# UK NATIONAL EXTERNAL QUALITY ASSESSMENT SERVICE FOR HISTOCOMPATIBILITY AND IMMUNOGENETICS' (UK NEQAS for H&I) EDUCATIONAL SCHEME - THE FIRST 10 YEARS



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# UK NATIONAL EXTERNAL QUALITY ASSESSMENT SERVICE FOR HISTOCOMPATIBILITY & IMMUNOGENETICS

# **Educational Scheme**

UK NEQAS for H&I established an 'Educational Scheme' in 2002. This was prompted by its participants who wanted to HLA type a small number of 'challenging' samples each year in addition to testing routine external quality assessment material.

The scheme, which is gratis to participants of its HLA Phenotyping and/or DNA HLA Typing schemes, is not assessed and caters for both serological and DNA-based typing.

Participants can compare their findings with those of about 20 laboratories reporting serologically-defined types and some 35 laboratories submitting types using DNA methods.

#### Scheme's material

Samples are derived from consenting donors on the Welsh Bone Marrow Donor Registry's panel of some 75,000 HLA typed, largely north-west European Caucasoid, blood donors. This panel has been the source of 33 new alleles and 60 confirmatory sequences over the last 20 years – many of which have subsequently featured in this scheme.

Over the last decade four whole blood samples (in three instances DNA extracts) were provided each year.

Samples have possessed:

□rare and recently described alleles, e.g. B\*08:02 and B\*40:92

□expression variants, e.g. A\*03:01:01:02N and B\*08:19N

□serologically 'difficult' specificities, e.g. those of B\*44:09 and B\*44:14

□unfamiliar allele/specificity combinations, e.g. A\*24:17, B\*15:02 and B\*40:01 with C\*01:02.

# Novel alleles identified

The scheme has enabled the identification of two novel alleles, namely:

HLA-A\*11:15 (Tissue Antigens 2006, 67, 153-6)

HLA-DQB1\*02:01:04 (Tissue Antigens 2011, 78, 296)

# **Testing comparisons**

Some notable alleles were provided twice in separate years. For example:

B\*27:23 - sent in 2003 and 2006

For B\*27:23 some 80% of laboratories failed to detect B27 using serology in both years' testing.

However, there was a significant increase in the successful detection of  $B^*27$  by DNA-based methods in the later send-out (77% to 94%, p<0.01) with a similar improvement in the assignment of  $B^*27:23$  (67% to 74%).

B\*15:42 - sent in 2005 and 2010

For B\*15:42 some 45% of laboratories identified B15 by serology in both send-outs indicating that B\*15:42 will be missed by many laboratories using serology alone.

Using DNA-based typing, 69% and 83% of laboratories identified B\*15:42 in the different years, respectively, with only a minority of laboratories failing to assign B\*15.

# **Further information**

Detailed information on the findings for all 40 samples can be found in UK NEQAS for H&I's Annual Reports at <a href="www.neqashandi.org.uk/reports.asp">www.neqashandi.org.uk/reports.asp</a>. Further information on UK NEQAS for H&I schemes - and its Prospectus - is available at <a href="www.neqashandi.org">www.neqashandi.org</a>.