Predictions of Residual Astigmatism from Surgical Planning for Topographic-Guided LASIK Based on Anterior Corneal Astigmatism (LYRA Protocol) vs the Phorides Analytic Engine

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Purpose: To identify the laser programming strategy that achieves the highest refractive astigmatism outcomes accuracy between LYRA surgical planning and Phorides surgical planning.

Methods: One hundred one eyes successfully treated with Contoura with LYRA Protocol were retrospectively surgically planned with Phorides to determine its accuracy. Eighteen eyes that had inaccurate 3 month outcomes with LYRA were also retrospectively planned with Phorides. Two patients with one eye treated with LYRA and one with Phorides surgical planning were analyzed for outcomes.

Results: Retrospective Phorides surgical planning deviated from already successful treatments with LYRA Protocol 51% of the time, with an average astigmatic deviation of 0.69 diopters (D) and an average deviation of 7.1 degrees. The percentage of eyes of the 101 successfully treated with LYRA Protocol then planned with Phorides differed by 0.5D in 19.8%, by 0.75D in 23.8%, and by 1D in 7.9%. A retrospective analysis with Phorides of patients treated with LYRA Protocol with residual post-operative astigmatism demonstrated that Phorides would have increased accuracy in only 33% of patients, and not helped or decreased accuracy in the rest.

Conclusion: Phorides was significantly less accurate in surgical planning than the LYRA Protocol overall and a 51% of primary patients could have a likelihood of significant residual astigmatism. Phorides would also have provided a less accurate outcome in the majority of patients that needed secondary enhancement.

Keywords: Contoura, topographic-guided ablation, LYRA Protocol, Phorides, uniform cornea

Introduction

The advent of topographic-guided ablation has brought with it a discussion on the most accurate way to plan surgically for accurate refractive outcomes while reducing light scattering higher order corneal aberrations. The initial protocols in the FDA study for Alcon WaveLight (Wavelight Inc, Erlangen, Germany) Contoura (the market name for the Alcon WaveLight topographic-guided ablation system) approval used the manifest refraction, but patients were chosen for which the Contoura measured astigmatism did not deviate dramatically from the manifest refraction.¹ The authors published in 2017 the first protocol to create a more

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uniform cornea by using the Contoura measured astigmatism to treat the anterior corneal elevation, the LYRA Protocol (Layer Yolked Reduction of Astigmatism).¹⁻³ Although Wallerstein et al published a study showing manifest refraction obtained good results, the study itself has issues with how it arbitrarily divides the data, does not address the topographic consequences of utilizing manifest refraction, the subjective patient quality of vision outcomes, and directly contradicts the vector astigmatism analysis study that demonstrates that manifest refraction does not work as well.

Subsequently, the Phorides Analytic Engine (Mark Lobanoff, MD Minneapolis, Minnesota) was created and is now being supported by Alcon, Inc for use in surgical planning with Contoura.⁴⁻⁷ Phorides takes the corneal higher order aberration map from the Contoura surgical planning software, re-analyzes it, requires the entry of the posterior corneal astigmatism from the Pentacam or the Gallelei, and then use other proprietary algorithms to calculate posterior ocular astigmatism which is combined with the refractive effect of the anterior HOA map in an attempt to create a more accurate refractive outcome. As Dr. Lobanoff describes in his paper. The Phorides Analytic Engine thus considers the anterior corneal astigmatism, the topographic abnormalities that create higher order aberrations, the posterior corneal astigmatism, and the lenticular astigmatism ...⁷ It is an intriguing attempt to simulate in a surgical planning engine a similar goal to Ray Tracing technology (which is a different methodology utilizing hardware and software that attempts to correct all anterior or posterior astigmatism/aberrations that to focus a point of light perfectly on the retina), to correct ocular astigmatism posterior to the anterior corneal surface. This correction takes place on the anterior corneal surface in both methodologies.

To determine the accuracy of these surgical planning algorithms for clinical outcomes, this study retrospectively examined 3 categories of eyes: 101 eyes treated with Contoura with LYRA Protocol and retrospectively planned with Phorides; 18 eyes that had inaccurate outcomes at 3 months with LYRA and retrospectively surgically planned with Phorides, and a small sample of 2 patients with 1 eye performed with LYRA and 1 eye with Phorides which was part of a study aborted due to results.

Materials and Methods

Patient data were retrospectively analyzed if they had myopic/astigmatic LASIK with Contoura with the LYRA

Protocol (Table 1) and at least 3 month results had been obtained for stability.¹⁻³ Patients who had greater than or equal to 0.50 diopters (D) of difference between manifest and Contoura measured astigmatism were selected. This difference of 0.5D would elucidate the differences between LYRA and Phorides as patients with a small difference seem to result in surgical planning too close to elucidate differences between the two protocols. Each eye of each patient was required to have a plano result (in the case of a few eyes nearly plano but without subjective visual complaints) after treatment with Contoura with LYRA Protocol, and the Phorides software was used for surgical planning in these patients retrospectively.

