

## POLICY

# EMQN/CMGS best practice guidelines for the molecular genetic testing of Huntington disease

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Huntington disease (HD) is caused by the expansion of an unstable polymorphic trinucleotide (CAG)<sub>n</sub> repeat in exon 1 of the *HTT* gene, which translates into an extended polyglutamine tract in the protein. Laboratory diagnosis of HD involves estimation of the number of CAG repeats. Molecular genetic testing for HD is offered in a wide range of laboratories both within and outside the European community. In order to measure the quality and raise the standard of molecular genetic testing in these laboratories, the European Molecular Genetics Quality Network has organized a yearly external quality assessment (EQA) scheme for molecular genetic testing of HD for over 10 years. EQA compares a laboratory's output with a fixed standard both for genotyping and reporting of the results to the referring physicians. In general, the standard of genotyping is very high but the clarity of interpretation and reporting of the test result varies more widely. This emphasizes the need for best practice guidelines for this disorder. We have therefore developed these best practice guidelines for genetic testing for HD to assist in testing and reporting of results. The analytical methods and the potential pitfalls of molecular genetic testing are highlighted and the implications of the different test outcomes for the consultand and his or her family members are discussed.

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## DESCRIPTION OF THE DISEASE

Huntington disease (HD, OMIM #143100) is a progressive neurodegenerative disorder that presents with motor symptoms, cognitive impairment and psychiatric disturbances. The first symptoms usually manifest between 35 and 50 years of age and the duration of the disease is between 15 and 20 years.<sup>1,2</sup> A small number of cases present before the age of 20 (juvenile onset) and about 25% of cases present after 50 years of age (OMIM). HD is inherited in an autosomal dominant fashion with an incidence of 3–10 in 100 000 in populations of Western European descent. It is much less frequent (0.1–0.4: 100 000) in other populations.<sup>1</sup>

HD is characterized pathologically by loss of specific neuronal populations in many brain regions, although the pathology is not limited to neurons. Neuropathological features include selective degeneration of neurons in the caudate and putamen and less severe loss in the cerebral cortex. More detailed information on the pathogenesis can be found in Ross and Tabrizi.<sup>3</sup> Individuals homozygous for HD expansions appear to have a similar age of onset, but may exhibit an accelerated rate of disease progression.<sup>4</sup>

## THE GENE AND THE MUTATIONS

The gene involved, the huntingtin (*HTT*) gene (NM\_002111.6; NG\_009378.1) previously known as IT15, is located on chromosome 4p16.3, spans 180 kb and consists of 67 exons. The *HTT* gene is widely expressed and is required for normal development. It is expressed as two alternatively polyadenylated forms displaying different relative abundance in various fetal and adult tissues. The larger transcript is approximately 13.7 kb and is expressed

predominantly in adult and fetal brain, whereas the smaller transcript of approximately 10.3 kb is more widely expressed.<sup>5</sup>

HD is caused by the expansion of an unstable polymorphic trinucleotide (CAG)<sub>n</sub> repeat in exon 1 of the *HTT* gene, which translates into an extended polyglutamine tract in the protein. Alleles with <27 CAG repeats are classified as normal, whereas alleles with ≥36 repeats are detected in patients (see Table 1). So far, the smallest number of CAG repeats described in patients with confirmed clinical features of HD is 36. Alleles with 27–35 repeats (called mutable normal or intermediate alleles) are not associated with disease symptoms but can expand into the affected range upon (predominantly paternal) germline transmission and thus cause HD in offspring. Repeats of 36–39 CAG are incompletely penetrant and can be found in affected individuals as well as individuals who show no clinical symptoms in an advanced age (≥70–80 years). The CAG-repeat number correlates inversely with the mean age of onset of symptoms. Generally, individuals with longer CAG repeats have an earlier age of onset. This is supported by findings that individuals with very large CAG repeats (>60) present with juvenile HD, and individuals with shorter CAG repeats (36–39) can remain asymptomatic. However, the number of repeats accounts only for approximately 70% of the variance in age at onset.<sup>6</sup> This implies that not all juvenile cases have ≥60 repeats. Numerous predicting models discussing the statistical relationship between the CAG-repeat length and the age of onset have been published over the last 15 years. A review and validation study of these statistical approaches can be found in Langbehn *et al.*<sup>7</sup>

