The guidance on age equality from the Royal College of Psychiatrists' Faculty of the Psychiatry of Old Age and General & Community Psychiatry Faculty1 is a breath of fresh air. It explores the implications of the Equality Act 2010. Under the Act, from April 2012 healthcare and other services in the UK must not discriminate by age and must be appropriate to the age of the service users. This is not the same as making services age inclusive. Equity of provision does not mean treating everyone the same, but rather recognising that different needs require different provision, distributed fairly. For example, putting older people with functional disorders on all-age wards where physical frailty, confusion or coexisting physical illness puts them at a disadvantage is unacceptable,2 as is inadequate training of staff working with them. Some people develop physiological and pathological changes usually associated with old age before the customary arbitrary dividing line of 65 years, others much later. Using a fixed chronological age cut-off for services in healthcare may be construed as ageist under the Act. A fixed age for accessing services also risks promoting ageism, by encouraging definitions of old age based on political and economic models rather than medically relevant ones. Flexibility to meet needs is essential, but there is then the risk that people falling into the ‘grey areas’ may be declared nobody’s responsibility. No style of service provision is without its drawbacks.

Tackling age discrimination

Some services for older people have changed their referral criteria to emphasise clinically based parameters, to ensure that service delivery is best tailored to the patient’s needs and least likely to be intrinsically ageist. However, the shape of an ideal service for older people is still open to debate. New service designs need further evaluation, but it is vital that they encourage and provide skills to treat functional and organic disorders in the presence of other illness and physical frailty. That breadth of approach has long been a valued characteristic of successful older people’s services. Discarding what works well may also be ageist if it leaves a sector of the community without dedicated, needs-appropriate services.

Clinicians are not going to openly admit to ageism, but ageism, possibly unintentional, is deeply embedded in healthcare. A survey by the British Geriatrics Society and Age UK concluded that the National Health Service (NHS) is institutionally ageist.3 Ageism includes underestimating older people’s ability to benefit from interventions, for both mental and physical illness. The latter was highlighted in a recent report about under-treating older people with cancer on the basis of chronological age rather than according to their overall health and fitness and likelihood of benefiting from treatment.4

The route to achieving non-discriminatory services is carefully outlined in the College statement.1 However, a number of pitfalls lie in our path and we must not be complacent.

Law might not be enough

Legislation alone does not guarantee enforcement. The NHS itself is enshrined in law by the National Health Service Act 1946. It was meant to be universal and provide for all age groups according to need. Laws to prevent discrimination on the grounds of disability, sexual orientation, gender, race and religion are still not fully implemented. Interpreting the law depends on the culture, expectations and understanding of society, not just legal matters. Society includes the
The ‘£2 billion gap’

The College’s guidance on the Equality Act points to an incredible £2 billion gap in the funding of health services between older and younger adults. This has accumulated over many years.

Negative attitudes towards older people have contributed to delegating them to second place in healthcare provision. This preceded the NHS, but was explicit in the planning of the welfare state when Sir William Beveridge stated in 1942:

‘It is dangerous to be in any way lavish to old age, until adequate provision has been assured for all other vital needs, such as the prevention of disease and the adequate nutrition of the young.’

Fears at a time of high infant mortality and declining birth rate may have influenced that stance. But it also seems relevant to ongoing reluctance to provide age-appropriate services.

In 1947, the British Medical Association (BMA) committee on the care and treatment of the elderly and infirm commented about delays in the provision of services:

‘Owing to the present restrictions on building construction and the shortages of trained staff, progress in achieving an improved medical service for the elderly will inevitably be slow. Many years must pass before any scheme ... can be brought fully into operation throughout the country.’

In 1956, a government-appointed committee reported on the cost of the NHS. It warned against under-resourcing older people’s services:

‘[T]he health authorities concerned should make sure that the needs of the aged are given their due priority in the allocation of additional resources and are not overlooked amid the pressure of other competing needs.’

Providing additional resources for old age was regarded as necessary, affordable and appropriate, but it was not forthcoming.

In the 1970s, health economist Nick Bosanquet reiterated the need to prioritise older people and prophesied:

‘For the psycho-geriatric service a special financial incentive is needed. Health authorities should be asked to submit plans and they should be financed by a special earmarked allocation of funds. Otherwise the DHSS [Department of Health and Social Security] will very probably be reissuing the same document in 1991 as it issued in 1959 and 1972.’

The mental health charity Mind urged the government in 1979 ‘to introduce legislation compelling social services departments to provide an early assessment and intervention service’. That never happened. It is, however, reminiscent, 30 years later, of the National Dementia Strategy which aimed to achieve ‘early diagnosis and intervention’. However, the Strategy unfortunately also stated that, ‘There is no expectation ... that all areas will necessarily be able to implement the strategy within 5 years’. Now, over 3 years into that 5-year period, despite improvements in some places, elsewhere little has been achieved. Clinically sound, well-intentioned objectives have been tempered by the laissez-faire acceptability of delays in implementation, as feared by Mind, and all too similar to the BMA statement in 1947.

Providing adequate mental health services for older people may need to be made obligatory; giving providers discretion has not worked in the past. The Commissioning for Quality and Innovation (CQUIN) payment framework giving financial rewards for implementing clinical innovations may have that effect; the national dementia CQUIN to identify dementia in older people admitted to general hospitals may be a step in the right direction. However, the alarm has been raised by some geriatricians doubtful whether service capacity is sufficient for following up those identified. Nevertheless, the CQUIN is a welcome first step.

In 2009, there was a 24-fold variation in activity between the highest and lowest performing primary care trusts for provision of dementia services. Age discrimination and the ‘postcode lottery’ both need to be addressed. The Faculty of the Psychiatry of Old Age and its predecessors have repeatedly tried to improve this state of affairs. Collaboration with other organisations such as Alzheimer’s Society has helped, but old age service advocates are fighting widespread prejudices of society and have not yet achieved their goals. In March 2012, we heard the Prime Minister declare that dementia care is a ‘national crisis’; once more deficits are acknowledged, will they be remedied?

Is this rhetoric, with action impeded by ageism?

Today, with the unfavourable economic climate, many might be tempted to reiterate the statement by the Ministry of Health in 1950 about proposals for better services for ‘mental infirmity’ in old age: ‘It is recognised that the present conditions of financial stringency limit opportunities for action’. Repeatedly putting older people to the end of the queue is discriminatory and has
not worked; the crisis remains. The Equality Act may provide opportunities to argue for and implement age-appropriate schemes on a par with those for younger people, and result in better health and cost savings.

We cannot afford complacency

We have a moral and ethical obligation to support people who are dependent because of age, illness or disability. The Equality Act should help us achieve that. It will be disastrous if the Act is allowed to join a catalogue of other ineffective legislation and powerless Department of Health proposals. The Act is well intentioned and idealistic with respect to older people. We need to grasp the opportunity it gives us and ensure it succeeds.

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No scope for complacency: time to improve healthcare for older people
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