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**Coverage with evidence development, only
in research, risk sharing or patient access
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CHE Research Paper 77

Coverage with evidence development, only in research, risk sharing or patient access scheme? A framework for coverage decisions

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Abstract

Context

Until recently, purchasers' options regarding whether to pay for the use of technologies have been binary in nature: a treatment is covered or not covered. However, policies have emerged which expand the options - for example, linking coverage to evidence development, an option increasingly used for new treatments with limited/uncertain evidence. There has been little effort to reconcile the features of technologies with the available options in a way that reflects purchasers' ranges of authority.

Methods

We developed a framework within which different options can be evaluated. We distinguished two sources of value in terms of health: the value of the technology *per se*; and the value of reducing decision uncertainty. The costs of reversing decisions are also considered.

Findings

Purchasers should weigh the expected benefits of coverage against the possibility the decision may need to be reversed and the possibility adoption will hinder/prevent evidence generation. Based on the purchaser's range of authority and the features of the technology different decisions may be appropriate. The framework clarifies the assessments needed to establish the appropriateness of different decisions. A taxonomy of coverage decisions consistent with the framework is suggested.

Conclusions

A range of coverage options permit paying for use of promising medical technologies despite their limited/uncertain evidence bases. It is important that the option chosen be based upon not only the expected value of a technology but also the value of further research, the anticipated effect of coverage on further research, and the costs associated with reversing the decision.

Key words: health technology assessment, cost effectiveness, comparative effectiveness, only in research, coverage with evidence development, patient access scheme

Introduction

Historically, new health care technologies such as pharmaceuticals and devices gained access to countries' health care systems and coverage by public or private third-party payment systems directly following regulatory approval or licensing. In the case of pharmaceuticals, this required evidence of the safety and efficacy of their products (European Medicines Agency 2009; U.S. Food and Drug Administration 2009). However, many health systems, in an effort to stem rising health care costs, now require a higher standard: that the expected additional health benefits of a new technology justify its additional cost (Freemantle and Hill 2004). This so-called "fourth hurdle" (Taylor et al. 2004) has changed the type and quantity of evidence required by purchasers before technologies are made available for widespread use and manufacturers are able to receive payment for use of their products. Even in systems which do not explicitly consider cost and cost-effectiveness, there is often a focus on the magnitude of health benefits, considering the effectiveness and safety of new products, and informally weighing this against cost - for example, evidence suggests that Medicare in the USA does not automatically cover new, more costly medical technologies without consideration of the *magnitude* of their incremental health benefits (Chambers, Neumann, and Buxton 2010; Neumann and Tunis 2010).

Thus, in most systems purchasers (sometimes called 'reimbursement authorities') are increasingly balancing the potential for improved health outcomes offered by new technologies against acquisition costs. This challenge is accentuated by the fact that such decisions are having to be made earlier, often shortly following licensing, partly as a result of pressure from patients, lobby groups, politicians and manufacturers (Boseley 2006; Cooksey 2006; Evans and Boseley 2006; House of Commons Health Committee 2005; National Institute for Health and Clinical Excellence 2008). However, the type of evidence available at a product's launch is largely focused on what is necessary to inform regulatory authorities (e.g. for pharmaceutical licensing) rather than coverage decisions, and there is typically little information on relative effectiveness in routine use or against existing interventions, let alone about relative cost-effectiveness. New technologies often appear promising, offering improvement over existing interventions in the same disease area. However, the evidence of long term benefits and costs is inevitably not available at launch, and there is often considerable uncertainty about the consequences of their widespread use.

A health care technology may be considered valuable if the additional benefits it generates are expected to exceed the additional costs associated with its use. In a budget-constrained health care system like the UK's NHS, a technology can be regarded as valuable if the expected health benefits exceed the health expected to be forgone as other NHS care is displaced due to any additional costs of the new intervention (Claxton et al. 2008). In open health care systems (those without an explicitly fixed budget constraint), such as in the US, the expenditure on a more expensive new technology may displace either health, other non health uses of expenditure as a result of higher taxes or insurance premiums or a mixture of both. As such, a new technology would be regarded as valuable only if its health improvement outweighs the benefits of those things it has displaced.

