Appendix C4. Smoking cessation services model

Smoking cessation services (SCS): Quantifying the impact of smoking cessation services on smoking cessation rates

Introduction

There are essentially two main approaches to help reduce the smoking prevalence among a population:

1) preventing initiation, and 2) increasing the cessation rate. For this report we focused on smoking cessation services, a means to increasing the cessation rate among a population.

Many smokers find it hard to quit smoking on will power alone (termed self-help) – many smokers make multiple attempts to quit but fail because self help appears to have at best a small effect on success (1). Various types of aided, cessation strategies exist ranging from counselling behavioural therapy to first-line and second-line medications (2), all of which are known to increase the long-term success of quit attempts (3, 4) (Table 1).

Effectiveness of smoking cessation services

The effectiveness of smoking cessation services depend on a number of factors, such as the type of medication used, the type of behavioural intervention used, the smoker's psychological state and environmental influences. The Cochrane Collaboration conducted a series of reviews which looked into the effectiveness of various smoking cessation interventions (Table 2). Of the various smoking cessation pharmacological interventions, varenicline appears to be the most effective in achieving long-term abstinence rates. Varenicline (trade name Champix) is a nicotinic receptor partial agonist. A randomised controlled trial found that the one year continuous abstinence rate was 10% for placebo, 15% for bupropion and 23% for bupropion (5). A Cochrane systematic review concluded that varenicline improved the likelihood of successfully quitting smoking by two- to three-fold relative to pharmacologically unassisted attempts. Varenicline was more efficacious than bupropion in this regard but not statistically superior to NRT (3).

Table 1. Effectiveness of smoking cessation interventions (results from the Cochrane Collaboration review series)

Smoking cessation intervention	RR for	95%	No. of participants
	cessation	confidence	
		interval	
Combined pharmacotherapy & hebayioural intervention	3.88	3 35 - 4 50	5 887 (1 study)

(1)			
Group behavioural intervention	1.98	1.60 - 2.46	4,375 (13 studies)
(6)			
Pharmacotherapy (nicotine replacement therapy*) intervention	1.60	1.53 - 1.68	50,000 (117 studies)
(7)			
Telephone counselling intervention	1.37	1.26 - 1.50	24,000 (9 studies)
(8)			
Self-help material intervention	1.32	1.20 - 1.42	28,189 (25 studies)
(9)			

RR: risk ratio for cessation; * includes any form of NRT: gums, patches, lozenges, inhaler and nasal spray.

The importance of creating a national network of smoking cessation services

In the UK, the government has already demonstrated a strong commitment to reducing smoking prevalence through the creation of a highly effective national network of smoking cessation services – known as NHS stop smoking services (10). This support is designed to be widely accessible within the local community and is provided by trained personnel. However, recently there have been decline in the number of smokers attempting to quit through the NHS stop smoking service, as well as a fall in the number of smokers successfully quitting (11). It is important to ensure that a national network of smoking cessation service continues to be easily accessible for smoker in the UK, and to ensure that such a policy intervention can be established in the other EConDA countries.

Table 2. Various smoking cessation treatments are available

Types of smoking cessation treatments

Nicotine replacement therapy (NRT)

Skin patches

Chewing gum

Inhalator

Tablets, strips, lozenges

Nasal spray

Mouth spray

Smoking cessation medication

Bupropion

Varenicline

Nortriptyline

Electronic cigarettes

Behaviour change techniques

Written material

Group counselling sessions

Individual counselling sessions

Telephone counselling sessions

Combination of the above

Project aim

The project aims to make a quantitative comparison of the health and economic impact of smoking cessation service over time, by comparing the policy scenario with the 'baseline' scenario (i.e. no change as compared to the current situation).

