Newborn and childhood screening programmes: criteria, evidence, and current policy

D A C Elliman, C Dezateux, H E Bedford

“The systematic application of a test or enquiry, to identify individuals at sufficient risk to benefit from further investigation or direct preventive action, amongst persons who have not sought medical attention on account of symptoms of that disorder” (Wald)

Screening is offered to apparently well individuals to identify those at high risk of a specific condition, for whom early treatment, more effective treatment, or information may be offered in order to improve health outcome or to provide opportunities for informed decision making. By definition, screening tests are not diagnostic tests and therefore cannot separate reliably those with a specific condition from those without. Hence, those with a negative screening result will include some affected individuals, usually referred to as “false negatives”, who may be falsely reassured by the screening result. Conversely, those with a positive screening result will include unaffected individuals, usually referred to as “false positives”, who may be worried unnecessarily or be exposed to the risks of subsequent diagnostic tests. Therefore the threshold for a positive test will be determined by balancing the goals of screening and the perceived disbenefits of missing affected individuals or falsely labelling healthy individuals.

In the past screening tests have often been introduced as a result of the enthusiasm of one or more individuals without evidence of effectiveness or careful consideration of the wider implications of false reassurance or false labelling. In the UK, this has led to the refinement of the historic “Wilson and Jungner” criteria, so that all proposed screening programmes are assessed against specified criteria.

THE PRESENT FRAMEWORK

To introduce a more systematic approach, the National Screening Committee (NSC) was established in 1996 with the remit that it should advise the Minister of State for Health on screening issues. Even though there are now separate health administrations, the NSC still has a role across the whole of the UK. One of the first tasks of the NSC was to define criteria for an acceptable screening programme. In recognition that the whole programme and not just the screening test has to be acceptable, there are now 19 criteria, although all may not have to be met for each programme. They cover the condition, the test, the treatment, and broader aspects of the programme. These criteria provide a framework for appraising evidence in relation to current or proposed screening programmes, which in turn form the basis of policy recommendations. For a current programme, these may include recommendations to continue screening, perhaps with modifications, or to discontinue the programme. For a proposed programme, these may include advice either to implement, or not to introduce. It is not within the remit of the committee to advise on issues to do with screening of particular high risk groups, for example, screening for hearing loss or hypothyroidism in children with Down’s syndrome.

In 1998, Antenatal and Child Health Subgroups (ANSG and CHSG) were set up, in recognition of the large number of screening tests that are offered during pregnancy and early childhood. The CHSG advises the UK NSC on three main aspects of screening programmes: first, on the implementation, development, review, modification, and where necessary, cessation of UK childhood screening programmes; second, on the need for research in relation to screening, and for analytical work to help focus and make best use of research; and third, on quality management issues. In its first two years, the CHSG considered many conditions, often by holding workshops with relevant professionals and parent groups. Where relevant, these were held jointly with the ANSG to reflect the fact that screening strategies for some conditions could be offered before conception, during pregnancy, or after birth, albeit with often different goals. There is not room to describe all the recommendations in detail, but the main points are set out below.

PHENYLKETONURIA, CONGENITAL HYPOTHYROIDISM, AND MEDIUM CHAIN ACYL COA DEHYDROGENASE DEFICIENCY

In the UK, screening for phenylketonuria was introduced in 1969 and, for congenital hypothyroidism, in 1981. These programmes have been successful in identifying infants with phenylketonuria or congenital hypothyroidism before irreversible neurological damage has occurred, thereby preventing lifelong disability. The advent of tandem mass spectrometry (MS/MS) enables many compounds, including amino acids, to be assayed on the dried blood spots. In the UK, MS/MS is being increasingly used to screen for phenylketonuria, replacing older methods, and has the capacity to detect many other compounds. These include acyl carnitines which may be used to identify other inborn errors of metabolism, such as organic acidemias and disorders of fatty acid oxidation, of which medium chain acyl CoA dehydrogenase deficiency (MCADD) is one of the most important. In appraising the criteria for MCADD screening, the CHSG has drawn on primary research carried out in the UK, systematic reviews commissioned by the UK NHS Research and Development programme, and reports from established or pilot programmes in North America, Australia, and Europe. Despite international experience of screening well over a million newborn infants, important questions and uncertainties remain about performance and outcome of newborn screening for MCADD. In particular, there has been no report of a systematic follow up of longer term outcome in affected infants detected by screening. In view of this, the NSC recommended that any new programme should be carried out within a UK setting to determine programme performance and early outcome.

