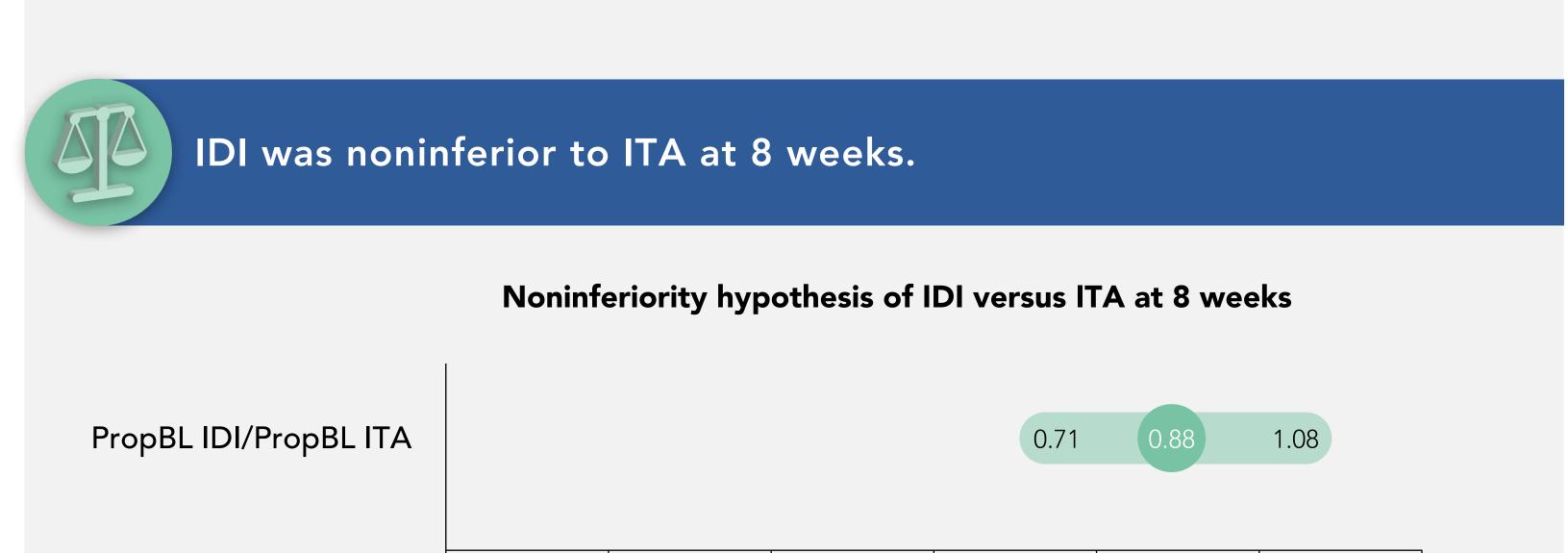




PropBL ITA/PropBL PTA

HR = Hazard ratio; CI = Confidence interval.

0.1 0.2 0.5 0.7 0.3 0.4 0.6 0



0.4

0.2

-5 letters greater

P< .003

0.91

1.0

improving and resolving uveitic ME.

Primary outcome interval

0.5

1.83

2.0

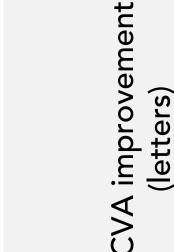
1.29

1.5

0

The ITA group had a BCVA improvement of 5 letters

greater than the PTA group at week 8.



10-

8

6

2

ITA/PTA

IDI/PTA

100 -

Eyes (Periocular)

Eyes (Intravitreal)

Eyes (Dexamethasone) 79

20

10

74

80

Eyes (Periocular)

Eyes (Intravitreal)

Eyes (Periocular)

Eyes (Intravitreal)

Eyes (Dexamethasone) 79

74

82

Eyes (Dexamethasone) 79

100 -

90-

80-

74

80

0 ITA PTA IDI PTA

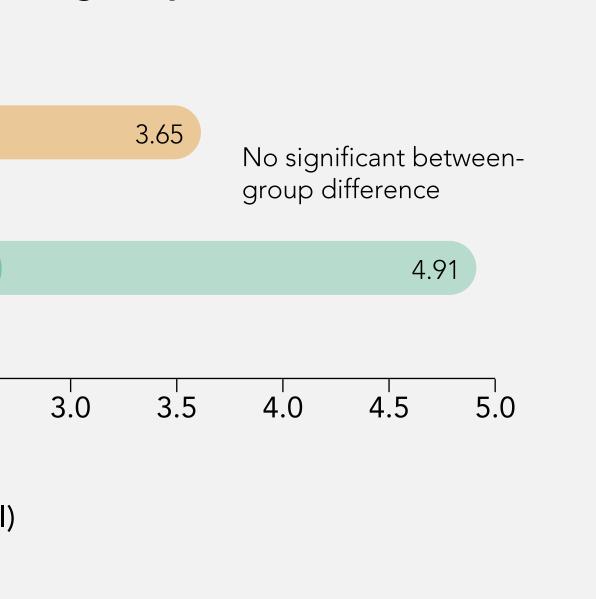
The risk of having intraocular pressure (IOP) ≥24 mmHg was higher in the

intravitreal treatment groups compared with the periocular group; however,

there was no significant difference between the 2 intravitreal treatment groups.

