RESEARCH

RPE65-associated inherited retinal diseases: consensus recommendations for eligibility to gene therapy

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Abstract

Background: This research aimed to establish recommendations on the clinical and genetic characteristics necessary to confirm patient eligibility for gene supplementation with voretigene neparvovec.

Methods: An expert steering committee comprising an interdisciplinary panel of Italian experts in the three fields of medical specialisation involved in the management of RPE65-associated inherited retinal disease (IRD) (medical retina, genetics, vitreoretinal surgery) proposed clinical questions necessary to determine the correct identification of patients with the disease, determine the fundamental clinical and genetics tests to reach the correct diagnosis and to evaluate the urgency to treat patients eligible to receive treatment with voretigene neparvovec. Supported by an extensive review of the literature, a series of statements were developed and refined to prepare precisely constructed questionnaires that were circulated among an external panel of experts comprising ophthalmologists (retina specialists, vitreoretinal surgeons) and geneticists with extensive experience in IRDs in Italy in a two-round Delphi process.

Results: The categories addressed in the questionnaires included clinical manifestations of RPE65-related IRD, IRD screening and diagnosis, gene testing and genotyping, ocular gene therapy for IRDs, patient eligibility and prioritisation and surgical issues. Response rates by the survey participants were over 90% for the majority of items in both Delphi rounds. The steering committee developed the key consensus recommendations on each category that came from the two Delphi rounds into a simple and linear diagnostic algorithm designed to illustrate the patient pathway leading from the patient’s referral centre to the retinal specialist centre.

Conclusions: Consensus guidelines were developed to guide paediatricians and general ophthalmologists to arrive at the correct diagnosis of RPE65-associated IRD and make informed clinical decisions regarding eligibility for a gene therapy approach to RPE65-associated IRD. The guidelines aim to ensure the best outcome for the patient, based on expert opinion, the published literature, and practical experience in the field of IRDs.

Keywords: Gene therapy, Inherited retinal diseases, RPE65, Voretigene neparvovec

Background

Inherited retinal diseases (inherited retinal dystrophies; IRDs) are a heterogeneous group of ocular neurodegenerative disorders resulting from mutations in any one of over 250 causative genes [1]. They are mostly characterised by progressive retinal degeneration that leads to severe visual impairment and blindness [2–6]. Inherited
Retinal dystrophy caused by confirmed biallelic mutations in the *RPE65* gene, which encodes all-trans retinyl ester isomerase, an enzyme critical to the visual cycle, is a serious and sight-threatening autosomal recessive genetic disorder that causes a severe form of rod-cone mediated IRD that eventually progresses to complete blindness [3–5]. The spectrum of *RPE65*-mediated IRD exhibits common clinical findings, initially characterised by nyctalopia (night blindness), present from early childhood and due to a primary effect on the rod photoreceptors [7, 8]. The visual function of individuals affected with IRD declines with age, with deteriorating visual acuity (VA) and progressive loss of retinal structure and function (retinal sensitivity) on visual field testing by Goldmann kinetic perimetry (GVF), often leading to blindness in young adulthood [8–11]. The disease course may include earlier or later onset, nystagmus, along with night blindness and loss of vision. Indeed, individuals with biallelic *RPE65* mutations may be given one of a variety of clinical diagnoses. Depending on the time of disease onset, severity, rate of progression and presenting phenotype, the most common diagnoses are Leber congenital amaurosis (LCA) and early-onset severe retinal dystrophy (EOSRD) [5, 8]. These forms are thought to be responsible for approximately 5% of cases of severe IRDs [7]. However, a smaller proportion of patients exhibit a milder phenotype with a slower progression, possibly associated with hypomorphic alleles [11–13].

Initially considered incurable, as the understanding of the pathophysiological mechanisms underlying the subtypes of IRD has expanded, a number of therapeutic approaches to treating IRDs have been proposed, the most advanced of which is gene supplementation therapy [6]. Monogenic ocular diseases are good candidates for gene transfer therapy, as the eye has favourable anatomical and immunological characteristics, providing a contained physical space protected by the blood-ocular barrier that is particularly suited for local delivery [14]. Remarkably, *RPE65*-associated IRD represents a successful model for the development of ocular gene supplementation therapy applied to monogenic diseases.