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**Table 1** Summary of the implications of different repeat ranges for the individual tested and family members

| Number of repeats                      | Implications for individual/patient                              |  |   |
|--|--|--|---|
|  | Diagnostic test  | Predictive test  | Implications for family members                     |
| 6–26 normal allele                     | Diagnosis not confirmed or diagnosis of HD excluded              | Will not develop HD  | No increased risk for HD                            |
| 27–35 intermediate allele              | Diagnosis not confirmed or diagnosis of HD excluded <sup>a</sup> | Will not develop HD <sup>b</sup>                             | Increased risk for HD<br>(few %; <10%) <sup>c</sup> |
| 36–39 incomplete penetrance allele     | Diagnosis of HD confirmed  | May or may not develop HD;<br>in range of reduced penetrance | Increased risk for HD                               |
| 40 and over complete penetrance allele | Diagnosis of HD confirmed  | Will develop HD  | Increased risk for HD                               |

<sup>a</sup>At the higher end of the range one should use: 'diagnosis of HD is very unlikely'.<sup>b</sup>At the higher end of the range one should use: 'it is very unlikely that the consultand will develop HD'.<sup>c</sup>The figure depends on the type of intermediate allele (IA). General population IAs have a much lower (<1%) risk than IA that have already shown an expansion in the family.

Huntingtin is a protein of 3144 amino acids with a predicted molecular mass of 348 kD. The polyglutamine tract starts at residue 18 (c.52 p.18) and, when abnormally expanded, is thought to acquire a novel deleterious function. This eventually leads to neuronal dysfunction and neurodegeneration. The polyglutamine expansions also result in the formation of neuronal intranuclear inclusions containing huntingtin and ubiquitin.

## REPEAT SIZE RANGES

### Normal alleles

The CAG repeat is highly polymorphic in the population. Alleles in the range of 6–26 CAG have never been found to be associated with HD and there has been only one single report about a normal allele that was unstable on transmission to the next generation.<sup>8</sup>

### Disease size range

The smallest number of CAG repeats described in patients with confirmed clinical features of HD is 36.<sup>9,10</sup> This repeat size has been observed in more than six documented HD cases. Although it is possible that HD cases with fewer repeats exist, only few ( $n=5$ ) possibly affected cases with 29–34 repeats have been reported.<sup>11–14</sup> Such cases are difficult to ascertain due to the paucity of alleles of this size. Although we cannot exclude the possibility that HD may be associated with these smaller repeat sizes, the published data are not conclusive and alternative diagnoses need to be carefully considered in such cases with HD-like signs.

### Range of reduced penetrance

There have been many reports with elderly asymptomatic individuals with HD alleles of 36–39 repeats which define the range of incomplete or reduced penetrance of the mutation.<sup>9,10,15,16</sup> Many carriers of these alleles remain asymptomatic until a very late age, although exceptions exist.

### Intermediate or mutable normal alleles

Inconsistencies and confusion exist in literature regarding intermediate alleles despite the existence of published guidelines<sup>17</sup> (personal observations during external quality assessment (EQA) for European Molecular Genetics Quality Network (EMQN)). These alleles of 27–35 repeats have been designated 'intermediate' and are also referred to as 'mutable normal' or 'large normal' alleles. They are defined as 'being below the affected range but having the potential to expand into the disease range in the next generation'. The lower limit is defined as the shortest size ever reported to expand into the HD range after one

transmission. Intermediate alleles are relatively common in the general population with frequency estimates between 1 and 7%.<sup>17,18</sup>