NEONATAL SCREENING FOR SICKLE CELL DISEASE

Sickle cell disease predominantly, though not exclusively, affects particular ethnic minority populations in the UK. There is evidence from non-UK populations that prophylactic penicillin given following early diagnosis can reduce invasive infections with encapsulated organisms, in particular the pneumococcus, and reduce mortality. In the light of this evidence and the findings of two commissioned systematic reviews, the NSC has recommended newborn screening for sickle cell disorders and antenatal screening for haemoglobinopathies, including thalassaemia. This
was announced as a government priority in the NHS plan and details of the implementation are currently being considered by a working party.

CYSTIC FIBROSIS
A systematic review of screening for cystic fibrosis, commissioned by the NHS Research and Development programme, concluded that “the ability of [newborn] screening to alter prognosis has not been conclusively proven”, that evidence available was “either predominantly short-term or subject to strong statistical bias”, but that there was “some circumstantial evidence favouring a benefit”. At the same time, the authors concluded that antenatal screening should be offered to pregnant women using a DNA based test with the aim of identifying couples with a 1 in 4 risk of an affected child, who could then be given information regarding their risk and reproductive choices. The NSC reviewed the evidence from this and other international reviews, from information published subsequently, and from workshops and meetings with relevant professional and parent groups. The NSC concluded that there was currently insufficient evidence of longer term benefit, specifically in relation to pulmonary function, to support implementation of newborn screening. Subsequently, a ministerial decision was made in July 2001 to formally introduce screening in England, followed shortly by a similar decision for Scotland. An implementation plan for England is being formulated. Currently antenatal screening is only carried out by a single centre in Scotland, but this policy is currently under active review.

DUCHENNE MUSCULAR DYSTROPHY
There is no treatment for Duchenne muscular dystrophy. The main value of screening is therefore to warn parents of the possibility that future boys in the family may be affected. This allows for the opportunity for prenatal counselling. Further evidence will become available from the experience of screening programmes in Wales, but, at present, there is not enough evidence to support a recommendation that screening should be universal.

DEVELOPMENTAL DYSPLASIA AND CONGENITAL DISLOCATION OF THE HIP
The term developmental dysplasia of the hip (DDH) refers to a spectrum of developmental hip disorders which include partial or complete displacement of the femoral head from the acetabulum, particularly in children with congenital dislocation of the hip. This spectrum also includes acetabular dysplasia with or without displacement. Developmental displacement is associated with impaired growth and development of the hip and, usually, with subsequent abnormalities of gait and premature degenerative changes in the hip joint. A national screening programme based on clinical screening to identify neonatal hip instability was introduced in the UK in 1969. There is however scientific uncertainty about the effectiveness of this programme, reflecting the fact that it has never been assessed in a randomised trial. Since the mid 1980s there has been increasing use of ultrasound to image newborn infants’ hips, although in the UK this has been principally offered to those with risk factors or clinically detected hip instability. While neither clinical nor ultrasound screening programmes have been satisfactorily evaluated in relation to longer term outcomes, evidence suggests that universal ultrasound programmes identify similar proportions of affected children compared with screening by expert clinical examiners. However, experience from other European countries with universal ultrasound screening programmes suggests that uncertainties in the ultrasound criteria for treatment result in very high treatment rates. This raises concerns about over treatment of infants with false positive screening results. Sources of information for the policy review included research carried out for the MRC Working Party on Congenital Dislocation of the Hip, and systematic reviews from the USA and Canada. The NSC has recommended that clinical screening should continue, but with renewed emphasis on the standards and training for the clinical screening examination, with ultrasound used to image infants with clinical hip instability and recognised risk factors for DDH. Two national workshops have been held to develop an implementation plan. However, the NSC recognises that these policies are based on limited research evidence and plans to keep this policy under review.