The proof of principle of gene therapy for *RPE65*-associated IRD was demonstrated in murine and canine models of LCA [15–17], in which a recombinant adeno-associated viral vector serotype 2 (AAV2) gene replacement therapy produced encouraging improvements in visual function. This led to a clinical trials programme that confirmed the safety, durable efficacy, and favourable benefit-to-risk profile of voretigene neparvovec (AAV2-hRPE65v2, voretigene neparvovec-rzyl, LUXTURNA™; Spark Therapeutics, Inc, Philadelphia, PA, USA, Novartis, Basel, Switzerland), administered as a (one-time) subretinal injection, in improving retinal and visual function in *RPE65*-mediated IRD [18–30].

Voretigene neparvovec received marketing authorisation for the US in 2017 [31] and the European Union in 2018 [32] for the treatment of adult and paediatric patients with vision loss due to IRD related to confirmed *RPE65* biallelic mutations and who have sufficient viable retinal cells [32].

A precise genetic diagnosis is necessary to establish eligibility for treatment of *RPE65*-associated IRD and to optimise the use of a precision therapeutic intervention such as voretigene neparvovec in a clinically and genetically heterogeneous group of IRDs. Not only is there a lack of shared criteria for the selection of patients suitable for *RPE65* gene therapy, but the cost and complexity of the procedure mean that an equitable and transparent process for evaluating the urgency to treat for eligible patients is also necessary. In the absence of specific national guidance in this area, the goal of this project was to develop a clinical pathway algorithm that sets forth a stepwise process for ophthalmologists and geneticists to make decisions about the correct diagnosis and treatment with voretigene neparvovec of patients with *RPE65*-associated IRD. Herein, we report the outcomes of a consensus process by a group of Italian experts in IRDs to establish recommendations on the clinical and genetic characteristics necessary to confirm patient eligibility for gene therapy with voretigene neparvovec.

**Methods**

A steering committee (the authors of this paper) comprising an interdisciplinary panel of Italian experts in the three fields of medical specialisation involved in the management of *RPE65*-associated IRD (medical retina, genetics, vitreoretinal surgery) was established to investigate the correct identification of patients with the disease, identify the fundamental clinical tests to determine the correct diagnosis and to evaluate the urgency to treat patients eligible to receive treatment with voretigene neparvovec. The Delphi technique is a recognised and reliable means of consensus-building utilising a series of precisely constructed questionnaires to collect data from an external survey panel with recognised experience in the field of interest [33, 34]. A standard Delphi process was chosen to reach consensus among a wider external panel of experts comprising ophthalmologists (chosen from the applicable sub-specialties of retina specialists, paediatric ophthalmologists and vitreoretinal surgeons) and geneticists with extensive experience in IRDs in Italy who were identified and invited to participate in the survey rounds. The survey participants, named as the Italian IRD Working Group and listed in Additional file 6: Appendix, were carefully selected to minimise the risk that, in such a multidisciplinary field, there could be skills gaps between the respondents.
Delphi process
The extensive clinical expertise of the steering committee was drawn upon to build broad and open-ended statements for the first Delphi questionnaire in order to elicit further qualitative input on the diagnostic and treatment approaches of the participating experts to be considered for inclusion in a second Delphi round. The initial questionnaire was supported by a comprehensive review of the literature to identify the current understanding of the disease and the place of voretigene neparvovec gene supplementation in its treatment [35]. The categories covered in the literature review included: RPE65-related IRD, IRD screening and diagnosis, gene testing and genotyping, ocular gene therapy for IRDs, and voretigene neparvovec. The candidate items were finalised to ensure that they were understandable and exactly captured the information needed for designing the decision pathway. Then, they were reviewed by a small, independent validation panel of experts comprising one expert from each specialty (genetics, ophthalmology) and not part of the population surveyed (see Additional file 6: Appendix 1) before the questionnaire was distributed for administration to the wider survey panel.