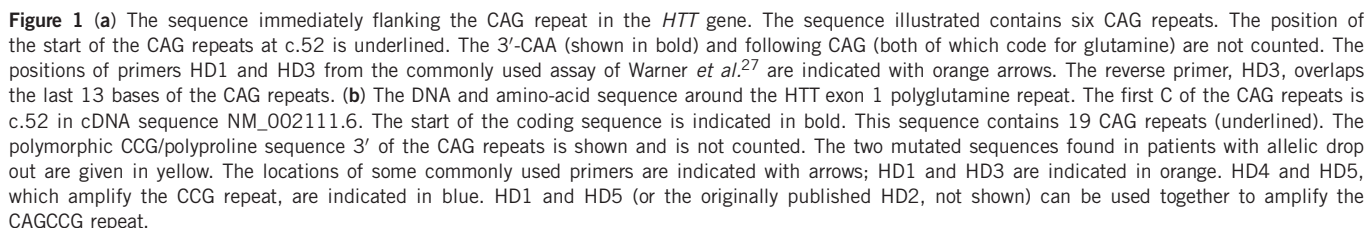
The risk of expansion of an intermediate allele into the disease range has been estimated at 0.1–1% per generation,<sup>18,19</sup> (see also next paragraph). Individuals who are found to carry alleles in this range should be counseled about the possibility of prenatal diagnosis and the risk for other family members. The small risks are difficult to quantify but may be more significant for males transmitting alleles in this range. Where intermediate and pathogenic alleles are found in the same individual, the interpretation should clearly distinguish the risks associated with each allele.

## REPEAT INSTABILITY

Although the rate of new mutations in HD is low, these have been described and originate from intermediate alleles of 27–35 repeats, mainly through paternal transmission.<sup>20,21</sup> Up to now, only one maternal case has been documented.<sup>22</sup> Factors influencing repeat instability include size of the allele, sex and age of the transmitting parent, family history and gene sequence and haplotype on which the intermediate allele is located. Alleles with a larger number of CAG repeats have a higher likelihood of expanding into the affected range upon transmission to the next generation.<sup>17</sup> The risk that offspring will develop HD is the highest for fathers carrying these alleles who are >35 years of age.<sup>23</sup> Intermediate alleles can be coincidentally identified when healthy family members in an HD family undergo CAG-repeat sizing ('general population intermediate alleles'). In contrast, they can also be ascertained from new mutation families and these new mutation alleles are more prone to repeat expansion when compared with similar-sized alleles in the general population. The genetic variability near the repeat tract can influence its stability. The c.7934\_7936del p.Glu2645del (known as the  $\Delta 2642$ ) polymorphism is vastly overrepresented on HD chromosomes (38%) relative to the general population (7%). Finally, repeat tracts in which the 3'-CAA repeat has changed to a CAG are markedly more unstable (see Figure 1a). Both polymorphisms tend to cluster in specific haplotypes.<sup>24</sup> Additional information is needed from various populations to determine a more precise and critical risk assessment for offspring of intermediate allele carriers.

## Anticipation

Anticipation is the phenomenon in which increasing disease severity and/or decreasing age of onset is observed in successive generations. In HD, it occurs more commonly through paternal transmission. The phenomenon arises because of the expansion of the unstable CAG repeat during spermatogenesis. Also large expansions, for



member with HD is advisable. Presymptomatic testing of minors (<18 years of age) is considered appropriate only in exceptional circumstances.<sup>25,26</sup> Comprehensive genetic counseling is required, and informed consent must be provided in accordance with local practices.

According to the international guidelines, predictive testing for HD should only be requested by a clinical geneticist.<sup>25,26</sup> Presymptomatic testing can be offered to individuals at an *a priori* 50 or 25% risk. On very rare occasions, an individual at 12.5% risk may be tested. Confirmation of HD at the molecular level in at least one family

Prenatal testing should be requested by a clinical geneticist. The testing method is identical to presymptomatic testing with some exceptions (see section on analytical methods). Preimplantation genetic diagnosis (PGD) may be a valuable option for couples who wish to avoid transmitting the disease without revealing their own status or to avoid termination of a pregnancy. It is only offered in a limited number of specialized centers and will not be discussed here (see also Rij van *et al*<sup>27</sup>).

In cases where an individual with a 25% risk of developing HD requests predictive testing but the parent, who has a 50% risk, does not want to be informed about his or her risk, an exclusion test can be offered. This test is nowadays almost exclusively used in prenatal testing where the fetus is at 25% risk and the future parent at 50% risk. It is performed with linked polymorphic markers (see also interpretation section). Exclusion testing may not be available in all countries.