HEARING IMPAIRMENT
The authors of a recent systematic review of neonatal hearing screening to identify children with permanent congenital hearing impairment concluded that the current infant distraction test (IDT) fails to fulfil the criteria for a good screening test. Even in ideal circumstances the IDT has poor sensitivity and specificity and cannot identify children before the age of 7–9 months. As early identification results in better outcome for children, and targeted neonatal screening only detects about 50% of cases, it was recommended that universal neonatal screening should be introduced. The CHSG held two workshops and established a universal neonatal hearing screening steering group to consider the evidence, consult with both health professionals and parents, and take forward an action plan. Twenty pilot sites have been set up around the country as the first step in implementation of universal neonatal hearing screening. Screening in most of the pilot sites will be hospital based, but the effectiveness of screening in other settings will also be addressed.

VISION DEFECTS
Currently babies are examined at birth and at 6–8 weeks for evidence of ocular abnormality, predominantly congenital cataract. At subsequent ages, an attempt is made to detect strabismus, reduced visual acuity, and abnormalities of colour vision. The evidence in favour of most of these activities is surprisingly sparse. The NSC has recommended that the programme of screening for retinopathy of prematurity should continue, as should newborn screening for media opacities and other eye anomalies, and a repeat examination between 4 and 6 weeks. However, training and supervision must be improved. The NSC also recommended that orthoptists should screen children between 4 and 5 years of age, with the aim of testing all children by the age of 5, but that all other routine vision testing in children less than 5 should cease, as should the test for 7 year olds. The major resource implication of these proposals is that orthoptists will be undertaking more tests. The feasibility of this needs to be examined in more detail.

It is tempting to recommend that all other tests of vision in school aged children should be abandoned because the evidence in their favour is poor and one might rely on self referral to the local optician. However, as there is some evidence that vision screening may reduce, if not eliminate, social gradients in age at presentation with amblyopia, it is possible that discontinuation of existing programmes may result in increased health inequalities. Therefore all other tests of vision in school aged children should continue unchanged—for further details see the Health For All Children website (http://health-for-all-children.co.uk).

GROWTH DISORDERS
There is a lack of formal evidence as to the value of routine growth monitoring; its use in detecting abnormal growth velocity has low sensitivity and specificity. In 1998 a multiprofessional group met to develop a consensus on this issue. At this meeting it was agreed that the potential benefits of growth monitoring include identification of chronic disorders, giving reassurance to parents, monitoring the health of the nation’s...
children, and supporting future re-
search. Weight monitoring in infancy and among older children does not ful-
fill the criteria for screening. Single height
measurements, with a cut off point at the 0.4th centile, come closest to satisfying the criteria for screening. It has been
suggested that it is good practice to
weigh infants and young children at
routine contacts, for example, at 6–8
weeks and when they attend for
immunisations. It is recommended that
height and weight should be measured at
school entry. At no other age does the
measurement of height satisfy the crite-
ria for a screening test and so cannot be
recommended as a routine in well
children. Recently, attention has been
drawn to the increasing prevalence of
obesity; however, there is insufficient
evidence to support the introduction of
routine screening for obesity at any age in
childhood.11