LYRA is an acronym for Layer Yolked Reduction of Astigmatism.¹⁻³ This is a recognition of the fact that the Aberration Removal Layer is yolked/linked to the Refraction Correction layer. Contoura calculates the results of that linkage. The protocol is as follows:

1. Enter the manifest/cycloplegic refraction into Contoura during Pre-surgical planning.
2. Zero out the astigmatism and sphere to see ablation pattern for the Aberration Correction Layer.
3. Enter Contoura measured astigmatism and axis for the final correction. The ablation map at this point should be similar to the Pentacam anterior elevation map. This will assist understanding the ablation when there is a significant discrepancy between manifest vs measured astigmatic power and axis.
4. The sphere is now entered after adjustment for the spherical equivalent of the change in astigmatism.

The large difference between manifest and measured astigmatism (0.50 D or greater) was chosen to create a situation that accurately was able to demonstrate the outcomes of the different forms of planning, and to create a “stress test” for the Phorides software to determine how the addition of calculated posterior data changed the outcome of the procedure. One hundred one (101) eyes were included in this part of the study, and each of these eyes had already been treated with Contoura with LYRA Protocol and had resulted in a successful result (ie, plano or within 0.5 D of a plano result). In addition, it was necessary for each patient to also be subjectively satisfied with their vision and have no subjective complaints. We have noted in patients who have more uniform corneas that the quality of the vision results in patients noting even small corrections or visual flaws. They rarely tolerate even

Table I Difference (in Diopters) Between Refractive Outcomes with Contoura Using the LYRA Protocol versus the Phorcidex Software

Manifest (Rx Wet)	LZ TX(LYRA)	PHORCIDES	Target Goal	Post-Op RX	VA	Cyl Difference	Axis Difference
-3.50-1.25x105	-3.69-0.87x173	-3.89-0.77x164	plano	plano	20/15	0.1	9
-2.75-1.00x100	-2.90-0.71x159	-2.22-0.37x147	-1.00	-1.00	20/30 J1 +	0.34	12
-1.00	-0.54-0.93x002	-1.01-0.18x002	plano	plano	20/15	0.74	0
-1.00	-0.40-1.21x005	-0.95-0.30x012	plano	plano	20/20	0.91	7
-1.50-0.75x105	-1.79-0.17x132	-1.76-0.53x137	plano	plano	20/20	0.36	5
-2.25-1.00x100	-2.75-1.00x100	-2.54-0.62x104	plano	plano	20/15	0.53	4
-1.00-1.25x098	-1.44-0.37x092	-1.34-0.78-0.94	plano	plano	20/15	0.41	2
-1.00-1.00x080	-1.17-0.66x032	-1.38-0.43x037	plano	plano	20/15	0.23	5
-4.00-2.75x008	-3.63-3.50x002	-4.10-2.74x006	plano	plano	20/20	0.72	4
-3.50-2.50x003	-3.25-3.00x179	-3.62-2.46x003	plano	plano	20/20	0.71	4
-3.00-1.25x169	-2.60-2.05x176	-3.10-1.25x169	plano	plano	20/15	0.8	7
-4.25-1.00x006	-3.84-1.83x008	-4.43-0.87x013	plano	plano	20/15	0.96	5
-2.50	-2.12-0.77x171	-2.44-0.31x165	plano	plano	20/15	0.46	6
-2.50	-2.00-1.01x015	-2.48-0.24x010	plano	plano	20/20	0.77	5
-4.75-1.50x053	-5.03-0.93x063	-5.43-0.92x023 and WFO plano-0.50x113	plano	plano	20/20	0.49	40
-6.00-1.75x152	-6.41-0.93x150	-6.15-1.21x153	plano	plano	20/15	0.28	3
-1.25	-0.41-1.68x014	-1.06-0.69x016	plano	plano	20/20	0.99	2
-3.50	-3.09-0.83x152	-3.42-0.45x136	plano	plano	20/20	0.38	16
-3.75	-3.40-0.70x001	-3.80-0.01x177	plano	plano	20/20	0.69	4
-5.00	-5.00-0.10x145	-4.95-0.14x034	plano	plano	20/15	0.04	111
-5.50	-5.18-0.64x005	-5.35-0.30x010	plano	plano	20/20	0.4	5
-2.00	-1.43-1.13x172	-1.98-0.25x171	plano	plano	20/15	0.88	1
-1.75	-1.60-0.81x010	-1.98-0.33x015	plano	plano	20/15	0.48	5
-5.75-0.50x090	-5.73-0.55x040	-5.78-0.26x066	plano	plano	20/15	0.29	26
-6.00-0.25x090	-5.67-0.81x165	-5.91-0.30x163	plano	plano	20/15	0.51	2
-1.50	-1.16-0.71x009	-1.56-0.07x019	plano	plano	20/20	0.64	10
-1.25	-0.85-0.82x167	-1.22-0.26x166	plano	plano	20/20	0.56	1
-0.50-1.25x115	-0.79-0.67x140	-0.78-0.99x140	plano	plano	20/15	0.32	0
-0.50-1.25x055	-0.73-0.80x023	-0.76-0.52x041 and WFO plano-0.50x063	plano	plano	20/15	0.22	18
-2.00	-1.75-0.50x006	-1.98-0.24x007	plano	plano	20/15	0.26	1
-2.25-0.25x030	-1.93-0.90x180	-2.22-0.50x001	plano	plano	20/15	0.4	1
-2.50-0.50x080	-2.28-0.95x025	-2.72-0.26x028	plano	plano	20/15	0.69	3
-2.50-0.50x080	-1.38-0.73x161	-0.74-0.33x148	-1.00	-0.75-0.50x100	20/25 J1	0.4	13
-5.25-0.75x105	-5.14-0.97x167	-5.44-0.44x162	plano	-0.25D/S	20/20	0.53	5
-4.50-0.50x145	-4.14-1.22x177	-4.57-0.53x001	plano	plano	20/20	0.69	4
-1.50-2.00x120	-1.78-1.45x133	-1.82-0.93x135 and WFO plano-0.75x109	plano	plano	20/15	23	2
-0.25-3.00x070	-0.82-1.86x061	-0.66-2.37x069	plano	plano	20/20	0.51	8
-9.75-0.50x020	-8.88-0.73x180	-8.51-0.43x001	plano	plano	20/20	0.3	1
-8.00D/S	-7.71-0.58x022	-8.09-0.40x027	plano	-0.50D/S	20/20	0.18	5
-2.00-1.25x090	-2.37-0.52x039	-2.40-0.64x039	plano	-0.50D/S	20/20	0.12	0
-2.00-1.25x085	-2.39-0.48x100	-2.36-0.73x091	plano	plano	20/15	0.25	9
-4.25D/S	-3.82-0.86x004	-4.26-0.21x001	plano	-0.50D/S	20/20	0.65	3
-3.00-0.75x168	-2.53-1.69x167	-3.05-0.95x169	plano	-0.25D/S	20/20	0.74	2
-6.50-1.50x003	-6.29-1.93x175	-6.17-1.78x179	plano	plano	20/15	0.15	4
-3.00-2.50x170	-3.65-2.31x165	-3.36-2.08x173	plano	plano	20/15	0.23	8