The number of repeats is established by PCR analysis of the region encompassing the CAG repeat (Figure 1a), followed by fragment

sizing through capillary or gel electrophoresis at sufficient resolution to allow separation of alleles with one repeat difference.<sup>28,29</sup> Other methods with comparable resolution can also be applied. Regardless of the PCR-based strategy selected, it is important that the assay conditions are optimized to ensure the accurate and unambiguous determination of the number of repeats. Laboratories must validate the test by measuring allele sizes and calculate uncertainty for their tests. Genotyping results from the yearly EQA scheme for molecular genetic testing of HD from the EMQN show that over the last 3 years (2008–2010), 3–9% of alleles fall outside the error limits set by the EMQN. For 2010, 51 out of 570 alleles were typed outside the error limits (which, at the time, were set at  $\pm 1$  for alleles  $< 40$  repeats, and  $\pm 3$  repeats for alleles  $> 39$  CAG repeats). In several cases, this was because of the fact that CAGCCG repeats are reported instead of CAG repeats. Two other studies reported similar results.<sup>30,31</sup>

### Controls

PCR products containing CAG repeats migrate anomalously in electrophoresis, making conventional size ladders unreliable. Instead, control samples with well-defined repeat sizes at the borders of normal, intermediate, reduced penetrance and expanded repeats, as well as a large expansion ( $> 60$  repeats), should be used for allele sizing. The number of repeats in these controls should preferably be determined by DNA sequencing and/or by calibration against a certified reference material (see below).

A prenatal test is technically identical to a presymptomatic test but the maternal or both parental samples should be analyzed in the same run. If the fetal genotype is identical to the maternal genotype, maternal contamination should be excluded using polymorphic microsatellites or equivalent.

### Reference materials

Certified reference materials are essential aids to the accurate and traceable calibration of measurement systems. The only certified reference material available for HD testing is SRM 2393 from the US National Institute of Standards and Technology. This panel of six genomic DNAs contains alleles ranging from 15 to 75 CAG repeats characterized by Sanger sequencing. A useful panel of well-characterized DNAs is available from the Coriell CDC Repository.<sup>32</sup> A cell line has been established from a patient shown by sequencing to carry 24 and 35 repeats. This cell line (or DNA prepared from it) is available from the ECACC (<http://www.ecacc.org.uk/>). It is cell line number CM0034, ECACC ref. no: 95090133.

### Sizing accuracy

It is the laboratory's responsibility to empirically determine the error limits (precision) of their assay. Acceptable error limits are  $\pm 1$  repeat for alleles  $\leq 42$  and  $\pm 3$  repeats for alleles  $> 42$ . The CAG repeat in the *HTT* gene is adjacent to a 3'-positioned CCG repeat, which is also polymorphic in length (Figure 1b).<sup>6,29</sup> The original primer sets, used to size the CAG repeats, also included this CCG polymorphism and can thus result in misclassification of alleles.<sup>33</sup> Therefore, this assay should not be used for routine sizing of the CAG repeats, but may be very helpful to resolve two homozygous normal HD alleles with identical numbers of CAG repeats but different numbers of CCG repeats.

Annual participation in an EQA scheme is not only necessary to allow the comparison of allele sizing with other laboratories but also for quality assurance, continuous validation, evaluation of reporting and continuous education. HD testing should only be performed in

laboratories that are accredited according to ISO 15189 or equivalent.<sup>34</sup>

To exclude the presence of a very large expansion that would be missed by PCR in a case with homozygosity of a normal CAG-repeat allele, the use of a southern blotting protocol (*Pst*I-digested DNA hybridized with probe 4G6P1.7) or TP-PCR is recommended (also see the next paragraph).<sup>28</sup> PCR of large ( $> 100$ ) CAG-repeat tracks in DNA from fresh or frozen tissue samples is feasible.<sup>35</sup> Because individuals with  $> 60$  repeats will present with a juvenile or early-onset form of HD, it will be unlikely that adult onset cases are missed by PCR analysis alone.