A 5-point Likert scale in which 1 = strongly disagree, 2 = disagree, 3 = neither agree nor disagree, 4 = agree and 5 = strongly agree was proposed to rate the answers generated in the Delphi rounds. Disagreement was defined as a Likert response of 1, 2 or 3; Agreement as a Likert response of ≥ 4, and Consensus was defined as a level of agreement of ≥ 70%. Although a universally-agreed proportion has not been established for the Delphi process [36], a 70% cut-off was chosen as a rigorous way of determining consensus, with agreement from over two-thirds of the panel considered to provide a reliable indicator of consensus. The stability of the data was tested by assessing the change in the degree of consensus between Round 1 and Round 2 [33].

As part of the questionnaire format, potential challenges relating to the multidisciplinary nature of the survey population were addressed by further analysing the responses through stratification of the sample according to specialty (clinical, genetic, surgical). Taken together with the participants’ self-assessed degree of confidence in the specific sub-topic, section by section, a more precise interpretation of the questionnaire responses was facilitated.

Refinement of Round 1 results
The results of the Round 1 questionnaire were further developed by the steering committee, considering all of the important differences in responses among the medical specialties, and modifying or re-proposing statements that did not achieve consensus in Round 1. The second-round questionnaire, therefore, built on the statements in each section and incorporated additional statements designed to capture the range of information necessary to establish the diagnostic algorithm. The revised statements were circulated electronically and underwent a second Delphi consensus round. The Round 2 survey results were then ordered and prioritised towards the development of the diagnostic algorithm designed to refine the selection of patients and optimise treatment outcomes.

Results
Key consensus recommendations
Response rates by the survey participants were over 90% for the majority of items in both Delphi rounds. Key consensus recommendations from the two Delphi rounds are summarised in Table 1 and the consensus findings of the Round 1 and Round 2 questionnaires are detailed in Additional files 1, 2, 3, 4 and 5. The following is a summary of the consensus of the indications obtained in the course of the Delphi approach.

Clinical Manifestations of RPE65-associated IRD (Additional file 1: Table S1)

1. When assessing a patient with suspected RPE65-associated IRD, the anamnesis (medical history) must include:
   - Inquiry about consanguinity, the existence of other affected family members, history of infectious diseases and other diseases, and pedigree.
   - The taking of general ophthalmological history, symptoms at onset, age at symptom onset, other previous visits to a general ophthalmologist or to a retina specialist, previous ophthalmological clinical assessment (see fundamental clinical diagnostic tests below), signs of disease progression previous, and current therapy for ocular diseases.
   - Presence or absence of neurological or extra-ocular symptoms, general pharmacological history and ongoing medical treatments, patient expectations, professional activities and lifestyle (for adult patients).

2. The role of clinical diagnosis in reaching a level of suspicion that justifies genetic testing in the presence of a hereditary retinal dystrophy was emphasised. RPE65-associated IRD should be suspected in individuals with the following clinical findings:
Symptomatic onset between birth and 5 years, nystagmus or roving eye movements, profound nyctalopia and decreased central VA.

• Fundus appearance that tends to be normal in infancy and then ranging from RPE mottling to pigmentary retinopathy with attenuated vessels and optic nerve pallor [37].
• Full-field electroretinogram (ERG) barely detectable or severely abnormal.
• Severely diminished or absent fundus autofluorescence (FAF) and a relatively preserved central retinal structure at optical coherence tomography (OCT) [38].

3. The order of clinical manifestations should be undertaken to give them priorities and weights depending on their frequency in patients and to characterise their importance for diagnosis. Night blindness, narrowing of the visual field and reduced VA are the three symptoms that are present; in infants, light staring (photoattraction) is characteristic of RPE65- or LRAT-associated IRDs. Nystagmus is often associated with this condition, but not all patients have it.