(Continued)

Table I (Continued).

Manifest (Rx Wet)	LZ TX(LYRA)	PHORCIDES	Target Goal	Post-Op RX	VA	Cyl Difference	Axis Difference
-1.00D/S	-0.87-0.60x006	-1.02-0.26x015	plano	plano	20/15	0.34	9
-1.25D/S	-0.97-0.57x180	-1.22-0.35x000	plano	plano	20/15	0.22	0
-0.50-3.50x095	-1.05-2.40x099	-0.84-3.12x098	plano	plano	20/20	0.72	1
plano-4.00x090	+0.00-2.46x086	-0.48-3.26x087	-1.00	-0.75D/S	20/20 JI +	0.8	1
-3.75-0.50x175	-3.43-1.15x003	-3.76-0.69x003	plano	plano	20/15	0.46	0
-2.75D/S	-2.28-0.94x001	-2.65-0.43x011	plano	plano-0.50x100	20/20	0.51	10
-4.25-1.00x180	-4.06-1.39x178	-4.37-1.09x178	plano	plano	20/15	0.3	0
-4.25-0.75x175	-3.80-1.66x171	-4.32-0.84x173	plano	plano	20/15	0.82	2
-5.50-1.00x175	-4.99-2.02x004	-5.31-1.48x006	plano	plano	20/15	0.54	2
-6.00-1.75x165	-5.76-2.23x169	-5.79-2.04x169	plano	plano	20/20	0.19	0
-4.25-1.75x105	-4.62-1.02x120	-4.52-1.54x105	plano	plano	20/20	0.52	15
-4.75-1.50x073	-5.07-0.86x048	-4.96-1.28x059	plano	plano	20/20	0.42	11
-3.75D/S	-3.23-1.05x170	-3.56-0.67x172	plano	plano	20/15	0.38	2
-4.00-0.50x010	-3.83-0.83x003	-4.09-0.52x007	plano	plano	20/10	0.31	4
-1.75D/S	-1.54-0.43x166	-1.74-0.22x169	plano	plano	20/15	0.21	3
-1.75-0.75x157	-1.52-1.21x170	-1.69-1.09x164	plano	plano	20/15	0.12	6
-4.00-1.25x098	-4.63+0.00x098	-4.31-0.83x138	plano	plano	20/15	0.83	9
-4.00-1.25x080	-4.54-0.17x069	-4.26-0.93x080	plano	plano	20/15	0.76	11
-1.50D/S	-1.03-0.94x007	-1.60-0.09x007	plano	-0.25D/S	20/15	0.25	0
-1.25D/S	-1.02-0.46x173	-1.26-0.7x174	plano	plano	20/15	0.19	1
-4.25-3.00x010	-3.88-3.75x002	-4.35-3.14x006	plano	plano	20/15	0.61	4
-2.25-3.25x165	-1.75-4.25x166	-2.52-3.00x165	plano	plano	20/15	1.25	1
-3.75D/S	-3.49-0.53x011	-3.84-0.02x013	plano	-0.25D/S	20/20	0.51	2
-3.75D/S	-3.50-0.49x169	-3.78-0.13x065	plano	plano	20/15	0.36	104
-4.50-1.00x010	-4.14-1.72x003	-4.47-1.22x005	plano	plano	20/15	0.5	2
-3.75-1.00x165	-3.47-1.56x170	-3.76-1.19x169	plano	plano	20/20	0.37	1
plano-4.25x015	+0.00-4.75x011	-0.40-3.74x014	plano	plano-0.50x060	20/25	1.01	3
plano-4.00x160	+0.00-4.75x169	-0.44-3.42x162	plano	plano-0.50x045	20/25	1.33	7
-1.25-0.75x090	-0.50-0.26x046	-1.46-0.54x108	plano	plano	20/15	0.28	62
-1.25-1.00x075	-1.39-0.73x078	-1.36-0.98x072	plano	plano	20/15	0.25	6
-2.25-1.25x005	-1.86-2.02x001	-2.30-1.44x0001	plano	plano	20/15	0.58	0
-3.75-1.25x165	-3.46-1.83x175	-3.68-1.59x169	plano	plano	20/20	0.24	6
-2.50-3.00x035	-2.62-2.77x031	2.86-2.48x034	plano	plano	20/15	0.29	3
-3.50-3.25x147	-2.70-2.84x156	-2.83-2.89x152	-1.00	-1.00,-0.50 x 120	20/15 JI +	0.05	4
-1.25-1.00x115	-1.44-0.62x137	-1.44-0.81x131	plano	plano	20/15	0.19	6
-1.50-1.00x060	-1.72-0.56x039	-1.71-0.78x057	plano	plano	20/20	0.22	18
-3.00-0.25x028	-2.44-1.37x009	-2.93-0.59x006	plano	plano	20/15	0.78	3
-3.00-0.25x125	-2.58-1.09x174	-3.06-0.99x174 and WFO plano-0.75x091	plano	plano	20/15	0.65	0
-9.00-0.25x060	-8.00-0.22x027	-8.61-0.08x047	plano	plano-0.50x171	20/20	0.14	20
-8.25-1.25x088	-8.00-0.22x102	-8.01-0.88x089	plano	plano-0.25x101	20/15	0.66	13
-5.00-0.75x170	-4.93-0.90x174	-5.02-0.75x172	plano	-0.25D/S	20/20	0.15	2
-5.50-0.25x030	-5.75-0.76x013	-5.43-0.28x026	plano	-0.25-0.25x095	20/20	0.48	13
-2.50D/S	-1.73-1.55x007	-2.37-0.46x008	plano	plano	20/20	1.09	1
-2.50-0.25x175	-1.85-1.55x173	-2.36-0.73x173	plano	plano	20/20	0.82	6
-3.00D/S	-2.52-0.96x015	-2.83-0.64x018	plano	plano	20/15	0.32	3
-3.00D/S	-2.55-0.91x179	-2.88-0.54x178	plano	plano	20/20	0.37	1

