

ESCRS

EUREQUO

Annual Report 2017







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Annual Report 2017

based on data from the European Registry of Quality Outcomes for Cataract and Refractive Surgery

EUREQUO Steering Committee

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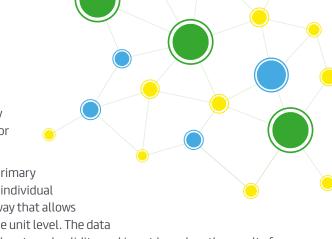
1. Background

What is the Purpose of a Registry

In health care, most countries have well-developed and functioning systems to monitor economic and human resource activities. Corresponding systems have not been developed for working with patients, although this is the actual core and the ultimate aim of provider organizations. The traditional patient record systems have not facilitated the compilation and analysis of data needed for quality improvement. Although increasingly more records are electronic, they essentially continue to be note pads that individual caregivers use for memory support in treating individual patients.

Quality registries have been developed to fill the gap left by the lack of primary monitoring systems. The quality registries collect information on individual patient's problems, interventions, and outcomes of interventions in a way that allows the data to be compiled for all patients and analysed anonymously at the unit level. The data

from registers is invaluable; as the conclusions are evidence based with external validity and is not based on the results from a small clinical trial or a case series. In summary, the quality registries are created and ran by professionals within areas where a need for specific information not covered by textbooks or scientific literature has been identified.



EUREQUO Success and Future

The ESCRS believes that a continuing audit of surgical outcomes is necessary to ensure the best care for patients, by making comprehensive data available for comparison of visual outcomes.

EUREQUO is one of the largest international IT projects in ophthalmology, connecting surgeons all over the world and building a network to facilitate the exchange of expertise.

The EUREQUO Steering Committee is constantly working to update the platform with new parameters to reflect the development of new surgical techniques and trends in the field. The committee is also committed to analysing the aggregate data of the database and developing new scientific papers and has released eleven publications to date. This work will continue.

The EUREQUO Platform

The European Registry of Quality Outcomes for Cataract and Refractive Surgery (EUREQUO) provides a platform to audit surgical results and encourages surgeons to adjust their techniques and improve their outcomes. The EUREQUO platform contains important clinical parameters but also patient-reported outcome measures for both cataract and refractive surgery.

One important use of the database is benchmarking. To date, more than 2.6 million cataract surgeries and 115.000 refractive surgeries have been recorded in the system.

EUREQUO is funded by the ESCRS, with initial support from the EU. The platform has been launched globally, and all ESCRS members can access it free of charge.

EUREQUO Benefits and Benchmarking

By collecting data, surgeons will be able to monitor their own results over time, and anonymously compare their results with other colleagues, clinics and countries.

- EUREQUO is a convenient web-based registry. Cataract, refractive and patient-reported outcomes are all in one platform.
- EUREQUO provides a unique opportunity to monitor and compare results.
- Collecting data will support surgeons to make an audit report.
- EUREQUO is linked to patient-reported outcomes.
- EUREQUO allows clinical improvement using patient-reported and clinical data.

Only anonymous data are stored within EUREQUO, and only clinical data without any patient ID data are mandatory within the EUREQUO datasets.

For more information visit the EUREQUO website at: www.eurequo.org

2. Cataract Surgery

Development of the Database

Data presented here is based on the database standing at 31st May 2018. The number of cataract extractions reported to the EUREQUO database is shown in Table 1.

Table 1.

Number of annually reported cataract extractions and the accumulated number of cases in the database.

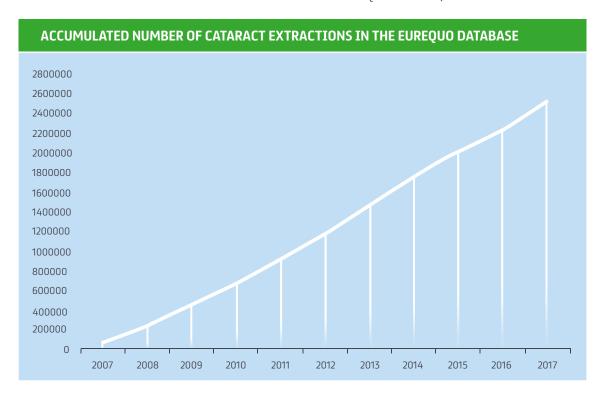
Year	Number of reported cataract extractions	Accumulated number of cataract extractions in the database
2007	57,552	57,552
2008	165,647	223,199
2009	201,355	424,544
2010	227,283	651,837
2011	249,474	901,311
2012	260,525	1,161,836
2013	280,090	1,441,926
2014	297,564	1,739,490
2015	259,662	1,999,152
2016	203,955	2,203,107
2017	296,787	2,499,894

The growth of the database is shown graphically in Figure 1.