Methodology

EConDA model

The intervention chosen for this study was a 12-week smoking cessation service involving the administration of varenicline alongside face-to-face counselling. This was based on the Maudsley model which is an evidence based approach to treating dependent smokers using a combination of regular meetings (with a trained advisor using structured, withdrawal-oriented behavioural therapy) combined with smoking cessation medications such as nicotine replacement therapy (NRT), bupropion or varenicline (10). Many clinical practice guidelines recommend the use of both classes of interventions for smokers who are prepared to make a quit attempt (1). This is based on the assumption that the two types of treatment have complementary modes of action, and may independently improve the chances of maintaining long-term abstinence. However, surveys suggest that the proportion of people who use both types of treatment when attempting to quit smoking is low (12).

Varenicline was used for all of the EConDA countries except for the Netherlands whereby bupropion was used as the pharmacological intervention of choice (due to availability of data). Varenicline, a relatively new drug (approved by the FDA and EMA in 2006), was evaluated instead of bupropion as it is known to deliver higher smoking cessation rates, be more cost effective and is relatively safe and well tolerated (13, 14) – hallmarks of a pharmacological intervention that would make it ideal for rolling out nationwide. The EConDA model requires three types of input data:

- Effectiveness of the intervention
- Reach of the intervention
- Cost of the intervention

Effectiveness of the intervention

Effectiveness of the intervention in terms of cessation rates was expressed as 12-months continuous abstinence. Only cessation rates that were biochemically validated through the measurement of the smoker's carbon monoxide levels (as opposed to self-reported data) were included in the model. Given that these type of data were not available for most countries, proxy data from other countries were used (Table 4). It was deemed appropriate to use proxy data where necessary based on the assumption that the pure biological effect of a drug can be expected to be the same, irrespective of the country (15). Studies by West et al. found that the first 28 days since quitting is the most crucial period for likelihood of relapse. Thus it was deemed appropriate for this model that the rate of relapse was negligible following the use of 12-months continuous abstinence rates.

'Reach' of the intervention

Typically, various demand- and supply-side constraints contribute to the overall 'reach' of a public health intervention within a given population. This means that even if an intervention is rolled-out on a national scale, the intervention may only go on to be taken up by a fraction of the target population.

Whereas fiscal smoking policies can be imposed on an entire population (i.e. a population reach of 100%), non-mandatory interventions have a smaller reach since their demands are affected by, for example, the smokers' willingness to quit smoking and their desire to reach out for professional support. Data on 'willingness to quit smoking' was publicly available from four of the EConDA countries (Table 3) – these figures were then incorporated into the model to reflect the demand-side constraint of the 'reach' of the intervention. In some countries, smoking cessation drugs have to be paid for by the service user through the purchase of a prescription. Table 5 lays outs which of the smoking cessation drugs are free a country-by-country basis. For simplicity, however, in the model, the cost of the service at the point of delivery was assumed to not act as a barrier to the uptake of the SCS by the target population (i.e. service would be free for any smoker taking up the service), given that data on the relationship between the cost and demands of the SCS were not available. In the model, the SCS was free for all patients, in that the payer (National Health Service or national/federal health insurance) covered the total cost of the service (16), keeping in line with making the economic case for providing public health interventions that are free at the point of delivery.

It was assumed that only 50% of those wanting to quit smoking would actually participate in the intervention owing to supply-side constraints, such as the supply of healthcare professionals and current availability of intervention infrastructure (17). This figure was applied across all of the eight EConDA countries (Table 3), since country-specific data in this area was lacking.

Cost of the intervention

The intervention cost, expressed as total cost per quit attempt, was based on estimates of real resource use. Unless otherwise stated, the price typically covered the duration of a 12-week course of varenicline tablets as well as the administrative costs incurred by healthcare professionals leading the counselling sessions. Costs of adverse effects were assumed to be negligible. Costs varied considerably between the EConDA countries (Table 3) – in countries where cost data were not available, proxy data from another country were used in its place (Table 4).

