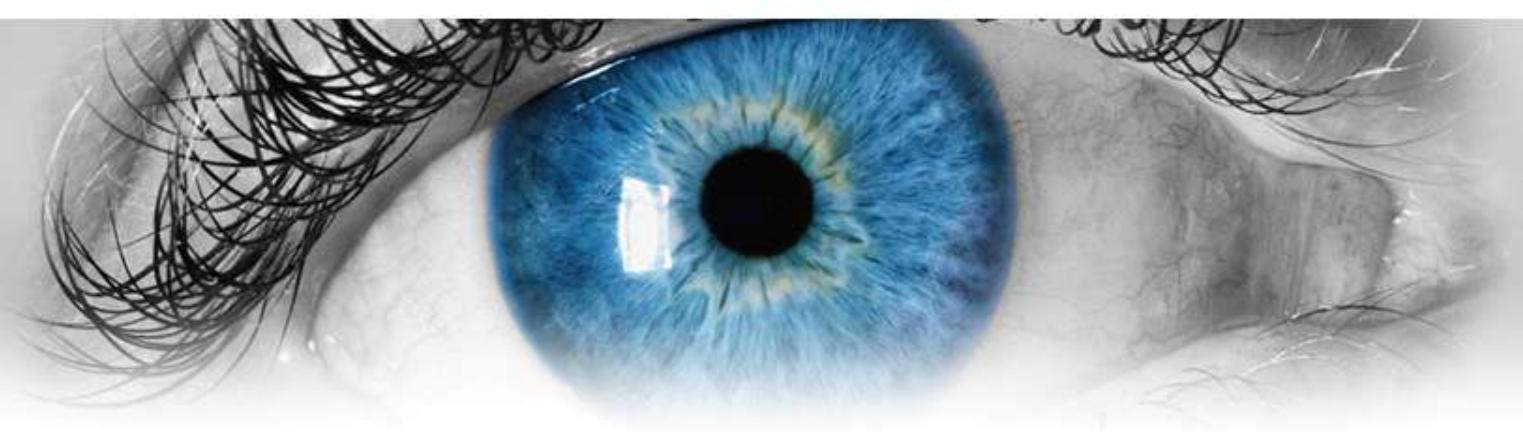


# The Great Debate – How often is enough?



BARS Annual Conference, 23 September  
2010

Dr Deborah M Broadbent



This house believes that  
patients should be screened  
annually for diabetic  
retinopathy



# Format

- Straw poll
- Present the pros and cons
- Expert panel discussion
- Open debate
- Repeat poll

A close-up, high-resolution photograph of a human eye with a vibrant blue iris. The pupil is dark and centered. The surrounding sclera is white, and long, dark eyelashes are visible at the top and sides of the eye. The image has a soft, slightly blurred background.

# Straw poll

- Annually
- 2 yearly for R0
- Individualised
- Undecided





# Recommended screen intervals

- The ENSPDR current recommendation is annual screening for all PWD aged  $\geq 12$  years\*
- Recommendations for alternative screening intervals have been made by national & international groups based on expert opinion / consensus rather than direct evidence.

*\* Workbook version 4.3 [www.retinalscreening.org.uk](http://www.retinalscreening.org.uk)*



- European Retinopathy Working Party recommends screening at least 2 yearly after diagnosis and at least yearly or more frequently if retinopathy develops [1]
- ADA recommends yearly or more frequently for type 2 DM [2]
- AAO recommends yearly screening for no DR / BDR and 6-12 monthly screening for mild PPF without maculopathy [3]

1. *Diabet Med* 1991;8:263–67

2. *Diabetes Care* 1998;21:157–59. 3

3. <http://www.aao.org/ppp>



# Evidence for longer intervals...

- Incidence data
- Cost-effectiveness
- Patient “costs”



# Cumulative incidence of STDR in Type 2 diabetes

7615 patients underwent 20,570 screen events

- Progression to STDR in year 1
  - BDR 5%
  - Mild PPF 15%
- 95% likelihood of remaining free of STDR:
  - No DR 5.4 years
  - BDR 1.0 years
  - Mild PPF 0.3 years





# Cumulative incidence (CI) of STDR in Type 1 diabetes

501 patients underwent 2742 screen events

- CI of STDR in patients without baseline DR:
  - 0.3% at 1 year
  - 3.9% at 5 years
- 95% likelihood of remaining free of STDR:
  - No DR                      5.7 years
  - BDR                        1.3 years
  - Mild PPF                0.4 years



# Conclusions

- Patients with both type 1 and type 2 diabetes and no DR at baseline could safely be screened at longer intervals (up to 3 years) unless:
  - duration > 20 years
  - insulin use in patients with type 2 diabetes
- Patients with BDR or the above risk factors need to be screened annually
- Patients with mild PPR need to be screened 4-6 monthly

A close-up, high-resolution photograph of a human eye with a vibrant blue iris and a black pupil. The eye is looking slightly to the right. The eyelashes are dark and thick, framing the eye. The skin around the eye is fair and has a soft, natural texture.