Figure 1.

Number of accumulated cataract extractions in the EUREQUO database over time. One million cases were achieved in 2011 and two million cases in 2016.

Since launch the number of participating countries has varied between 14 and 16 and the number of reporting sites between 185 and 221. In 2017 clinics from 14 countries entered data to the EUREQUO database, 12 from the EU and 2 from outside the EU.



Trends in Outcomes Over Time

Visual outcome, postoperative corrected distance visual acuity, LogMAR. Cases with follow up data.

Year	N	Mean	SD
2008	64,751	0.088	0.23
2009	100,302	0.081	0.23
2010	110,037	0.080	0.23
2011	114,036	0.076	0.23
2012	122,024	0.069	0.22
2013	118,625	0.062	0.20
2014	142,591	0.069	0.24
2015	140,220	0.059	0.23
2016	91,725	0.041	0.16
2017	148,680	0.057	0.26

Refractive outcome, biometry prediction error (spherical equivalent). Cases with follow up data.

Year	N	Absolute mean	SD	within ±0.5D
2008	63,568	0.478	0.54	67.3%
2009	100,628	0.457	0.52	69.0%
2010	111,091	0.443	0.49	70.2%
2011	114,414	0.433	0.48	71.2%
2012	122,367	0.419	0.52	72.3%
2013	118,889	0.416	0.47	72.4%
2014	142,575	0.435	0.55	72.3%
2015	140,201	0.415	0.48	73.2%
2016	91,934	0.412	0.48	73.5%
2017	148,939	0.403	0.46	73.7%

Any surgical complication

Year	N	Number	% of all surgeries
2007	85,711	2768	3.2
2008	137,543	3402	2.5
2009	201,287	4593	2.3
2010	226,471	4973	2.2
2011	244,639	6005	2.5
2012	256,855	4908	1.9
2013	261,100	4354	1.7
2014	293,554	7103	2.4
2015	254,838	3323	1.3
2016	197,391	2447	1.2
2017	296,787	3445	1.2

The trends show an improvement for visual outcome, refractive outcome and a reduced number of surgical complications.

Data for 2017

Preoperative and surgical data

The number of cataract extractions reported into the database for 2017 was 296,787. The mean age of the operated patients was 73.0 years and 57% were females and 43% males.

Preoperative visual acuity

The preoperative corrected distance visual acuity (CDVA) in surgery eye was 0.1 (6/60) or worse in 10.6% of all cases. In 42.6% of the cases the preoperative CDVA was worse than 0.5 (6/12). This means that in 57.4% the visual acuity was 0.5 (6/12) or better.



Ocular co-morbidity

In 27.3% there was an ocular co-morbidity in the surgery eye. Such co-morbidities were Glaucoma (6.7%), Macular Degeneration (10.7%), Diabetic Retinopathy (2.7%), Amblyopia (1.1%) or another sight-threatening eye disease (8.7%).

Surgical difficulty

In 10.6% the surgeon reported a surgical difficulty. Such a difficulty could be: Previous corneal refractive surgery (0.20%), White/Brown cataract with need of capsular staining (2.22%), Small pupil with need for mechanical dilatation (3.13%), Corneal opacities (0.9%), Previous vitrectomy (0.67%) and other surgical difficulty (4.38%).

Surgical experience

This was a new parameter for 2017. This parameter was reported for 13,302 surgeries, and 246 (1.8%) of these surgeries were performed by a trainee.

Type of operation

A phacoemulsification with implantation of a posterior chamber intraocular lens (IOL) was the most frequent type of surgery (97.1%). In 5943 (2.0%) cases a planned extracapsular cataract extraction was performed, in 167 (0.1%) cases an anterior chamber IOL was implanted and in 52 (0.02%) cases a cataract extraction was combined with a filtering procedure. A femtosecond laser-assisted cataract extraction was reported in 603 (0.2%) cases. In 1906 (0.6%) cases an "another" type of operation was performed.

Type of IOL material

The dominating type of IOL material was an acrylic hydrophobic IOL, 87.7% received such an IOL. An acrylic hydrophilic IOL was implanted in 9.2%. In 273 cases (0.1%) no IOL was implanted and the eye was left aphakic.

Additional refractive quality IOL

Additional refractive quality IOL was reported for 124,523 cataract extractions. In 2066 (1.7%) cases a multifocal IOL was implanted and in 480 (0.4%) a toric or multifocal toric IOL was implanted.