Model assumptions

The following assumptions in the model were made:

• An individual eligible for the intervention is selected at random from the entire population distribution of smokers. To determine to determine whether or not the intervention takes place, an

application-generated random number assigned to an individual in the simulation is compared against the threshold probability (composed of the 'reach' and 'abstinence' rates)

- A smoker is defined as an individual who has smoked for at least a year
- All smokers in the model are eligible for the intervention (but in reality, for example, smokers who
 present with a known history of epileptic seizures, brain tumour, renal disease, hepatopathy, severe
 hypertension or suicidal ideation would be ineligible for a course of bupropion medication)
- The willingness to quit and the effectiveness of the intervention are the same across age, sex, severity
 of addiction and socioeconomic gradients
- A smoker's willingness to quit smoking stays the same throughout the entire simulation period i.e. no other changes in cultural or political trends would occur that might alter the smoker's willingness to quit smoking over time
- The 'reach' of the intervention stays the same throughout the entire simulation period i.e. no other changes in the supply or demand of the intervention is expected to occur within the time horizon
- The 'reach' of the intervention is the same across age, sex, socioeconomic gradients and geographical areas
- Once a smoker quits smoking as a result of the intervention, the smoker stays an ex-smoker throughout the rest of the time horizon (the relapse rate is captured within the 12-month continuous abstinence rate)
- For both the baseline and the policy intervention scenario, smokers can also quit smoking by means other than that of the intervention e.g. unassisted attempt to quit (smokers who quit smoking via these routes may still relapse and become a smoker again at some point in the future)
- The cost of the intervention is free at the point of delivery i.e. it is paid in full by the national health service, local government or national/federal health insurance
- A smoker cannot use the intervention more than once in any given year, but has the potential to use the SCS at the start of each new year within his/her lifetime regardless of the number of times he/she has had the intervention

Limitation of the model assumptions

Cessation of smoking is known to slow the progression of COPD in patients who had been smoking (18), and thus this intervention scenario could possibly reduce the number of patients with severe stage COPD and reduce direct healthcare costs associated with COPD – this effect was not factored into the model

Table 3. Summary input data for the smoking cessation service intervention

	Bulgaria	Finland	Greece	Lithuania	Netherlands	Poland	Portugal	UK
Reach								
Willingness to quit smoking								
(%)	59%	59%	65%	59%	40%	59%	59%	68%
Accessibility of the	3770	3770	0370	3770	1070	3770	3770	0070
intervention (%)	50%	50%	50%	50%	50%	50%	50%	50%
Overall reach (%)	30%	30%	33%	30%	20%	30%	30%	34%
Impact of the intervention								
Type of pharmacological drug	Varenicline	Varenicline	Varenicline	Varenicline	Bupropion	Varenicline	Varenicline	Varenicline
12-month abstinence rate		, , , , , , , , , , , , , , , , , , , ,		, , , , , , , , , , , , , , , , , , , ,				
(%)*	34%	34%	22%	34%	17%	34%	34%	34%
Long-term relapse rate (%) **	0%	0%	0%	0%	0%	0%	0%	0%
Outcome criteria ±	Continuous	Continuous	Continuous	Continuous	Continuous	Continuous	Continuous	Continuous
Validation method \P	Biochemical	Biochemical	Biochemical	Biochemical	Biochemical	Biochemical	Biochemical	Biochemical
Cost								
Cost (cost/quit-attempt)	429 лв	€ 248	€ 220	€ 621	€ 282	621 zł	€ 209	£ 164

Grey shading indicates the use of proxy data (more information available in appendix A1 to A4 and A6 to A8) * as a % of the service users; ** as a % of the service users (>1 and <5 years post cessation); * either point prevalence or continuous abstinence; ¶ either self-reported or validated by biochemical testing