4. The clinical diagnosis is mainly LCA or EOSRD/ EORP. Nevertheless, RPE65 biallelic mutations can be also associated with Fundus albipunctatus (FA) a rare form of stationary night blindness [39].

5. Visual acuity may vary and is heterogeneous between patients at the onset. Generally, central VA is worse when the onset is before age 1 year compared with onset between 1 and 5 years of age.

6. Fundus examination can be quite variable and can appear normal at presentation.

7. Other fundus findings include RPE mottling, white dots at the level of the RPE, pigmentary retinopathy with attenuated vessels and optic nerve pallor.

8. The basic clinical diagnostic tests to guide towards a level of suspicion that justifies genetic testing (i.e. tests with the highest diagnostic value) are fundus examination, full-field electroretinogram (ERG), GVF and OCT. Tests for further investigation should include FAF, full-field light sensitivity threshold (FST) testing and microperimetry.

Order of testing for viable retinal cells (Additional file 2: Table S2)
The clinical tests able to assess viable retinal cells in order of priority are the following: (1) OCT, followed by (2) standard ophthalmological examinations including VA, posterior segment biomicroscopy, then (3) GVF testing, microperimetry, colour picture, FAF and full-field ERG.

Diagnostic genetic testing: Sanger sequencing versus targeted, multi-gene NGS panel (Additional file 3: Table S3)

1. Genetic testing must be prescribed and performed in all cases where there is suspicion of an inherited retinopathy. Genetic testing must be carried out by a certified laboratory affiliated with medical geneticists and inherited retinal diseases specialists from other national and international centres.

2. A strong consensus was reached on the utility of a next-generation sequencing (NGS) approach for the genetic diagnosis of potential RPE65-associated IRD cases over a Sanger approach limited to the analysis of the RPE65 gene. The consensus recommendation was to use a targeted multi-gene NGS approach, including all the genes known to be responsible for IRDs, both isolated and syndromic forms. The use of a larger panel (i.e. either a clinical exome or a whole-exome sequencing) is not excluded but, due to the issue of possible incidental findings, requires a more careful pre-test counselling.

3. Regarding segregation analysis, the issue of collecting a sample from the parents was identified and segregation defined as a necessary step.

4. The issue of variants of uncertain significance (VUS) is important. Genotypes including pathogenic and likely pathogenic variants constitute a straightforward eligibility indication for the treatment. However, genotypes, including VUS, provide more problems in that respect [40–46]. The panel agreed that they should not be excluded altogether but evaluated on a case-by-case basis. Among additional criteria that can possibly be considered when VUS arise:

Patient eligibility and prioritisation (Additional file 4: Table S4)

1. In general, it is important to give priority to the treatment of paediatric patients, in order to provide them with an opportunity for maintaining better VA (as they are likely to be in a less advanced stage of the disease) and enjoying normal social growth. In accordance with the current level of evidence from clinical trials, a paediatric patient should be consid-
Consideration of surgical issues (Additional file 5: Table S5)

1. On the basis of a comprehensive anamnesis suggestive of RPE65-associated IRD, patients must be referred to a clinical centre where ophthalmologists are specialised in the diagnosis of IRDs and have the ability to prescribe IRD genetic testing and genetic counselling from a certified laboratory with consolidated expertise in the genetics of IRD.

2. On confirmation of eligibility to voretigene neparvovec gene therapy by an IRD-specialised centre, the procedure must be carried out in a paediatric hospital setting that provides children-centred care appropriate for children aged 3–6 years and that is able to deliver and properly manage paediatric anaesthesiology procedures.

3. The vitreoretinal surgeon who performs the subretinal injection must have surgical experience with paediatric and adult patients with IRD and must be affiliated to an IRD centre certified for the use of gene therapy with voretigene neparvovec.

4. To corroborate patient eligibility for surgery and to assess the risks of subretinal injection, the retina specialist must discuss with the certified surgeon the expectations of the patient regarding the clinical outcome of the procedure while considering the age of the patient, retinal thickness, the presence of any other eye disorders, eligibility for general anaesthesia and an evaluation of the risk/benefit ratio of treatment.