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Table 1 (Continued).

Manifest (Rx Wet)	LZ TX(LYRA)	PHORCIDES	Target Goal	Post-Op RX	VA	Cyl Difference	Axis Difference
-0.75-1.00x023	-0.38-1.75x175	-0.75-1.30x171	plano	plano	20/20	0.44	4
-2.75-0.50x175	-2.25-1.49x175	-2.70-0.79x178	plano	plano	20/15	0.7	3
-1.50-0.50x175	-1.00-1.52x167	-1.38-0.93x169	plano	plano	20/20	0.59	2
plano-2.25x022	+0.00-2.28x027	-0.18-2.18x021	plano	-0.50-0.75x164	20/20	0.1	6
-1.25-3.75x010	-1.65-3.00x006	-1.60-3.25x010	plano	plano-0.50x180	20/25	0.5	4
-2.75-3.75x165	-3.15-3.00x166	-3.02-3.52x166	plano	plano-0.25x180	20/20	0.28	0
-4.50-1.00x142	-2.56-1.89x174	-3.06-1.75x171 and WFO plano-1.00x098	-1.50	-1.25D/S	20/30 J1	0.86	3
-4.00-0.50x035	-3.41-1.68x178	-4.06-1.65x179 and WFO plano-1.00x080	plano	plano	20/20	0.97	1
-2.50-0.75x010	-2.05-1.65x010	-2.39-1.17x007	plano	plano	20/15	0.48	3
-2.25-0.75x005	-1.90-1.43x175	-2.16-1.13x175	plano	plano	20/15	0.3	0
Total OU=101							

Notes: Green denotes those instances when Phorcides would have provided a substantially better result. Yellow denotes a potentially better results with Phorcides and red denotes a poorer visual outcomes using Phorcides compared with the LYRA Protocol.

Abbreviation: Cyl, cylinder.

0.50 D of astigmatism, and only tolerate small amounts of sphere or astigmatism (such as 0.25 D). The majority of patients with 0.50 D of residual astigmatism requests enhancement in our clinic. Therefore, accuracy within 0.5 D of astigmatism became our standard benchmark. During the analysis comparing Phorcides planning with LYRA Protocol treatment results, a residual astigmatism close 0.5 D was rounded up to be significant, but anything less than 0.4 D was not deemed to be significant. This was performed by looking at eyes that had successful results not only by post-operative manifest refraction, but had no visual complaints that would indicate residual refractive error or incomplete aberration treatment with visual abnormalities caused by incoming light scatter.

Final manifest refraction pre-operatively and post-operatively was performed by the surgeon for the purposes of this study review, these were not performed only by technicians or ancillary staff.

Each eye was retrospectively surgically planned with Phorcides to determine if Phorcides was accurate in all of these patients (Table 1). Eighteen eyes that had residual astigmatic errors 3 months after initial correction with Contoura with LYRA protocol were then retrospectively planned with Phorcides to determine if a more accurate outcome would have resulted (Table 2). Finally, 2 patients were treated with LYRA Protocol in the right eye, and Phorcides in the left eye (Table 3).