DISORDERS OF DEVELOPMENT
AND BEHAVIOUR
The terms “developmental screening” and “developmental assessment” are
often incorrectly used interchangeably.
As a result of this lack of clarity, tests which may be useful as assessment tools
have often drifted into use as screening
tests. Law et al concluded that there was
insufficient evidence to introduce screening tests for speech and language
delay.12 A trial of a screening instrument for autism in toddlers—the CHAT test—
has recently been completed, which
has reported a low sensitivity. This test was
performed at 18 months (norms are not available for other ages). Of the 38
children who matched the criteria for
high risk for autism and who were
re-screened a month later, 12 continued
to meet the criteria on re-testing. At 42
months, 10 of these 12 had diagnoses on
the autistic spectrum, one had language
delay, and one was normal. However, the
sensitivity was only 18%. A satisfactory
screening test for autism is therefore
currently unavailable. A systematic
review of randomised controlled trials of
occupational therapy for developmental
coordination disorders failed to provide
evidence that occupational therapy im-
proved the child’s ability to function in
everyday life.13 There is no evidence to
support population screening for de-
velopmental delay.

ASSURING QUALITY IN
SCREENING PROGRAMMES
Newborn and childhood screening pro-
grammes depend on high quality ser-
vices, including appropriate information
for families who are offered testing, to
ensure that informed choices are made, and
to ensure that care is delivered in an effective
and timely fashion. A national audit of
newborn screening for phenylketonuria
and congenital hypothyroidism,10 to-
gether with a commissioned review of
quality management in screening,2 have
resulted in the establishment of a
UK-wide National Newborn Screening Pro-
gramme Centre. This centre started work
in April 2002 and is overseeing all
screening programmes based on testing
the heel prick sample. The centre has a
remit to monitor, and facilitate improve-
ment in, the quality of newborn screen-
ing processes and their outcomes for
parents and their babies. A key feature
will be the integration of this quality
assurance programme for newborn
screening into mainstream NHS quality
management through a national frame-
work for standard setting, training, and
performance measurement.

CONCLUSION
Screening tests are only one component
of a properly coordinated screening pro-
gramme and should not be introduced
until there is evidence that the potential
benefits of the whole programme out-
weigh the harms in place. It is important
that there is continued moni-
toring of the programme to ensure the
maintenance of standards and that
unanticipated problems are not arising.
Unless there are good reasons to the
contrary, no new screening programmes
should be introduced on a local rather
than national basis, except as a time lim-
ited research project. The menu of
potential screening programmes is con-
stantly changing and new topics will be
considered by the NSC in the future. It is
only through this process of reviewing
evidence against explicit criteria that we
can ensure that the benefits of new pro-
grammes outweigh the harms.

Arch Dis Child 2002;87:6–9

Authors’ affiliations
D A C Elliman, Department of Child Health, St George’s Hospital, Tooting, London SW17
OQT, UK
C Dezateux, H E Bedford, Centre for Paediatric Epidemiology and Biostatistics, Institute of Child Health, 30 Guilford Street, London WC1 1EH, UK
David Elliman chairs the Child Health SubGroup of the National Screening Committee, Helen Bedford and Carol Dezateux are members of this subgroup

Correspondence to: Dr D A C Elliman, Department of Child Health, St George’s Hospital, Tooting, London SW17 OQT, UK, DavidElliman@compuserve.com
Accepted 5 April 2002