Surgical complications

In 1940 (0.65%) cases a posterior capsule rupture with or without vitreous loss occurred. For trainees this complication occurred in 7.3%. Other surgical complications were: Dropped nucleus in 84 (0.03%) cases, Iris damage in 349 (0.12%) cases, and "Other sight-threatening complication" in 1167 (0.39%) cases.

Follow up Data (data from clinics committed to report outcome data)

Complete follow up data was reported for 149,043 cataract extractions. The average time from date of surgery to date of follow up was 31 days. The mean age of follow-up patients was 73 years and 57% were females.

Visual outcome

A final CDVA of 1.0 (6/6) or better was achieved by 75.9% (95% Confidence Interval [CI]: 75.7-76.1) of all reported cases. A CDVA of 0.5 (6/12) or better was achieved by 95.2% (95%CI: 95.1-95.4) of all cases.

Refractive outcome

The average spherical equivalent error was -0.03D (SD 0.61). The absolute mean spherical equivalent error was 0.40D (SD 0.46). Within ±0.5D biometry prediction error (spherical equivalent) was 73.7% (95% CI: 73.4-73.9), and within ±1.0D 93.8% (95% CI: 93.7-93.9).

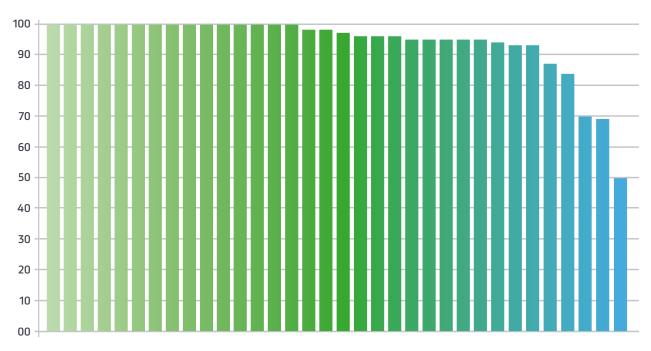
Postoperative complications

In 98.4% of the cases no postoperative complication was reported during the follow up time. In 181 (0.12%) cases there was a persistent corneal edema and in 57 (0.04%) cases uncontrolled elevated intraocular pressure. Eighteen cases (0.012%) of endophthalmitis were reported. This number does not include cases that occurred later than the follow up date.

Benchmarking Diagrams

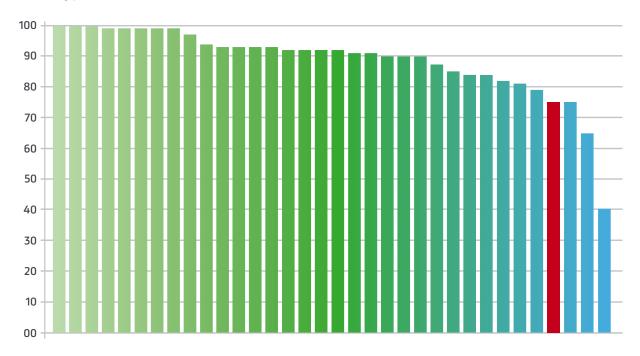
1. Visual outcome, percentage of cases with CDVA 0.5 (6/12) or better. Each bar represents one clinic. Only a part of the clinics is shown.

BCVA surgery eye >= 0.5 | (6/12) at final follow-up percent



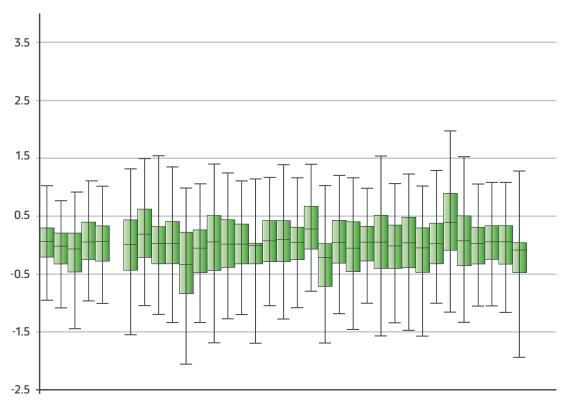
2. Biometry prediction error. Percentage of cases within ±1.0 D of deviation from intended refractive outcome. Every bar represents a unit. Red bar = requesting unit. Only a part of the clinics is shown.