Planning for both surgical planning systems was conducted according to the recommended protocols.

Phorcides planning is recommended to be performed by the technician, where the HOA pattern derived from the Contoura surgical planning station is entered into

Table 2 Residual Astigmatic Errors 3 Months After Initial Correction with Contoura with LYRA Protocol versus Retrospectively Planning with Phorcides

PHORCIDES	Cyl Delta	Axis Delta	ESX RX
-4.25-1.99x005	0.64	3	-0.25-0.75x080
-6.21-1.02x175	0.39	0	+0.75-1.00x095
-1.83-0.14x014	0.4	10	-0.50-0.75x093
-2.00-0.19x093	0.13	68	-0.25-1.00x110
-2.30-2.45x008	0.55	2	plano-0.75x123
-1.52-3.27x178	0.27	2	plano-1.25x075
-2.02-0.51x177	0.54	0	-0.25-1.50x080
-3.00-0.30x007	0.56	1	-0.25-1.00x095
-0.82-0.66x032	0.74	3	plano-1.00x090
-5.08-0.37x174	0.08	2	+1.00-1.25x125
-3.36-1.22x003	0.63	2	+0.25-0.75x115
-3.76-0.69x006	0.94	2	+0.25-1.00x106
-3.97-0.76x004	0.9	4	+1.00-1.00x090
-0.73-1.49x026	0.6	4	+0.25-1.25x110
-2.46-1.79x084	0.76	0	plano-1.25x080
-2.28-0.83x040 and WFO plano-0.50x060	0.23	9	-0.25-0.75x064
-1.22-1.86x101	0.96	1	+0.25-1.25x085
-0.62-1.32x079	0.76	36	-0.25-1.25x090

Notes: Green denotes those instances when Phorcides would have provided a substantially better result. Yellow denotes a potentially better results with Phorcides and red denotes a poorer visual outcomes using Phorcides compared with the LYRA Protocol.

Abbreviations: Cyl, cylinder; ESX, errors in astigmatism; Rx, prescription; WFO, wavefront optimized.

Table 3 Refractive Outcomes of 2 Patients (4 Eyes) Using LYRA Protocol on the Right Eye and Phorcidex on the Left Eye

LZ TX	Manifest	LYRA Protocol	PHORCIDES	Target Goal	Post-Op RX	VA	Cyl Delta	Axis Delta
LYRA	-4.75D/S	-3.98-1.55x002	-4.67-0.26x180	plano	plano	20/20	1.29	2
PHRCD	-5.00D/S	-4.2, -1.59 x 11	-4.87-0.40x0.14	plano	-0.25-0.75x003	20/20	1.19	3
LYRA	-1.50-0.50x040	-1.42-0.67x019	-1.66-0.37x024	plano	+0.25D/S	20/20	0.3	5
PHRCD	-1.00-0.50x170	-1.44, -1.06 x 179	-1.11-0.48x180	plano	plano-0.75x108	20/20	0.58	1

Abbreviations: Cyl, cylinder; LZ TX, laser treatment; PHRCD, Phorcidex; RX, refraction; VA, visual acuity.

Phorcidex, along with the Contoura measured astigmatism and also posterior astigmatism information from a Pentacam or Galilei Scheimpflug Analyzer. Phorcidex 2.0 was used for surgical planning. Phorcidex 3.0 has been released, but the author was unable to get a registered, functioning copy for this study as only Dr. Lobanoff can provide license keys and Alcon cannot. LYRA Protocol planning was done by the surgeon, eliminating disparate topographic images and then utilizing the Contoura measured astigmatism and axis along with the resultant calculated sphere to treat the patient.

Each surgery was performed by one surgeon (MM), at one center, utilizing the WaveLight EX500 laser.¹⁻³ Flap creation was performed with either the FS200 femtosecond laser or the Moria M2 microkeratome. Patients were followed up at 1 day, 1 week, 1 month, 3 months, 6 months, and 1 year post-operatively, and uncorrected visual acuity, refraction, best corrected visual acuity, topography, epithelial thickness mapping was obtained for each visit.⁸ At 3 months a post-operative Pentacam image was also obtained. All of this testing was also done pre-operatively.

Although the term topography modified refraction (TMR) has been used somewhat interchangeable with Contoura measured astigmatism (CMA), we hesitate to do so as we simply have no experience with other topography units/topographic-guided software besides the WaveLight Contoura system to be able to generalize the term. We will not use TMR in this paper, but rather use Contoura measured astigmatism to be more accurate.

Patients were excluded if they could not achieve 20/20 vision before surgery, had prior refractive surgery, or were not within the FDA approved treatment parameters WaveLight Contoura treatment. Anterior segment abnormalities such as corneal ectasia or keratoconus, recurring eye disease such as severe dry eye, uncontrolled diabetes or hypertension, and pregnancy also disqualified patients from participating in the study.

All patients signed written informed consent forms allowing their data to be used in this study. This study

falls under the exemption of the Health and Human Services (HHS) Policy for the Protection of Human Research Subjects 45 CFR 46.104 (d) for retrospective studies and 46.104 for exempt research, and thus, no Institutional Review Board approval was required.^{9,10} This study also conforms to the Declaration of Helsinki guidelines.¹¹ There were no safety-related incidents that occurred or were reported to Alcon Inc. or WaveLight concerning patients involved in this study.