In making the decision to cover a new technology, there is also value in reducing the uncertainty surrounding a decision through evidence generation: reducing the chance of error (e.g., deciding to pay for use of a technology which is ultimately not valuable) and making better decisions (Claxton, Cohen, and Neumann 2005). Also, making a wrong decision and having to reverse it as more evidence emerges often incurs a cost (Eckermann and Willan 2008; Palmer and Smith 2000). For example, a technology may require some investment in capital or staff training which cannot be recouped or, once a technology has been widely used, there may be a cost of withdrawing it (e.g. costs associated with changing common clinical practice, political pressure to maintain its coverage).

Purchasers need to be able to take all these factors into account when making decisions about the coverage of new health care technologies.

Until recent years, coverage decisions in many health systems have been considered largely binary in nature, based on the price set by the manufacturer and the evidence available at launch: that is, the purchaser would choose whether or not to pay for the product, for the entire indication or for a particular subgroup, based on existing evidence and current prices. However, new coverage options have emerged which have expanded the options available. Examples include coverage only *with* research (Carino, Sheingold, and Tunis 2004; Schuessmann et al. 2009; Tunis and Whicher 2009), only *in* research (Chalkidou 2006) and with performance linked payments, sometimes referred to as 'risk sharing' (Rutten, Uyl-de Groot, and Vulto 2009; Towse and Garrison 2010). These methods have been developed with the objective of allowing patients early access to promising and innovative health care technologies. They also aim to give manufacturers longer to make returns before patent expiration, and also to allow health care providers to give their patients access to promising technologies whilst reducing the risk payers face in making a wrong coverage decision and obtaining more evidence.

In the US, Medicare has begun to develop coverage policies aimed at reducing uncertainty about effectiveness through the generation of additional evidence. The term Coverage with Evidence Development (CED) is used by Medicare as a catch all term for two separate coverage options in which generating additional evidence is a condition of coverage: coverage with study participation (CSP) and coverage with appropriateness determination (CAD). CSP allows coverage of technologies for which the evidence is not adequate to support full coverage but where additional evidence gathered in the context of clinical care would clarify the benefits of treatment, therefore restricting coverage to those patients receiving the intervention as part of a clinical trial or registry. CAD allows for the collection of additional clinical information that would not be available on a claims form to determine the appropriateness of coverage, therefore not restricting the coverage to those patients participating in a clinical trial or registry but still providing additional evidence. These two schemes have also been referred to as 'only in research' (OIR), which restricts coverage to patients receiving the intervention as part of a clinical study or registry, and 'only with research' (OWR), which does not necessarily limit coverage to those patients participating in a study or registry, respectively (Carlson et al. 2010). The distinction between OIR and OWR is primarily the degree of coverage that the payer confers during the period of evidence generation.

There has been little effort to date to reconcile the plethora of terms used to describe these schemes or to distil their essential features. This paper provides a conceptual framework which allows these new coverage options to be understood more clearly, identifying the essential contribution each makes and the key assessments that purchasers should make to establish when each is most appropriately used. We show that appropriateness of coverage decisions depends on the combination of several circumstances, including the features of the particular technology and the range of authority granted to the purchaser. Finally, we present a taxonomy of the coverage options available and a classification of schemes observed in practice and we discuss implications for policy and the impact of widening the range of authority granted to purchasers.

Conceptual framework

Identifying the assessments that should be made when considering the coverage of a new health care technology is central to understanding the potential contributions that new coverage options offer. The appropriate coverage of a promising technology depends upon the assessment of three essential features: i) the value of the technology based on existing evidence; ii) the value of reducing uncertainty about the technology's benefits, risks, and costs by acquiring more evidence; and iii) the value of any investment or reversal costs resulting from an initial positive coverage decision. These are considered in turn below.

The value of a technology

A technology can be considered valuable if its expected additional benefits (in health or cost terms) justify its additional net costs. In a budget-constrained, or 'closed', health care system like the UK's NHS, funding a more expensive new technology requires other health care activities to be curtailed to accommodate the additional costs, leading to forgone health outcomes elsewhere which are incurred as 'opportunity costs'. In such a system, a technology is valuable if the expected health benefits to those who will use that technology exceed the health expected to be forgone by other NHS patients whose care would be displaced. The same principle applies to open health systems such as those in the USA but, in the absence of firm budget constraints on health, the opportunity costs may manifest themselves in terms of forgone non health expenditure through increased insurance premiums, taxation or copayments. Although the following exposition takes the perspective of a closed system, it readily generalises to open systems with the opportunities costs falling on non health expenditure rather than health.