Table 4. Data sources for the smoking cessation service intervention model

	Bulgaria	Finland	Greece	Lithuania	Netherlands	Poland	Portugal	UK
Reach								
Willingness to quit								
smoking (%)	FL proxy	(19)	(20)	FL proxy	(21, 22)	FL proxy	FL proxy	(23, 24)
Accessibility of the		()	(-)	r · J	(, ,		r - J	(-,)
intervention (%)	NL proxy	NL proxy	NL proxy	NL proxy	(17)	NL proxy	NL proxy	NL proxy
Overall reach (%)	N/A	N/A	N/A	N/A	N/A	N/A	N/A	N/A
Impact of the	,	,	,	,	,		,	,
intervention								
Type of								
pharmacological								
drug	N/A	N/A	N/A	N/A	N/A	N/A	N/A	N/A
12-month abstinence		,	,	2.,22	,			/
rate (%) *	UK proxy	UK proxy	(25)	UK proxy	(26)	UK proxy	UK proxy	(27)
Long-term relapse	p. o	on prony	(==)	512 p. 61.j	(==)	promy	promy	()
rate (%) **	N/A	N/A	N/A	N/A	N/A	N/A	N/A	N/A
Outcome criteria ±	N/A	N/A	N/A	N/A	N/A	N/A	N/A	N/A
Validation method ¶	N/A	N/A	N/A	N/A	N/A	N/A	N/A	N/A
Cost	11,11	11,711	11/11	11/11	**/ **	11/11	11,722	11,711
Cost (cost/quit-								
attempt)	NL proxy	Norway proxy	NL proxy	NL proxy	(26)	NL proxy	NL proxy	(27)

Grey shading indicates the use of proxy data; * as a % of the service users; ** as a % of the service users (>1 and <5 years post cessation); * either point prevalence or continuous abstinence; * either self-reported or validated by biochemical testing

Table 5. Accessibility of pharmacological drugs (16)

Country	Country		Finland	Greece	Lithuania	Netherlands	Poland	Portugal	UK
Is there a toll-free telephone quit line/help line with a live person available to discuss cessation with callers in your country? Is this product legally sold in the country?		Yes Yes	Yes Yes	No Yes	Yes Yes	Yes Yes	Yes Yes	Yes Yes	Yes Yes
Nicotine replacement therapy	Where and how can this product be legally purchased in your country?	In a pharmacy without a prescription	In a general store without a prescription	In a pharmacy without a prescription	In a general store without a prescription				
	Does the national/federal health insurance or the national health service cover the cost of this product? Is any NRT on the	No	No	Partially	No	No	No	No	Fully
	country's essential drugs list?	No	No	No	No	Yes	No	No	
	Is this product legally sold in your country?	No	Yes	Yes	Yes	Yes	Yes	Yes	Yes
Bupropion	Where and how can this product be legally purchased in your country?		In a pharmacy with a prescription	In a pharmacy with a prescription	In a pharmacy with a prescription	In a pharmacy with a prescription	In a pharmacy with a prescription	In a pharmacy with a prescription	In a pharmacy with a prescription
	Does the national/federal health insurance or the national health service cover the cost of this product?		No	Partially	No	Fully	No	No	Fully
Varenicline	Is this product legally sold in your country?	Yes	Yes	Yes	Yes	Yes	Yes	Yes	Yes

	Where and how can this product be legally purchased in your country?	In a pharmacy with a prescription							
	Does the national/federal health insurance or the national health service cover the cost of this product?	No	No	Partially	No	No	No	No	Fully
Is smoking	Health clinics or other primary care facilities	Yes in some	Yes in most	Yes in some	Yes in some	Yes in most	Yes in some	Yes in some	Yes in most
cessation support available	Hospitals	Yes in some	Yes in most	Yes in some	Yes in some	Yes in most	Yes in some	Yes in some	Yes in most
in the following places	Office of a health professional	Yes in some	Yes in some	Yes in some	Yes in some	Yes in most	Yes in some	Yes in some	Yes in most
in your country?	In the community	Yes in some	Yes in some	No			Yes in some	Yes in some	Yes in most
	Other	Yes in some							
Does the national/federal	Health clinics or other primary care facilities	Partially	Partially	Partially	Fully	Fully	Partially	Fully	Fully
health insurance or the national health service cover the cost of	Hospitals	Partially	Partially	Partially	Fully	Fully	Partially	Fully	Fully
	Office of a health professional	Partially	Partially	Partially	Partially	Fully	Partially	Fully	Fully
	In the community	Partially					No		Fully
this support?	Other	Partially	Partially	Partially	No	Fully	Fully	Partially	Fully

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