Results

Table 1 shows the results of the surgical planning of 101 eyes (Group 1) with the resultant outcomes. If Phorcidex is more accurate than the LYRA Protocol, then virtually all of these eyes should have had planning results similar to those using the LYRA Protocol. The magnitude of cylinder difference between the outcome of treatment with Contoura with LYRA Protocol and the theoretical planning with Phorcidex is denoted as “Cylinder Difference”, and the magnitude of axis difference is denoted as “Axis Difference.” The red highlights represent outcomes where Phorcidex would have been inaccurate by 0.5 D or greater of astigmatism and would have left residual astigmatism requiring treatment. Some of these cases also had a significant difference in axis measurement as well. About half (52) of the 101 eyes (51%) were in this category, with Phorcidex planning differing from LYRA by a substantial margin. The average difference in magnitude of astigmatism between Phorcidex and LYRA was 0.69 D (range 0.36–1.33 D), and the average difference in axis was 7.1 degrees (range 0–76 degrees). The low of 0.36 in the range was included because it also had the highest axis deviation (76), so therefore became significant.

The cylinder difference between LYRA and Phorcidex surgical planning outcomes for Group 1 was subjected to a 2-tailed *T* test and the outcomes were statistically significant $p < 0.05$ (-0.00000486). The axis difference between LYRA and Phorcidex was not statistically different when subjected to a 2-tailed *t* test, $p > 0.05$ (0.248).

The percentage of eyes of the 101 that varied between LYRA and Phoricides in this group by 0.5D was 19.8%, by 0.75D was 23.8%, and by 1D was 7.9%.

There were 2 eyes with a post-operative outcome of -0.50 D of astigmatism from target, and one eye with an outcome -0.25 D of astigmatism from target for which Phoricides would have been more accurate than LYRA. These are highlighted in green. These 3 eyes represent 3% of the total.

The yellow highlights represent patient that do not reach the level of 0.5 D of astigmatism, but still may have an astigmatic axis of large enough difference combined with enough residual astigmatism that the patient may possibly have noticed. These included 12 eyes out of 101 (11.9%). Thirty-four eyes (33.7%) had similar surgical planning results between Phoricides and LYRA. These would have resulted in the same clinical outcomes.

Table 2 shows the results of 18 eyes (Group 2) that required secondary enhancement after treatment with Contoura using the LYRA Protocol. Here we determined if Phoricides would have provided a result that was more accurate than LYRA. Pre-op manifest refraction, Contoura with LYRA Protocol Laser Treatment, and Phoricides surgical planning utilizing pre-operative patient data are provided. The Residual Astigmatism column denotes the astigmatism that was present after laser treatment with the LYRA Protocol that needed to be enhanced (all enhancements were performed with Wavefront Optimized Treatment). Cylinder and Axis Difference denote the difference between the LYRA Protocol and Phoricides surgical planning, and comparison to residual enhancement was made to determine if Phoricides would have provided a more accurate outcome.

The cylinder difference between LYRA and Phoricides surgical planning outcomes for Group 2 was subjected to a 2-tailed *T* test and the outcomes were not statistically significant, but were very close to being so $p > 0.05$ (0.5358). The axis difference between LYRA and Phoricides was not statistically different when subjected to a 2-tailed *T* test, $p > 0.05$ (0.536).

Green denotes those instances when Phoricides would have provided a substantially better result as the post-operative residual astigmatism was approximated successfully by the difference between LYRA and Phoricides surgical planning. This was the case for 6 eyes (33%) as Phoricides would have yielded a more accurate outcome. Yellow denotes a potentially better result and there were 3 (17%) of eyes in this category. Red denotes an outcome that Phoricides would not have provided a more accurate result, of which there were 9 eyes (50%).

Table 3 shows the refractive results of 2 patients (4 eyes) that had corrections using LYRA on the right eye and Phoricides on the left eye. Note that the planning outcomes differ significantly between Phoricides and LYRA Protocol on both eyes. The right eye on both patients was performed utilizing LYRA Protocol planning, and the left eye utilized Phoricides. Note the delta of 1.19 D on the left eye of patient 1 and the 0.58 D on the left eye of patient 2 between LYRA Protocol and Phoricides and how it relates to post-operative manifest refraction.

Discussion

In a multi-center study by Stulting et al that used vector analysis for surgical planning outcomes on patients utilizing manifest, LYRA Protocol, and Phoricides, it was noted that LYRA and Phoricides were more accurate than manifest refraction.⁶ Phoricides appeared to be more accurate but the study did not measure clinical outcomes, only theoretical ones. Theoretical outcomes are not the same as clinical outcomes, as the eye does not always function as a Gullstrand model and does have variability and biological reaction to treatment such as epithelial compensation. The differences between LYRA and Phoricides in this study were not large, but this study did seem to show that Phoricides was theoretically more accurate.