Methods to estimate the long term additional health gains and additional net costs associated with a new technology are well established and increasingly sophisticated (Drummond et al. 2005; National Institute for Health and Clinical Excellence (NICE) 2008). Whilst there is no universal agreement on the most appropriate measure of health, most would agree that it is important to capture both the effect on the quantity and quality of life. One such measure which is commonly used is the quality-adjusted life-year (QALY) (Brazier et al. 2007). We use this measure as the unit of health measurement for illustration.

The relationship between value, health benefits and additional cost is illustrated in Figure 1. The y-axis shows the additional cost of the new technology when it is priced at three different levels (price less than P^* , P^* , and greater than P^*), and the x-axis shows the expected health benefits, measured in QALYs. If the health care budget is fixed then additional costs will displace other health care and health will be forgone for other patients (although as stated previously, this can be generalised to open systems where it is non health expenditure rather than health which is displaced). Therefore, some assessment of how much additional cost leads to one QALY being forgone elsewhere is required. This is the concept of the cost-effectiveness threshold used by some budget constrained health care systems (Culyer et al. 2007). The rising diagonal in Figure 1 illustrates a threshold where every additional £20,000 is expected to displace one QALY. This analysis can easily be extended to consider an open system with the threshold instead representing the willingness of society to give up non health expenditure for a health improvement.

Suppose the new technology is expected to offer two additional QALYs for each patient treated. If the price is less than P^* the additional health care costs of £20,000 would only displace one QALY elsewhere so there would be a net health benefit (NHB) to the health care system of one QALY. Therefore, in these circumstances the new technology can be regarded as cost-effective (the incremental cost effectiveness ratio is £10,000 per QALY which is less than the threshold). If the price is higher, at P^* , the additional health care costs of £40,000 would be expected to displace two

QALYs elsewhere, just equal to the expected health benefits. Therefore, there is no NHB overall and P^* represents the maximum the health care system can afford to pay for this technology. If the price is set above P^* , the health care costs of £60,000 would be expected to displace three QALYs. The health expected to be forgone exceeds the health expected to be gained. If this technology were approved it would reduce overall population health outcomes since the NHB is -1 QALY based on existing evidence.

What should become clear from this illustration is the critical role that the price of the technology plays in determining the value of a technology and whether it should be paid for based on existing evidence.

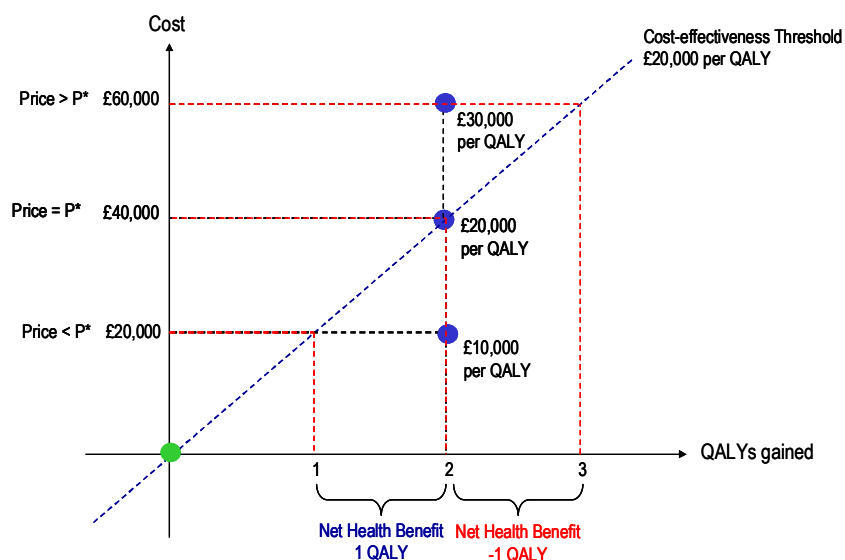


Figure 1: The value of a technology (source Claxton et al 2008)