Risk of having IOP ≥24 mmHg compared to PTA

improvement



Follow-up interval

24

72

81

75

24

72

81

75

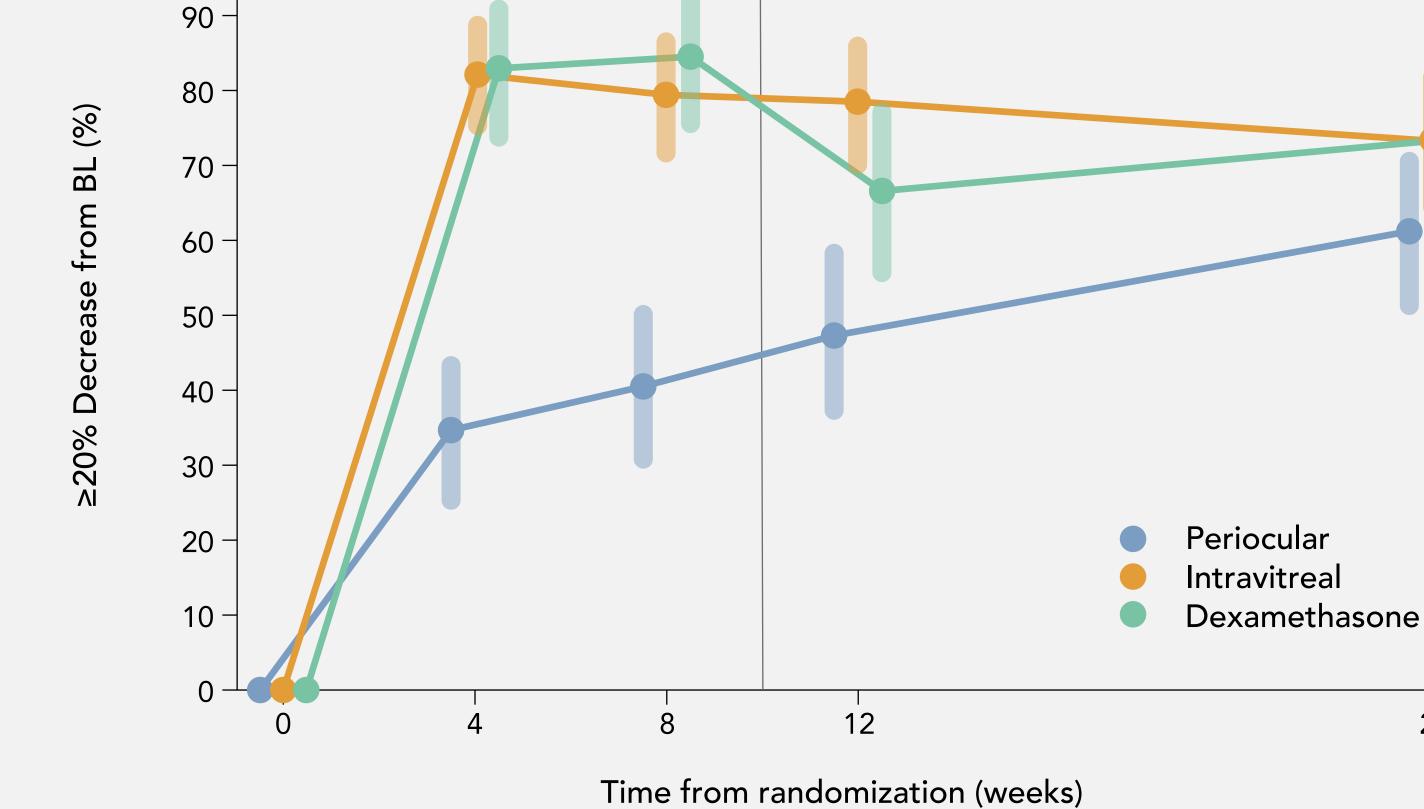


HR (95% CI) Both ITA and IDI treatments were superior to PTA treatment in

Proportion of eyes with improvement in ME

2.52

2.5



74

75

73

71

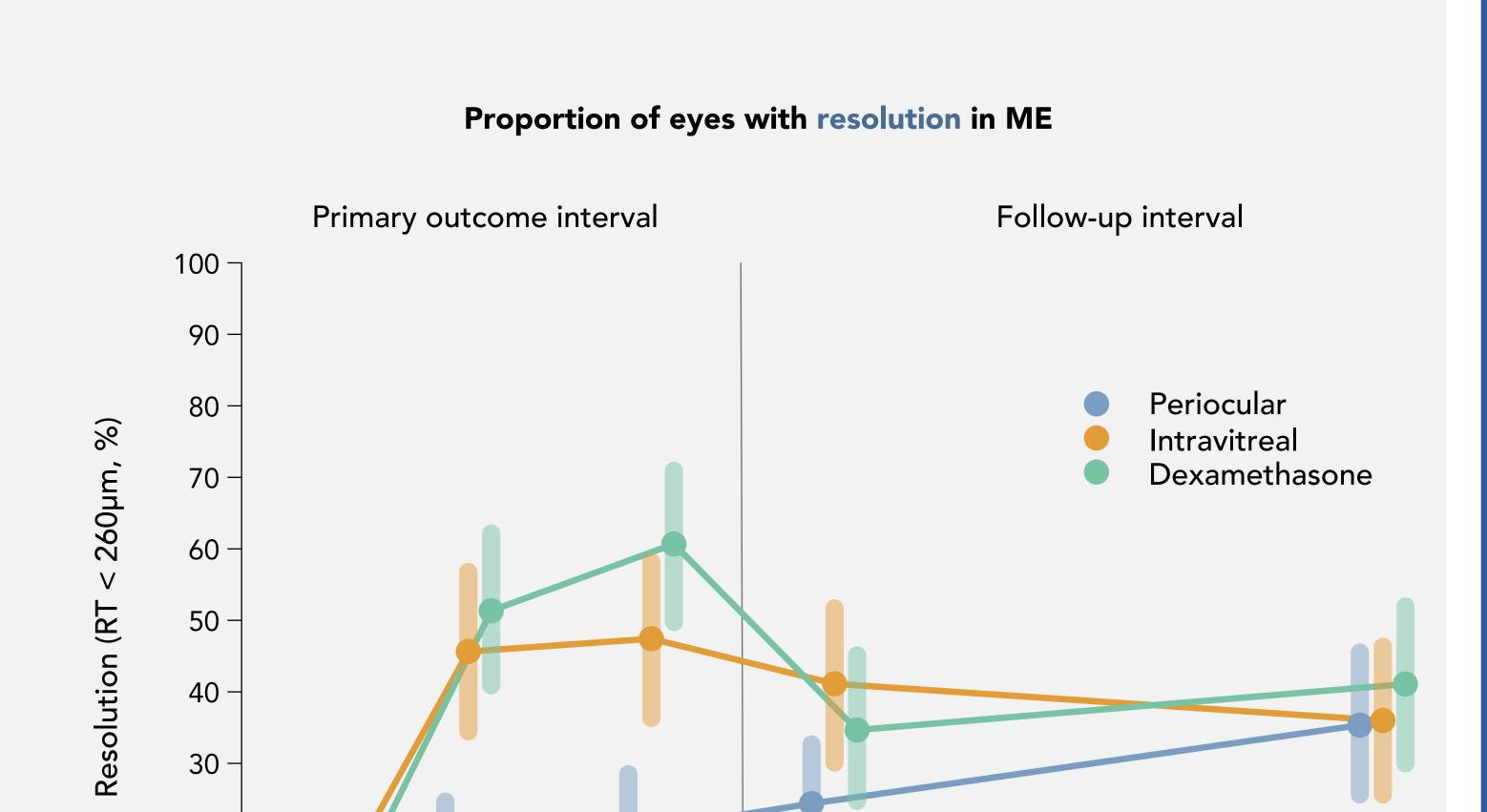
77

73

72

81

74



There were no significant differences in the use of IOP medications among the 3 treatment groups at any time.

Use of IOP medication at each visit

12

71

77

73

Follow-up interval

Periocular

Intravitreal

72

80

75

Dexamethasone

24

72

81

75

Time from randomization (weeks)

8

74

75

73

4

72

81

74

Primary outcome interval

P≥ .14 IOP medication use (%) 70 -60 50 40 30 20 10 0 8 12 20 4 0

74

77

76

72

81

76

Conclusions		

Both intravitreal triamcinolone acetonide and the intravitreal dexamethasone

implant were superior to periocular triamcinolone acetonide for treating uveitic

ME, with modest increases in the risk of IOP elevation.

Time from randomization (weeks)

72

77

73