5. The best surgical strategy for each patient candidate must be considered on an individual basis.

Diagnostic algorithm
The consensus findings formed the basis for a diagnostic algorithm that was constructed by the steering committee (Fig. 1). The algorithm is designed to illustrate the patient pathway leading from the patient’s referral centre all the way to the retinal specialist centre. The primary intention was to develop a simple and linear algorithm to guide paediatricians and general ophthalmologists who are not experts in this area to arrive at the correct diagnosis of RPE65-associated IRD and make informed clinical decisions regarding eligibility for a gene therapy approach to RPE65-associated IRD. The summary algorithm will ultimately be developed into an expanded form by incorporating supplementary material that will detail the decision-making process presented in the summary algorithm.

Discussion
The aim of this process was to provide practical guidance for the correct diagnosis of patients with RPE65-associated IRD, to identify the fundamental clinical tests to determine the correct diagnosis and to confirm patients eligible to receive treatment with voretigene neparvovec. The introduction of voretigene neparvovec as a gene supplementation therapy for RPE65-associated IRD is considered a milestone in the field of IRD. The pivotal phase 3 trial showed that a single sub-retinal injection of voretigene neparvovec is able to produce significant improvements in bilateral multi-luminance mobility test scores compared with controls at 1 year, with the beneficial effects maintained during the currently available 4 years of the planned 15 years of follow-up [48]. Furthermore, a recent evidence-based review by the UK National Institute for Health and Care Excellence (NICE) determined that the clinical benefits of voretigene neparvovec were important and represented a step change towards meeting the high unmet needs that have until now existed in RPE65-mediated IRD [49].
The heterogeneity and relative rarity of IRDs are further accentuated in \textit{RPE65}-associated IRD \cite{50}, and present challenges for practising ophthalmologists who may have little exposure to orphan diseases. Although not specific to Italy alone, the limited number of qualified retinal specialist centres and a lack of access to the molecular resources necessary for diagnosis further compounds the issue of ensuring optimal treatment of \textit{RPE65}-associated IRD. For example, a recent multinational survey by the European Vision Institute Clinical Research Network

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Fig. 1 Clinical pathway algorithm to evaluate eligibility for voretigene neparvovec gene therapy. *Refer to Table 1 for further information. FAF fundus autofluorescence, EORP early-onset retinitis pigmentosa, ERG electroretinography, FA Fundus albipunctatus; FST full-field light sensitivity threshold; IRD inherited retinal disease, LCA Leber congenital amaurosis, NGS next-generation sequencing, OCT optical coherence tomography, RP retinitis pigmentosa, RPE retinal pigment epithelium, VA visual acuity, GVF Goldmann visual field, VN voretigene neparvovec, VR vitreoretinal
benefit/risk profile of voretigene neparvovec could not be
or treatment response were identified, suggesting that the
between mutation subtype and baseline visual function
in 29 patients in the phase 3 study [41]. No correlations
neparvovec therapy, and adverse events were analysed
type and baseline visual function, response to voretigene
eration [40–46]. Correlations between mutation sub-
tions of treatment and the necessity to comply with
f.'s lives, optimising access to an
a life-
time beneficial effect is an important consideration. The
panel members reached valuable consensus findings that
direct attention to the relevant issues in identifying can-
candidate patients for gene supplementation with voreti-
gene neparvovec while clarifying matters that may be
misunderstood or are of lesser relevance. The taking of a
comprehensive anamnensis to establish the priorities and
tribution of the key clinical manifestations is critical,
and night blindness, reduced VA and narrowing of the
ed the three symptoms that are always present in RPE65-associated IRD. The
diagnosis, which is mainly LCA or EOSRD/EORP, needs
to be established using fundamental clinical diagnostic
tests (i.e. fundus examination, ERG, GVF examination
and OCT), supported by FAF, FST and microperimetry
for further investigation. Conducting a thorough assess-
ment to ensure that the patient understands the implic-
tions of treatment and the necessity to comply with
follow-up assessments post-treatment is also of the high-
est importance.