Whilst the above analysis focuses on the *explicit* use of cost-effectiveness analysis to assess value, it is recognised that not all payers use formal economic evaluation. However, no health system can completely ignore opportunity costs. For example, in the USA, some parts of the private sector are beginning to use formal cost-effectiveness analysis (Wellpoint National Pharmacy and Therapeutics Committee 2008), but Medicare does not undertake the explicit forms of economic analysis to inform decisions seen in many European countries, Canada and Australia. However, there is evidence that at least some Medicare coverage decisions reflect an *implicit* assessment of value for money because Medicare does not automatically cover every intervention offering an incremental benefit – expensive technologies yielding only small gains in effectiveness may not be approved for coverage, cost is likely to be one consideration in this (Chambers, Neumann, and Buxton 2010; Neumann and Tunis 2010). In other words, some assessment is made of whether the magnitude of the health benefits generated by a technology is sufficient to justify it being covered. Indeed, in the context of recent developments in comparative effectiveness research (Institute of Medicine 2009), one interpretation of the value of this initiative to policy and decision makers is that it can provide an assessment of the magnitude of the health improvement offered by new technologies against existing interventions which can be considered alongside their costs. Whether the assessment of value is made explicitly or implicitly, the conceptual framework remains relevant in considering the coverage of new technologies. Although the framework we offer focuses on formal assessments of NHB as the measure of value, our approach can be generalised to more implicit value assessment consistent with the emerging tenets of comparative effectiveness research, whereby some assessment is made of whether the magnitude of the health benefits generated by a technology justify its coverage. The remainder of the paper refers to the concept of the value of the technology in general but it should be emphasised that this can be based on an explicit evaluation of cost-

effectiveness (such as that made by NICE) or implicit assessment informed by comparative effectiveness research methods.

The value of evidence

The previous section has focused on the value of a technology assuming that the costs and benefits are known with certainty. When assessing the value of a new technology, however, gaps in information may mean that the estimates of the expected (i.e. average) costs, health outcomes and, therefore, overall NHB will be uncertain. Additional evidence can reduce this uncertainty and provide more precise estimates which can reduce the risk of incorrect decisions (i.e., coverage of a technology that does not actually provide a net health benefit).

This value of evidence is illustrated in Table 1, where a new technology B is compared with an existing technology A. There is uncertainty about the clinical effect (α) and, thus, about the expected NHB of both. For illustration, suppose that the current evidence indicates that α could take one of four possible values with equal probability, each producing a NHB for each technology.

Based on current evidence, technology B has the highest expected NHB (7.5 QALYs), even though the probability it meets a given cost-effectiveness threshold is only 0.25 (B only has the highest NHB for one of the four possible values of α). Nevertheless, assuming there are no costs associated with reversing the decision to cover the technology, B is expected to be valuable, offering an additional 0.5 QALYs compared to technology A given existing evidence (Claxton 1999).

However, if more evidence about the effects of the two technologies could resolve the uncertainty surrounding α , then better coverage decisions could be made, improving overall NHB. This is the basis for estimating the value of further research. Using the example in Table 1, if the uncertainty could be fully resolved then we can establish which treatment would be adopted given the true value of α : B would be chosen if α_1 was shown to be the true value, but A would be chosen if α_2 , α_3 , or α_4 where shown to be true, improving NHB by 1 QALY in each case compared to B. However, the results of further research are as yet unknown (i.e. we do not know *ex ante* which particular value α will take), so we need to allow for the probability that each value of α turn out to be true - here we assume each result is equally likely. Therefore, the expected NHB of treatment, if additional evidence could resolve this uncertainty, is found by averaging across the maximum NHB associated with each particular value of α (giving 8.25 QALYs).

Table 1: Net Health Benefits under uncertainty

	Net Health Benefit (in QALYs)			
Possible values of α	Technology A	Technology B	Maximum NHB	Technology chosen
α_1	10	15	15	B
α_2	8	7	8	A
α_3	6	5	6	A
α_4	4	3	4	A
Expected Net Health Benefit	7	7.5	8.25	

This simple example illustrates that evidence may be valuable for the same reasons as access to a new and effective technology: it can improve overall health benefits. In this case, evidence that could resolve uncertainty offers additional expected NHB of 0.75 QALYs compared to choosing B

based on existing evidence (8.25 - 7.5 QALYs), i.e., greater than the expected value of new technology itself given current evidence (0.5 QALYs). However, 0.75 QALYs represents an upper bound on the value of additional evidence, or the expected value of perfect information (EVPI), since research which could be conducted is unlikely to resolve all uncertainty surrounding the choice between A and B (Claxton 1999; Claxton and Sculpher 2006). Some assessment of uncertainty, its consequences and the need for further research is needed by purchasers. Such assessment of uncertainty can be informed through well established quantitative methods which can be extended to consider the value of proposed research designs (Ades, Claxton, and Sculpher 2006; Conti and Claxton 2009).