In this particular study, we examined 101 eyes successfully treated with the LYRA Protocol retrospectively with the Phoricides Analytic Engine. In particular, we chose eyes that had a significant difference in between manifest astigmatism and Contoura measured astigmatism to create a large enough difference to test the accuracy of Phoricides. If this difference is not large, it has been our experience that Phoricides results in treatment parameters very similar to LYRA Protocol, something that the recent Lobanoff et al study also demonstrated.⁷ With the understanding that Phoricides considers not just the anterior corneal aberrations and anterior corneal astigmatism that LYRA Protocol does, but in addition also considers the posterior corneal astigmatism and lenticular astigmatism, we expected to find that Phoricides would at least be similar in accuracy, and very possibly more accurate if these additional factors were significant and also accurately measured by Phoricides. It was not, with 51% of eyes evaluated likely would have resulted in worse outcomes with substantial residual astigmatism. Using Phoricides, the average deviation from LYRA was 0.69 D with an average axis deviation of 7.1 degrees. In some of these cases, a large axis difference could have incurred corneal higher order aberrations resulting in irregular astigmatism. We already know each degree of deviation of astigmatic axis during correction not only decreases

astigmatic correction efficiency by 3%, but also creates a new aberrant, irregular astigmatism along a different axis.^{2,10} This would result in optical abnormalities and require topographic-guided ablation to correct.

Five of the eyes had Phorides surgical outcomes that recommended a Contoura treatment followed by a WFO treatment, which would have required a second per procedure cost. As we had no access to the algorithm, we cannot explain why this would have been a recommended procedure. In none of these eyes did Phorides appear to offer a better refractive outcome than LYRA.

We then examined 18 eyes that needed secondary enhancement after the LYRA Protocol, to see if Phorides would be more accurate than LYRA. It was definitely more accurate in 1/3 of the eyes, but not accurate in 1/2 of eyes. If the remaining 3 eyes that the new analytic software could have possibly improved were included, Phorides could have improved visual outcomes at best on 1/2 of the eyes needing secondary enhancement.

If Phorides had been as accurate as LYRA in Group 1 then we could say that Phorides is more accurate clinically than LYRA since it did provide a degree of more accuracy in the patients requiring secondary enhancements. Unfortunately, Phorides provided worse visual outcomes than LYRA in Group 1, and at best only would have helped prevent enhancement in Group 2 about 50% of the time.

Interestingly, one of the patients that Phorides would have benefitted was a patient in which significant epithelial compensation of corneal higher order aberration had been documented, and was actually significant enough to warrant inclusion in a separate epithelial compensation of aberration manuscript.⁸ As far as we know, the algorithm for Phorides does not include any analysis of epithelial compensation of corneal higher order aberrations, so it is unknown if the increased accuracy on these two eyes is somehow coincidental.

The Phorides surgical planning on these patients was distinctly different from manifest refraction and LYRA Protocol planning. Since each patient already had successful correction with plano outcome via LYRA Protocol, it is not difficult to see that planning with manifest or Phorides would have resulted in a significantly different outcome. In our experience, patients treated with Contoura do not tolerate 0.50 D or greater of astigmatism post-operatively. These patients may be 20/20 and be considered a “success” via other standards, in our experience that patients are not satisfied with their vision and wish to be corrected. As previously mentioned, we find that patients with more uniform cornea and decreased corneal high order aberrations are less tolerant

of any flaws in their vision. Essentially residual aberration may cause blur that masks residual astigmatism, but we have seen a more uniform cornea seems to unmask every flaw that is residual, including significant flaws and astigmatism posterior to the anterior cornea.

If Phorides is used for smaller differences between manifest and Contoura measured astigmatism, it shows a relatively small delta with LYRA Protocol in magnitude of astigmatism and axis, and the result would likely be well tolerated. Why this is so is difficult to answer completely as the proprietary algorithm of Phorides is not known. The one factor that stands out with Phorides analysis, is that the astigmatism never exceeds the Contoura Measured Astigmatism (CMA) UNLESS that number is less than the manifest refraction. CMA exceeds manifest refraction in the vast majority of cases, and in these cases Phorides has always resulted in an astigmatism in between manifest and CMA. If Phorides accurately determines posterior astigmatism as a factor, there would be some case where the corneal and posterior astigmatism would interact in these patients and result in a number not between manifest and CMA, yet we have never seen that occur. Not having any access to the proprietary Phorides algorithm, it is difficult to say with any certainty why this is the case.

Lobanoff et al published a study in JCRS in 2020 comparing results utilizing Phorides surgical planning to treatment with manifest refraction. This is the first published clinical results for Phorides, but there are a series of issues with this study. First, although the authors state that there was a significant difference between manifest astigmatism patients and those treated with Phorides, the average difference was actually quite small, only about 0.3D or less. Furthermore, the Contoura measured astigmatism was almost exactly that of the Phorides treatment (1.07D vs 1.11D of astigmatism) demonstrating that this study did not test the Phorides system at all. Essentially this study shows that Phorides essentially tracks the same as LYRA Protocol when the differences between manifest astigmatism and Contoura measured astigmatism are not large, something that we already noted in this study. Unfortunately, this study does not provide axis differences so it is impossible to tell if that was significant, and it does not provide any statistical analysis to show if the differences in astigmatism between manifest, Contoura measured and Phorides were significant.

Second, the results show more eyes achieving 20/15 in the Phorides group vs the manifest astigmatism group, but interestingly, the refractive outcomes were worse for

Phorides. Based on this and the results of this particular study, Phorides may actually be worse than not only LYRA, but manifest treatment as well.