For homozygous (normal) CAG-testing results, several approaches may be taken:

1. As mentioned above, the CAGCCG repeat should be genotyped, which might demonstrate the presence of two normal alleles (with identical CAG-repeat counts but heterozygosity for the CCG repeat)
2. Samples homozygous for CAG and CCG repeats should be analyzed alongside an appropriate large CAG-repeat positive control (demonstrating the sensitivity of the test for larger CAG expansions)
3. In a symptomatic test setting, the age at onset of symptoms should be available and discussed as to whether young onset HD can be excluded clinically; that is, if the patient has onset in his/her 40 or 50s, a very large expansion is most unlikely.
4. In a presymptomatic test setting, although the proband's age can be taken into account, it may be advisable to offer genotyping for the proband's parents in order to confirm homozygosity or to check allele segregation by STR analysis.
5. In the juvenile HD setting, an apparent homozygous (normal) CAG result has to be treated with caution unless segregation analysis with polymorphic markers has indicated the presence of two normal alleles. If parental samples are not available, southern blotting or TP-PCR should be considered.<sup>36</sup>

Three very rare polymorphisms have been described in the 3' or 5' primer used in some CAG-specific assays.<sup>37,38</sup> These can potentially disrupt primer binding to an HD chromosome and result in the failure to amplify a pathogenic allele.<sup>37</sup> Table 2 lists the factors that can influence the analytical specificity and sensitivity of the CAG-sizing assay. Methods of mitigating false-positive and false-negative results are detailed. Finally, in some countries, it is common practice for presymptomatic tests to ask for a second independent blood sample to confirm the result. In general, measures to avoid sample swaps should be implemented in every laboratory offering a diagnostic service.

### CLINICAL SENSITIVITY AND SPECIFICITY

The absence of HD pathology in an individual with  $\geq 40$  repeats who died after living up to or past normal life expectancy has never been described. Therefore, a result of  $\geq 40$  repeats is a 100% diagnostic of HD.<sup>44</sup> CAG-repeat expansions account for  $> 99\%$  of cases of HD and, therefore, the test with a result of  $\geq 40$  repeats and all appropriate controls is  $> 99\%$  sensitive.<sup>44</sup>

### INTERPRETATION OF THE RESULTS AND REPORTING

The interpretation should always be done in the context of the clinical referral and is summarized in Table 1. It is important to make a clear distinction between a report for a diagnostic test on a patient with symptoms of HD and a report for presymptomatic testing. One has



**Table 2** Factors affecting diagnostic parameters in HD testing

| Parameter   | Possible risks (pitfalls)  | Technical requirements   |
|---|--|--|
| <p>Analytical sensitivity</p> <p>Sensitivity: The proportion of biological samples that are rightfully classified as positive by the test (= 1 – proportion of false-negative results).</p> | <p>As a sensitive HD test is designed to identify CAG-repeat expansions of unknown size, there is a risk of missing true positives (allelic drop out (ADO)).</p> | <ul style="list-style-type: none"> <li>• If the PCR test reveals two normal alleles in the sample, a false-negative result can be ruled out.</li> <li>• The PCR assay must have the capacity to detect alleles in the normal and expanded ranges (up to 100 repeats), and resolve alleles one repeat apart</li> <li>• Include positive control (&gt;60 repeats) in every experiment (or allelic ladder)</li> <li>• Primer design is crucial<sup>a</sup></li> <li>• Testing with primers spanning the CAGCCG repeat will help to distinguish two normal alleles in most cases</li> <li>• TP-PCR or southern blotting will exclude or confirm the presence of a large expansion</li> <li>• Exceptionally, an answer can be sought with segregation analysis with linked STR markers in extended pedigrees</li> <li>• Only a few observations of polymorphisms at primer-binding sites are reported<sup>b</sup></li> <li>• Caution with automated allele detection tools: expansion-related stutters can result in small peak heights. Ensure the electropherogram extends far enough to visualize large expansions.</li> </ul> |
| <p>Analytical specificity</p> <p>Specificity: The proportion of biological samples that is rightfully classified as negative by the test (= 1 – proportion of false-positive results)</p>   | <p>A pathological CAG expansion in the <i>HTT</i> gene is the only biological reason for HD</p>  | <ul style="list-style-type: none"> <li>• Electropherograms should be monitored thoroughly for the presence of triplet stutter peaks in order not to mistake electrophoresis artifacts for HD alleles.</li> <li>• The signals of PCR-amplified HD mutations in capillary electrophoresis represent a highly specific pattern, which is distinct from any other technical PCR or electrophoresis artifacts.</li> <li>• Standard negative controls (blank without DNA template) are sufficient</li> </ul>   |

