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Titolo	Aniridia and WAGR syndrome : a guide for patients and families / / edited by Jill Ann Nerby and Jessica J. Otis
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Descrizione fisica	1 online resource (xv, 191 pages) : illustrations
Collana	Oxford scholarship online
Altri autori (Persone)	NerbyJill Ann OtisJessica J
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Nota di contenuto	Preface; Contents; About the Authors; 1. Aniridia, WAGR Syndrome, and Associated Conditions; 2. Inspirations; 3. Aniridia-Epidemiology and Genetics; 4. Personal Experiences of Individuals with Aniridia; 5. Glaucoma Problems Associated with Aniridia; 6. Cornea and Lens Problems in Aniridia; 7. Low Vision and Aniridia; 8. Psychological Support; 9. Parents' Experiences; 10. Parents' and Families' Guide; 11. Teachers' and School Administrators' Guide; 12. Jill Nerby and Aniridia Foundation International; 13. Other Support Services; Appendix; Glossary; Index
Sommario/riassunto	When a child is born without a complete iris, it is usually a symptom of a broader condition. Known as aniridia, this condition can also be a sign other parts of the eye are underdeveloped as well. Moreover, recent research shows that the gene involved can also affect the kidneys, pancreas and forebrain, so aniridia can coincide with a range of symptoms known as WAGR syndrome. Until recently, however, there was very little information available on aniridia and WAGR Syndrome.

Even now, not all of the available information is current or correct, so that when a child is diagnosed with aniridia, the parents often find or are given information that is confusing and even frightening. Our hope is to enlighten and encourage those affected by aniridia and WAGR Syndrome by providing patient support and medical information.