Biometry prediction error, % within 1D deviation from intended



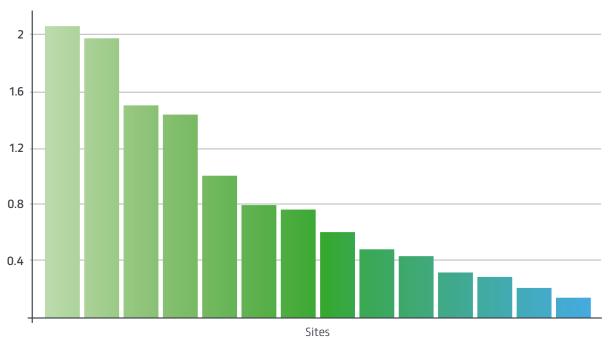
3. Boxplot diagram with biometry prediction error, correct sign. Every box represents a unit. Only a part of the clinics is shown.

Biometry prediction error, boxchart with correct signs



4. Reported capsule complications in per cent (posterior capsular break with or without vitreous loss) during surgery. Every bar represents a unit. Only a part of the clinics is shown.

Capsule complication in per cent



3. Refractive Surgery

Development of the Database

The number of refractive surgeries reported to the EUREQUO database is shown in Table 2.

Table 2.

Number of annually reported refractive surgeries and the accumulated number of cases in the database.

The database inherited historical cases from the Refractive Outcomes Information System collected until the end of 2007.

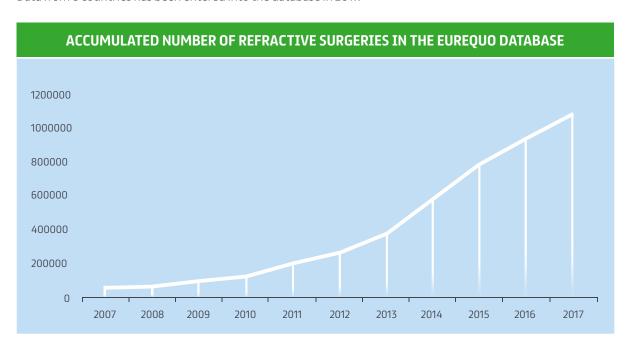
Year	Number of reported refractive surgeries	Accumulated number of refractive surgeries in the database
Inherited cases until the end of 2007	5773	5773
2008	492	6265
2009	1081	7346
2010	3100	10,446
2011	7766	18,212
2012	6837	25,049
2013	10,926	35,975
2014	19,711	55,686
2015	20,279	75,965
2016	15,470	91,435
2017	13,758	105,193

The growth of the database is shown graphically in Figure 2.

Figure 2.

Number of accumulated refractive surgeries in the EUREQUO database over time. 50,000 cases were reached in 2014 and 100,000 in 2017.

Data from 8 countries has been entered into the database in 2017.



Data for 2017

The number of refractive surgeries entered into the database for 2017 was 13,758.

Preoperative data

The mean age of the patients was 43.8 (SD 13.6) years and 48.4% were females and 51.6% were males.

The mean preoperative corrected distance visual acuity (CDVA) in the eye to be operated on was -0.004 (SD 0.06) LogMAR and the median value 0.0 LogMAR. The mean preoperative spherical refraction was -1.11 (SD 3.07) and the mean spherical equivalent was -1.50 (SD 3.09). A preoperative myopia of -3.0D or more existed in 26.6% of the eyes and a hyperopia of 2.0D or more in 11.0% of the eyes. A myopic spherical equivalent existed in 62.8% of the eyes, 1.2% were emmetropic and 36.0% had a hyperopic spherical equivalent before surgery.

Type of surgery

Most of surgeries were primary (99.2%). Only a small fraction were enhancements (0.6%) or additional (0.2%) surgery.

Surgery method

LASIK was performed in 4229 eyes, 691 by blade and 3538 by femtosecond laser. LASEK was performed in 1424 eyes and PRK in 1098 eyes. Another 108 unspecified laser surgeries were reported.

Altogether 185 incisions were reported.

Refractive lens exchange (RLE) dominated the type of refractive surgeries during 2017 in the database, 6289 eyes were reported. Of these surgeries 5727 were specified with type of IOL and 562 were unspecified. Among the specified RLE surgeries 5251 were trifocal IOLs with (1390) or without (3861) a toric IOL as well. A monofocal IOL was implanted in 468 cases with (215) or without (253) a toric IOL.

A phakic IOL in posterior chamber was implanted in 664 eyes and a phakic IOL in the anterior chamber was implanted in 53 eyes. An unspecified phakic IOL surgery was performed in another 27 cases.

Surgical complications

Very few surgical complications were reported. A corneal flap complication occurred in 20 eyes out of 3538 femtosecond LASIK procedures. Another cornearelated complication occurred in 17 cases. In single eyes of RLE a posterior capsular tear occurred (3 eyes).