# Norfolk Data

- Patients managed solely in general practice
- 1990-2006
- 20,788 people screened at least once - 63,622 screen episodes
- Screen intervals of 18-24 months cf 12-18 months were not associated with a higher risk of STDR
- For a screen interval of >2 years there was a 60% increase in likelihood of STDR being detected
- Complements data from Liverpool

# Individualised screen intervals

The screenshot shows a web browser window titled "ES\_4\_DR\_Risk\_Basic\_FF - Windows Internet Explorer". The address bar shows the URL "http://liverpool.ac.uk/Diabetes\_Risk\_Factors\_ES/ES\_4\_DR\_Risk\_Basic\_FF.htm". The browser window contains a form titled "Expert System for Risk Analysis in Sight-threatening Diabetic Retinopathy using MatSOAP". The form has two main sections: "Primary Risk Factor" and "Risk Prediction".

Primary Risk Factor	Risk Prediction
HbA1c [%] <input checked="" type="checkbox"/> $\geq 8.0$	Risk [%] 83.7
Diastolic BP [mmHg] <input checked="" type="checkbox"/> 160	Confidence Interval [40.3 ... 98.9]
Systolic BP [mmHg] <input checked="" type="checkbox"/> $\geq 200$	AUROC 0.769
Cholesterol [mmol/l] <input checked="" type="checkbox"/> $\geq 8.0$	<input type="button" value="Evaluate"/> <input type="button" value="Reset"/>
Disease duration [y] <input checked="" type="checkbox"/> 5	

At the bottom of the form, there are buttons for "Help" and "Advanced". Below the form, the text "contact: Tony Fisher [a.c.fisher@liverpool.ac.uk](mailto:a.c.fisher@liverpool.ac.uk)" is displayed. The browser window also shows a status bar at the bottom with "Trusted sites" and "100%" zoom.

Example of **v poor control** at 5 years

Risk = 83.7%

[www.liverpoolseye.org](http://www.liverpoolseye.org)





# Wilson and Jungner screening principles

- The cost of the case-finding programme (including early diagnosis and treatment of patients diagnosed) should be economically balanced in relation to possible expenditure on medical care as a whole

WHO 1968



# Cost-effectiveness

Liverpool incidence data suggested that 70% patients with no DR and no high risk criteria could be screened less frequently than annually, resulting in sizeable cost savings\*

\*this data is based on imaging using 35mm transparencies and it may be that digital imaging is more sensitive at detection of BDR



# Cost per QALY

- Cost utility analysis allows cost comparisons across different diseases
- Quality adjusted life years (QALYs) are used as a measure of the utility value for a health condition multiplied by the remaining years of life expectancy
- Interventions for diseases with onset at earlier ages show greater impact on QALYs – longer expected period of benefit (e.g. type I diabetes)
- Procedures with a cost per QALY between \$20,000 and \$50,000 considered beneficial



# Vijan et al

- Modelling – evaluation of progression of DR and cost
- High risk type 2 patients (younger and HbA1C >11) would have a cost of \$40,530 per QALY
- Low risk patients (older patients with HbA1C <7) cost an additional \$211,570 per QALY
- Screening every 2 years would reduce cost to \$107,510 per QALY. Screening every 3<sup>rd</sup> year would reduce to \$49,760 per QALY
- Did not take into account cost of effects of blindness





# Patient "costs"

- Reduced screening intervals would be more convenient for patients in terms of:
  - Fewer appointments
  - Inconvenience of dilatation
  - Time off work
  - Travelling costs
  - Time



# Evidence for annual screening...

- Change in risk factors
- Non-attendance
- Feasibility
- Acceptability
  - to patient
  - to health professionals
- Cost



# Changing Risk factors

- Worsening control
  - Adolescence
  - Stress / depression – family/ personal illness, bereavement, change in circumstances
- Tightening control
  - Nb. Insulin pumps (pregnancy)
  - Retinal worsening
  - Reduce HbA1C by  $\geq 3\%$  in 1 year



# Non-attendance

- Chronic disease: multiple appointments
- Failure to attend may relate to lack of appreciation by people with diabetes of the risk of visual impairment
- Increased risk of progression of disease
- Failure of programmes to meet the ENSPDR key performance indicator on compliance with screening





# Feasibility

- Are the software programmes able to manage screen intervals greater than / less than 12 months?
- Are admin teams able to manage screen intervals greater than / less than 12 months?



# Acceptability

- To patients
  - I am reassured by annual screening
  - What happens if something does develop?
- To health professionals
  - Patient safety
- Research data is not available on relationship between patient / health professional perceptions and screen interval.  
Qualitative research is required



# Cost of missed disease

- Litigation costs are significant
- Cost of supporting a visually impaired patient
- Cost on secondary health effects of blindness is scant
- Blindness has also been associated with increased length of hospital stay, nursing home placement, and hip fracture



Expert panel and open  
discussion





# Straw poll

- Annually
- 2 yearly for R0
- Individualised
- Still undecided



Thank you for taking part  
in the great debate!