The role of clinical diagnosis is to reach a level of sus-
picion that justifies genetic testing in the presence of an
IRD. There was a strong consensus for the value of tar-
targeted multi-gene NGS for diagnostic genetic testing, in
ference to single-gene Sanger sequencing. Variants of
uncertain significance (VUS) were identified as impor-
tant in the diagnostic process and justify evaluation on
a case-by-case basis rather than exclusion from consid-
eration [40–46]. Correlations between mutation sub-
type and baseline visual function, response to voretigene
eparvovec therapy, and adverse events were analysed
in 29 patients in the phase 3 study [41]. No correlations
between mutation subtype and baseline visual function
or treatment response were identified, suggesting that the
benefit/risk profile of voretigene neparvovec could not be
predicted by mutation subtype [41].

The treatment of paediatric patients is a high priori-
ity in order to take full advantage of the opportunity for
maintaining better VA and improving patients’ psycho-
logical well-being, relationships and family life. The safety
and efficacy of voretigene neparvovec in children aged
up to 4 years have not been established in clinical trials,
although age in itself should not be a reason to exclude
gene therapy. In the voretigene neparvovec clinical
development programme (n=41 patients), the average
age of included patients was 17 years, ranging from 4 to
44 years of age. Of the 41 patients, 25 (61%) were paedia-
tric patients under 18 years of age [28].

We identified a strong consensus that voretigene
eparvovec should be initiated and administered by a
vitreoretinal surgeon experienced in performing sub-
retinal/macular surgery in association with a centre spe-
cialised in managing patients with IRD and where there
is pharmacy capability for the handling of gene therapies.
Surgeons and pharmacists from the treatment centres
who meet these Risk Management Plan (RMP) criteria
need to attend mandatory surgery/pharmacy education
sessions in order to ensure the correct use of voretigene
eparvovec and to minimise the risks associated with its
administration and/or the administration procedure [28,
32].

Regarding the evaluation of viable retinal cells, based
on the mechanism of action of voretigene neparvovec,
the presence of sufficient viable retinal cells is consid-
ered necessary for therapeutic efficacy [52]. Although
there are no universally shared criteria to unques-
tionably establish this viability, studies in the clinical develop-
ment programme defined 101/102 and 301/302 approval
trials [26, 28] defined sufficient viable retinal cells as an
OCT showing more than a 100-μm thickness in an area of
retina within the posterior pole and/or retina without
atrophy or pigmentary degeneration within the poste-
rrior pole equal or larger than 3 disc areas and/or remain-
ing GVF within 30 degrees of fixation as measured by
a Goldmann isopter III-4e or equivalent. In this Delphi
project, a consensus was reached that the most appro-
iate approach to assess viable cells, through a structural
and functional evaluation, is to perform the clinical tests
separately according to the sequence: OCT, followed by
standard ophthalmological examinations including VA,
posterior segment biomicroscopy, then GVF testing,
microperimetry, colour picture, FAF and full-field ERG.
The OCT should be the preferred and first-performed
assessment to confirm eligibility for treatment. Indeed,
thickness measurements on OCT served as an inclu-
sion criterion in the phase 3 clinical programme [28], to
estimate whether sufficient viable retinal cells were pre-
sent for treatment. Of note, a correlation between OCT
findings and therapeutic efficacy of voretigene neparvovec has not been reported [52].