<sup>a</sup>Published primers: see refs 33,39,40<sup>b</sup>See refs 20,37,38,41–43

to bear in mind that the result is not only important for the consultand tested but is also relevant for the family members. Each laboratory has its own reporting format, but general reporting guidelines can be found on the EMQN website [www.emqn.org](http://www.emqn.org) linked to the CMGS guidelines for reporting ([http://www.cmgs.org/BPGs/Best\\_Practice\\_Guidelines.htm](http://www.cmgs.org/BPGs/Best_Practice_Guidelines.htm)). One should also adhere to the ISO 15189 and OECD guidelines.<sup>34</sup> A one-page report is the preferred format in which the test result and the answer to the clinical question should be easy to find and unambiguously formulated. The report is a stand-alone document that should not only be clear to the referring physician but also to other professionals involved in supporting and/or treating the patient. The reports should carry a clear key including a short clinical interpretation of the different size ranges. Alleles at the boundaries of size ranges should be determined precisely by the use of appropriate reference materials and/or by use of the most precise method available; if the resultant genotype still includes more than one range (eg,  $35 \pm 1$  CAG repeats), this should be reflected in the interpretation. The answer to the clinical question, the *take-home message*, should be stated clearly and unambiguously: for example, 'The diagnosis of HD either *is* or *is not* confirmed.' 'The consultand *will* or *will not* develop HD'. Although local policy can vary with regard to reporting, some relevant HD-specific items are mentioned below.

### Nomenclature

The number of CAG repeats is measured as the *number of uninterrupted CAG repeats* (see Figures 1a and b). There are more glutamines than CAG repeats in this part of the gene, as the polyglutamine tract is coded for in most chromosomes by (CAG)<sub>n</sub> CAACAG (CAA also codes for a glutamine). The HGVS nomenclature is not considered the most appropriate for reporting the results

of trinucleotide repeat analysis. Instead, the number of uninterrupted CAG repeats is used as a result of the genetic test. The error limits of the test should always be mentioned. Reporting actual allele sizes is subject to local practice. A statement whether the allele is in the normal or the disease range can very well be sufficient. However, it is important to report actual allele sizes if this is relevant for the interpretation of the result; for instance, in the case of a reduced penetrance or intermediate allele, as well as in the upper range of expansions where onset might be at a juvenile or infantile age.

### Diagnostic testing

In the case of a CAG repeat  $\geq 36$  the diagnosis of HD is confirmed (or is consistent with a diagnosis of HD). In the case of a CAG repeat  $\leq 35$ , the diagnosis is excluded (or the result is not consistent with HD). However, recently a few possibly affected cases with 29–34 repeats have been reported.<sup>11–14</sup> Alleles in the reduced penetrance or intermediate (normal mutable) range have implications for family members, which should be discussed in the report. In these cases and in the case of a confirmation of HD, the family should be offered a referral for genetic counseling. In the case of a test result with two alleles in the normal range when disease symptoms are clearly present, the referring physician could be advised to consider reevaluating the clinical diagnosis and testing for HD-like diseases, such as HDL1 (PRNP gene), HDL2 (JPH3 gene), DRPLA (ATN1 gene) or SCA17. This may be mentioned in the report in more general terms.

### Predictive (presymptomatic) testing

In cases of a CAG repeat from 36 to 39, the individual is at risk of developing HD; in cases with a CAG repeat  $\geq 40$ , the individual will develop HD (or is at risk of developing HD). In cases of a CAG repeat  $\leq 35$ , the individual will not develop HD (or is not at risk of

developing HD). Implications of alleles in the reduced penetrance or intermediate (normal mutable) range should be discussed in the report. Where available and appropriate, prenatal testing and PGD should be offered. Data in which the likelihood that an individual with a particular size CAG repeat will be affected by a specific age have been published.<sup>7,16,45</sup> Although more accurate predictions can now be made, and these models are being used for recruitment for clinical trials, the 95% confidence intervals are still wide and they have not been validated for genetic counseling. Thus, extreme care should be taken applying this information in individual presymptomatic cases. In some cases of presymptomatic testing, the familial diagnosis is not confirmed by genetic testing. In such cases, an important caveat must be mentioned: a negative result excludes HD, but it does not necessarily exclude the familial disorder, which might be misdiagnosed. This must always be emphasized in the laboratory report.