Visual and refractive outcomes

Follow up visit and examination occurred on average 96 (SD 36.2) days after surgery.

In 63.6% of the eyes a final uncorrected distance visual acuity of 1.0 (6/6) was achieved and in 90.1% the same final distance visual acuity was achieved with best correction.

A final refraction within ±0.5D was achieved in 88.1% of the eyes.

Postoperative complications

In altogether 47 eyes a postoperative complication was reported. Sixteen eyes with significant haze, and 13 eyes with an optical error (>2.0D of error) were reported.

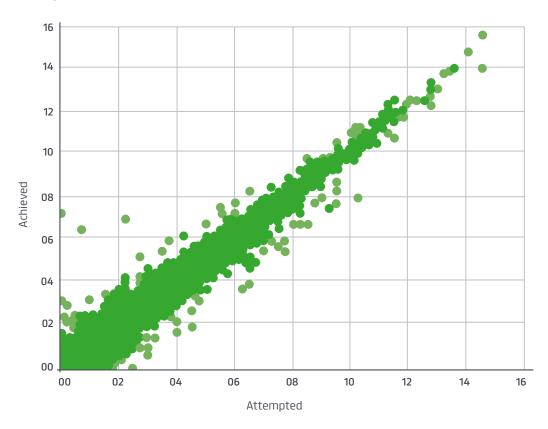


Benchmarking Diagrams

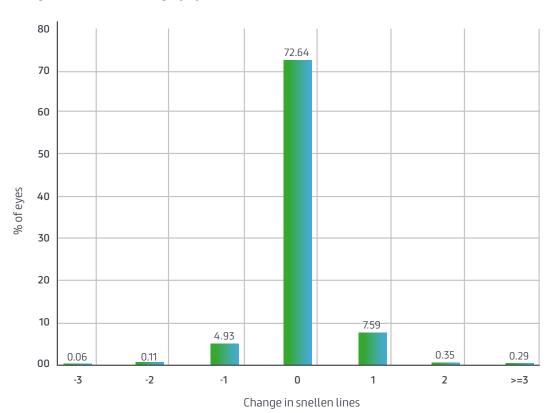
from the refractive surgery output report

The Graphs are from a Group of Reporting Units

Graph 1.Attempted versus achieved refraction



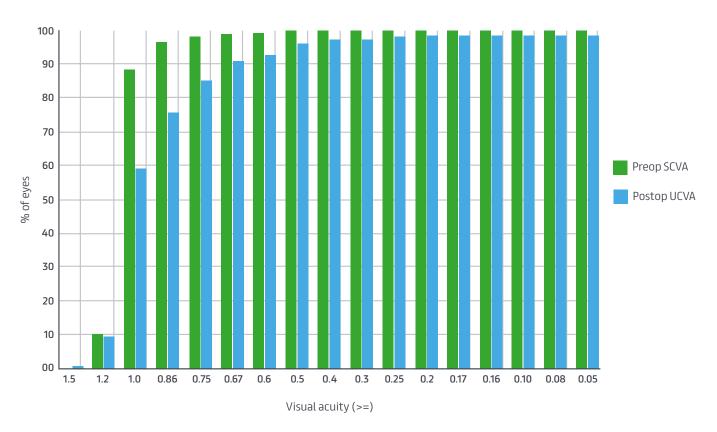
Graph 2.Change in Snellen lines, surgery eye



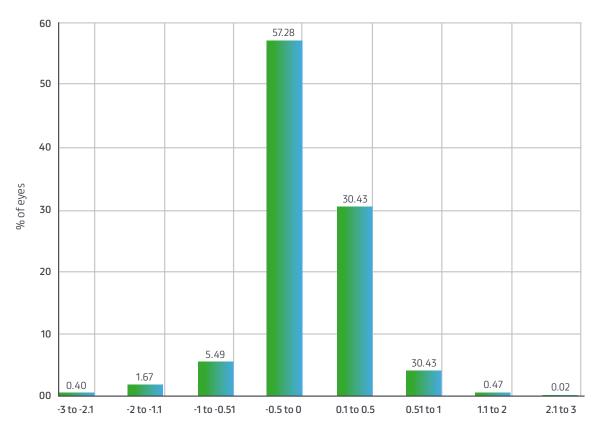
Benchmarking Diagrams

from the refractive surgery output report

Graph 3.Preoperative spectacle corrected VA and postoperative uncorrected VA



Graph 4.Postoperative spherical equivalent



Postoperative spherical equivalent refraction (D)

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