This study brought together the expertise of a multidisciplinary panel representative of experts experienced in the management of IRDs in Italy and whose support was demonstrated by the high response rate achieved during the two Delphi rounds. Although the Delphi process is not designed to result in the highest level of evidence, it is an appropriate means of gaining consensus within a community, particularly when conducted by a fully representative panel of experts and supported by a comprehensive literature review of the subject. In the absence of definitive evidence-based literature and treatment guidelines on this rare disease, our Delphi process obtained valuable information that provides the basis for clinically useful guidance of a more focused and specific nature than the more general guidelines on the assessment and management of patients with IRD [53, 54]. Recommendations that set out the conditions to be met prior to performing treatment with, and to optimize outcomes with, voretigene neparvovec are largely in agreement with those in the present paper. However, we feel that our rigorous stepwise Delphi process consensus approach by an interdisciplinary panel of experts, supported by an extensive review of the literature, strengthens the recommendations presented here. The statements presented in the Round 1 questionnaire of our study were carefully constructed to avoid ambiguity, validated before distribution, and appropriately modified for Round 2 based on the Round 1 responses. Analysis of the Round 1 and Round 2 responses was by an expert steering committee with considerable experience in the field of IRDs, and a consensus was achieved across a wide range of issues.

Conclusions

To summarise, this paper provides robust, evidence-based and consensus-driven guidelines that can be used by ophthalmologists and paediatricians to arrive at the correct diagnosis of RPE65-associated IRD and to make informed clinical decisions about eligibility for gene supplementation with voretigene neparvovec. This is particularly important in a rare condition, such as RPE65-associated IRD, where progressive visual impairment has a distressing impact on many aspects of patients’ lives. Although voretigene neparvovec makes available to clinicians a treatment with the potential to prevent blindness and provide a life-long beneficial effect for many patients with RPE65-associated IRD, its safe and effective use requires expertise across a multidisciplinary team that includes ophthalmologists, geneticists, surgeons, and patient support services.

The process of securing consensus among representatives of experts in the management of patients with IRDs has facilitated the development of a practical diagnostic algorithm to guide paediatricians and general ophthalmologists encountering IRDs in clinical practice. Although there are currently unmet needs regarding genetic counselling in RPE65-associated IRD, the limited availability of genetic testing and the long turnaround time in clinical practice, we consider that the information here presented can serve as a framework of care for the optimal treatment of these patients.

Abbreviations
cGH: Comparative genomic hybridisation; FST: Full-field light sensitivity threshold; EORP: Early-onset retinitis pigmentosa; EOSRD: Early-onset severe retinal dystrophy; ERG: Electroretinography; FA: Fundus albipunctatus; FAF: Fundus autofluorescence; GVF: Goldmann visual field; IRD: INHERITED retinal disease; LCA: Leber congenital amaurosis; MLMT: Multiluminance mobility test; MLP: Multiplex ligation-dependent probe amplification; MRI: Magnetic resonance imaging; NGS: Next-generation sequencing; OCT: Optical coherence tomography; RMP: Risk Management Plan; RP: Retinitis pigmentosa; RPE: Retinal pigment epithelium; SIGU: Società Italiana di Genetica Umana; VA: Visual acuity; VN: Voretigene neparvovec; VR: Vitreoretinal; VUS: Variants of uncertain significance.

Supplementary Information

The online version contains supplementary material available at https://doi.org/10.1186/s13023-021-01868-4.

Additional file 1: Table S1. Clinical manifestations of RPE65-associated IRD. Provides an overview of the consensus during the Round 1 and Round 2 questionnaires on statements relating to the clinical manifestations of RPE65.

Additional file 2: Table S2. Viable retinal cells: structure and function. Provides an overview of the consensus during the Round 1 and Round 2 questionnaires on statements relating to the structure and function of retinal cells.

Additional file 3: Table S3. Genetic testing. Provides an overview of the consensus during the Round 1 and Round 2 questionnaires on statements relating to genetic testing for IRD.

Additional file 4: Table S4. Patient eligibility and prioritisation. Provides an overview of the consensus during the Round 1 and Round 2 questionnaires on statements relating to patient identification and prioritisation of treatment.

Additional file 5: Table S5. Assessment of surgical strategy. Provides an overview of the consensus during the Round 1 and Round 2 questionnaires on statements relating to surgery for IRD.

Additional file 6: Appendix. Members of the panel of experts. List of all members of the panel of experts who participated in the Delphi consensus process.

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