REFERENCES
4 National Screening Committee. First report of the National Screening Committee. London: Health Departments of the United Kingdom, 1998.
5 Simons WF, Fuggle PW, Grant DB, et al. Educational progress and motor skills at 10 years in early treated congenital
hypothyroidism. Arch Dis Child 1997;77:219–22
6 Beasley MG, Costello PM, Smith I. Outcome of treatment in young adults with phenylketonuria detected by newborn
7 Pollitt RJ. Leonard JV. Prospective surveillance study of medium chain acyl-CoA dehydrogenase deficiency in the UK. Arch Dis
Child 1999;79:116–19
Child 1999;80:459–62
1997;1
1997;1
12 Liebl B, Fingerhut R, Roschinger W, et al. Model project for updating neonatal screening in Bavaria: concept and initial results [in German]. Gesundheitswesen
2000;62:189–95
13 Andreassen BS, Dobrowolski SF, O’Reilly L, et al. Medium-chain acyl-CoA dehydrogenase (MCAD) mutations identified by
MS/MS-based prospective screening of newborns differ from those observed in patients with clinical symptoms: identification
14 Carpenter K, Wiley V, Sim KG, et al. Evaluation of newborn screening for medium chain acyl-CoA dehydrogenase deficiency in
15 Zykovicz TH, Fitzgerald EF, Marsden D, et al. Tandem mass spectrometric analysis for amino, organic, and fatty acid disorders in
16 Davies SC, Craini E, Gill M, et al. Screening for sickle cell disease and thalassaemia: a systematic review with supplementary
1999;3:1–186
collection of the test results and social and psychological significance of the testing. Pediatrics 2000;106:1450–5
21 Godward S, Dezateux C. Surgery for congenital dislocation of the hip in the UK as a measure of outcome of screening. Lancet
1998;351:1149–52
dislocation of the hip. Arch Dis Child 2000;82(suppl 1):A2
23 Dezateux C, Elbourne D, Arthur R, et al, on behalf of the MRC UK Cleft Lip Lip
EUROPE CALLING

Shortage of vaccines: USA, Europe, who next?

We are all aware of the merits of preventing infectious diseases by vaccines and many of us have witnessed impressive examples of successful immunisation strategies: for example, the global eradication of smallpox, the elimination of poliomyelitis in most countries of the world, and the substantial decline of invasive Hib disease in the western world.

However, safety questions raised about certain vaccines—whether true (intussusception associated with rotavirus vaccine) or false (autism due to MMR)—have challenged paediatricians. Nowadays, it frequently requires long discussions, pronounced powers of persuasion and patience to convince parents about the benefits of vaccines. As if this were not enough, a further crisis has arisen: vaccine supplies are running short and the range of available vaccines is shrinking in the USA and several European countries. According to the Centers for Disease Control and Prevention, 8 of the 11 routine childhood vaccines are currently in short supply in the US, namely diphtheria, tetanus, and pertussis (DTPa); measles, mumps, rubella (MMR); varicella; and the pentavalent pneumococcal conjugate. The large demand for polysaccharides to be included in the latter may even negatively influence the availability of the familiar 23-valent pneumococcal polysaccharide vaccine, indicated for individuals aged 2 years or older with chronic diseases. In some European countries, formerly available “orphan vaccines” such as DT (without pertussis component) or monovalent diphtheria toxoid with reduced antigen content (“adult formulation”) have disappeared.

How could this happen? Several recent mergers of pharmaceutical companies have led to market consolidation. Only a few “global players” are still manufacturing vaccines and their perceived duty to shareholders has resulted in some ceasing to produce less profitable vaccines. Similarly, former state owned or subsidised vaccine manufacturers are increasingly being forced to show a profit. Unfortunately, inspections of some plants by regulatory authorities have revealed failures to maintain quality standards—so-called “good manufacturing practice” guidelines—have led to interruptions of production, thus intensifying the supply shortfall. Further, increasing numbers of different antigen components in modern vaccines make it more and more likely that a given production lot might not fully qualify criteria for all ingredients.

What is the way out of this dilemma? Logically, the more manufacturers of standard vaccines there are, the less likely is significant shortage of supply. Furthermore, companies producing vaccines should be required to announce ahead of time any withdrawal of vaccines from the market thereby allowing other manufacturers to increase their output. States should consider long term contracts with vaccine manufacturers to assure sufficient supply for their population and, perhaps most importantly, local vaccine producers in less developed countries need to be strongly supported to maintain or increase their output with respect to quality and quantity. The children of the world (and their parents) deserve no less.

Ulrich Heininger
European Editor