Third, this study shows a small but significant loss of best corrected visual acuity post-operatively with Phorides surgical planning, something that may be occurring as Phorides in attempting to treating lenticular astigmatism would not correctly treat the anterior elevation of the cornea, and likely leave residual higher order aberration behind. Notably, it would not make a more uniform if it differed significantly from the Contoura Measured astigmatism, and this may be the cause of best corrected visual acuity loss, something that we simply have not seen in any significant capacity in utilizing LYRA Protocol. Since this study did not look at the topographic effects at all, there is absolutely no information on the resultant corneal shape.

Fourth, the authors make some claims that I find curious, that Phorides utilizes corneal elevation implying that somehow that Contoura without Phorides does not. Contoura utilizes corneal elevation, and the Contoura measured astigmatism in the LYRA Protocol has been shown to treat the anterior corneal elevation creating a more uniform cornea.

Finally, in the Lobanoff study, there is no subjective patient data given at all, purely refractive and visual outcomes. The entire goal of reducing corneal aberrations is to improve vision both subjectively and objectively. No information was given as to which eye the patients preferred, the manifest or the Phorides treatment or equally. LYRA Protocol has already shown to provide extremely high levels of subjective patient satisfaction.

Jankov et al in 2006¹² also questioned the wisdom of

trying to correct the anterior corneal surface using the ablation profile based on the whole eye aberrations when the anterior corneal surface aberrations are clearly preponderant.

Furthermore, they also pointed out that posterior aberrations can change with age, and postulated that recreating the natural aspheric shape and a uniform cornea might well be the ideal.

Manns et al in 2002¹³ noted that because the corneal reshaping alters the path of light rays propagating in the eye, and although the lens and posterior corneal surface are unchanged after surgery, their contributions to the ocular aberrations will likely be different from pre-operatively.

Furthermore, as mentioned prior, the authors have noted that if the anterior elevation is not treated on a cornea, residual aberration may be left behind. In other words, anterior corneal aberrations may be left behind that cancel

out posterior astigmatism/aberrations. Unfortunately, as we are learning via the use of OCT epithelial thickness mapping, this may create a reaction via epithelial hyperplasia that may affect the refractive outcome.⁸ We have seen even small amounts of epithelial compensation to corneal HOA (2–3 μm) can cause residual astigmatism.

It is unknown if the Ray Tracing procedure ultimately will be subject to the same issues, as it also attempts to treat aberrations at a specific time to make each incoming light ray focus on the corresponding retina. It does not account for internal structures changing with age, nor does it account for the variability of the epithelial layer. If utilized in a LASIK procedure, Ray Tracing also cannot compensate for the biomechanical corneal changes that occur in some patients after flap creation that lead to an inaccurate outcome.

The author attempted to treat patients with a similarly large difference between manifest and Contoura measured astigmatism with one eye treated with LYRA and one treated with Phorides surgical planning. The first two patients both showed residual astigmatism via refraction post-operatively in the Phorides eye, even though they had good outcomes for both eyes based on solely vision results (20/20 or better). This measurable astigmatism was equivalent to the difference between LYRA Protocol and Phorides, and remained through the 6-month follow-up. This trial was then discontinued.

The authors had high hopes for Phorides initially, as the idea behind it has great appeal. It is difficult to analyze if further development of Phorides would increase accuracy as the underlying algorithm is not known to us nor has it been published anywhere. We are also unaware of any published clinical results. These data demonstrate that Phorides not only does not increase the accuracy of surgical outcomes, but actually decreases accuracy in the majority of patients to an amount unacceptable in our clinical practice. We do not yet understand why the vector calculations show a greater level of accuracy in the aforementioned study, but theoretical calculations are not as important as clinical outcomes.

Although Phorides would have helped with accuracy on a number of patients that needed secondary enhancements with LYRA, it showed significantly worse performance in Group 1 than the LYRA Protocol and would have resulted in large number of patients with significant residual astigmatism. As the axis was also different in many patients, the possibility of residual astigmatism was also present. Although the small sample size of two patients in Group 3 was not anywhere near statistical significance, the combination of the retrospective data

obtained, as well as the inaccurate outcomes of the first two eyes treated with Phorides surgical planning resulted in the discontinuation of Phorides use in our center.

Finally, the authors repeat the cautions of past authors of correcting posterior astigmatism/aberrations on the anterior corneal surface, as the fact remains that changing corneal structures with age, issues with the treatment of posterior astigmatism on the anterior surface, and epithelial compensation of residual anterior corneal aberration remain challenging issues for of treatment protocols that attempt to treat posterior astigmatism/aberrations on the anterior corneal surface. The authors agree with Jankov et al¹² and Manns et al¹³ that the best way to treat corneas is to attempt to create a uniform, aspheric shape to minimize light scatter and epithelial compensation, and to maximize visual clarity and accurate, stable corrections.

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Disclosure

Dr. Motwani has received a grant from Alcon for a previous unrelated study in 2017, during the conduct of the study. Dr. Motwani received non-financial support from Optovue, Inc. (loan of Optovue Avanti, clinical support in understanding analysis of data). Dr. Motwani has a patent pending on a theoretical device that could combine corneal HOA data and epithelial thickness data to achieve a more accurate refractive outcome; and reports patents pending for: the use of topographic-guided ablation for the creation of a uniform cornea; and the use of a device to utilize both epithelial thickness and corneal aberration to make a uniform cornea. The author reports no other potential conflicts of interest for this work.

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