### Prenatal testing

The interpretation and approach to reporting a prenatal test is essentially the same as a presymptomatic test. However, it should be immediately clear from the report that it concerns a prenatal test. In addition, if the genotype of the fetus is identical to the maternal genotype, maternal contamination must be excluded and the result reported.

### Exclusion test

If a person with an affected grandparent wants to know whether he/she has an increased risk of developing HD, but their linking parent does not want to know his or her status, an exclusion test can be proposed. Nowadays, it is almost exclusively used as a prenatal test. The test result reveals whether the fetus (individual) with an *a priori* 25% risk received the grandparental risk haplotype or not (see also reason for referral). If the grandparental risk haplotype is present, the risk for the fetus is increased to approximately 50%, without changing the risk for the 50% at risk parent. If the grandparental risk haplotype is not present, the fetus' risk is reduced to almost 0% (the recombination frequency between the markers and the gene should be taken into consideration). It is necessary to include DNA of at least one grandparent in the analysis. Confirmation of the diagnosis at the molecular level in at least one family member, preferably the grandparent with HD, is necessary. For the exclusion test, polymorphic markers in the *HTT* region on chromosome 4p16.3 can be used (see Table 3 for a list of STRs). Ideally, markers both proximal and distal to *HTT* need to be informative to have a secure result. It is

advisable to determine beforehand as to which markers are informative for the particular family under study. A pedigree with the result of the haplotype analysis should be included in the report.

### CONCLUDING REMARKS

These guidelines are based on the knowledge acquired from peer-reviewed and published data. The authors received additional information from several sources, which could not be included in the guidelines as it was unpublished. We encourage the diagnostic and scientific community to publish instructive cases or data sets to expand our knowledge of HD and its diagnosis. This will lead to a continuous improvement of our diagnostic services. These guidelines can only provide a snapshot of current knowledge at the time of publication. Readers are advised to keep up with the literature.

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### INTERNET RESOURCES

EMQN website [www.emqn.org](http://www.emqn.org): GeneReviews funded by NIH. Developed at the University of Washington, Seattle, USA <http://www.ncbi.nlm.nih.gov/bookshelf/br.fcgi?book=gene&part=huntington>: NCBI RefSeq Gene project [ftp://ftp.ncbi.nlm.nih.gov/refseq/H\\_sapiens/RefSeqGene/](http://ftp.ncbi.nlm.nih.gov/refseq/H_sapiens/RefSeqGene/): European Collection of Cell Cultures <http://www.ecacc.org.uk/>: Cell line number CM0034, ECACC ref. no: 95090133. European Collection of Cell Cultures, 2010.: CDC Centers for Disease Control and prevention Genetic testing reference materials program <link to GeT-RMs>: Coriell Cell and DNA repository <http://ccr.coriell.org/sections/collections/>: Clinical Molecular Genetics Society Reporting Guidelines: <http://www.cmgs.org/BPGs/Reporting%20guidelines%20Sept%202011%20APPROVED.pdf>

**Table 3 Suggested markers in 4p16.3 that are suitable for exclusion testing**

| Marker  | UniSTS | Ensembl<br>marker | Ensembl<br>location (hg19) | Het<br>frq | Alias<br>(Ref)     |
|---------|--------|-------------------|----------------------------|------------|--------------------|
| D4S2936 | 24920  | Z52740            | 692247-692420              |            |                    |
| D4S3038 | 42100  | Z51777            | 1099931-1100155            |            |                    |
| D4S1614 | 27925  | Z24429            | 2646689-2646866            |            |                    |
| D4S43   | 147240 | D4S43             | 2336363-2336628            | 0.7        | C39 <sup>46</sup>  |
| D4S127  | 149984 | D4S127            | 3038714-3038864            | 0.7        | P363 <sup>47</sup> |
| HTT     |        |                   | 3076408-3245687            |            |                    |
| D4S3034 | 38369  | Z51717            | 3325536-3325722            | 0.6        |                    |
| D4S412  | 9920   | Z16836            | 3380781-3380974            |            |                    |
| D4S2957 | 73817  | Z53093            | 3833487-3833597            | 0.6        |                    |
| D4S431  | 14923  | Z17175            | 6415645-6415795            |            |                    |

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