However, the generation of additional evidence may be costly, both in resources needed to do the research and in delays in approving a valuable technology for widespread use. Ideally, a promising technology that is expected on average to be of value would be paid for while further research is being conducted to resolve uncertainties. However, allowing patients early access to the technology may affect the prospects of the research, which is required to generate the additional required evidence, actually being conducted. For example, manufacturers would have less incentive to invest in additional research about a technology once it is covered; physicians might consider further clinical trials to be unnecessary and unethical; and patients might be unwilling to participate if they already have access to the new technology. Therefore, early coverage may mean that the benefits of additional evidence to inform treatment choice for future patients will be forgone. If the benefits forgone for future patients exceed the benefits to current patients of earlier access, then it may be better to withhold coverage of a technology, even if it is expected on average to be valuable given existing evidence, until further research is conducted (Griffin et al. 2010).

For example, in Table 1, the value of access to the new technology (B) for current patients is 0.5 QALYs (compared to standard practice, A), but the value of additional evidence for future patients is 0.75 QALYs. The purchaser should consider whether the benefits of early coverage exceed the future benefits which may be forgone. In doing so they should consider the value of evidence associated with research that might be undertaken if coverage is withheld, how long such research might take, and what other changes might occur during that interval (e.g., the entry of cheaper generics or of other new technologies). Such assessment will partly depend on the price of the technology. For example, if the price of the technology is reduced then there will be greater benefits of early access for current patients and, if the technology was already regarded as valuable at the original price, the value of additional evidence for future patients will tend to fall. Both of these effects would make immediate coverage more likely. Critically, there is an important relationship between the price of a technology, its value and the need for additional evidence (Claxton et al. 2008; Griffin et al. 2010).

Costs of investment and reversal

This discussion indicates that, if further evidence about the benefits of a technology would be forgone as a consequence of early coverage approval, then decisions should not be based only on expected cost, benefits and NHB. An assessment of uncertainty and its consequences is also required. The value of evidence that might be forgone can be thought of as an investment (opportunity) cost of approval. However, there are other more familiar investment costs such as capital expenditure on equipment and facilities or staff training and learning by doing. There may also be some costs of withdrawing coverage once a technology becomes widely used (e.g., the time and effort required to change common clinical practice or political costs of withdrawing treatment). In such cases uncertainty that might lead to subsequent withdrawal of coverage of a technology means that the decision should not be based on expected NHB alone. The source of uncertainty may be in the estimates of NHB for which further research could partly resolve. However, there are other sources of uncertainty that cannot be reduced by further research but that might,

nonetheless, resolve over time. For example, the price of the technology or its comparators might be expected to change over time (e.g., the entry of generics at patent expiry), or new future technologies might become available and make current technologies obsolete. Even if further research is not conducted, the existence of investment and reversal costs could mean that delaying coverage until uncertainties are resolved or reduced might be preferable even for technologies that are expected to be of value on average based on current evidence (Eckermann and Willan 2008; Palmer and Smith 2000).

The relationship between investment or reversal cost, uncertainty and immediate coverage or delay is illustrated in Table 2. The new technology D is compared to current clinical practice C. NHBs in each period are partly determined by an uncertain event (e.g., the price of future generics) λ , which is uncertain but has two equally likely values.

Table 2- Option values

Possible values of	Net Health Benefit per year (in QALYs)		
	Technology C	Technology D	Maximum NHB
λ_1 (p=0.5)	0	1	1
λ_2 (p=0.5)	0	-0.5	0
Expected Net Health Benefit (10 years)	0	2.5	

If there are no investment or reversal costs, then the purchaser can choose the alternative with the highest expected NHB. If the technologies offer the reported NHB in each of the next 10 years then D should be approved with its expected NHB of 2.5 $[(0.5 \cdot 10) + (0.5 \cdot -5)]$, compared with current practice which, for simplicity, we assume is 0 QALYs. If some investment is required to implement D which would displace 2 QALYs elsewhere, the expected NHB of D would be reduced to 0.5 QALYs $(2.5 - 2)$ but it should still be covered if the uncertainty surrounding λ cannot be resolved.

However, if, after one year, the uncertainty surrounding λ were to be resolved (e.g., the price of new generics became known), withholding coverage until the value λ is known might be worthwhile. In the first year C would be retained with expected NHB of 0 QALYs. If $\lambda = \lambda_1$ then D would be covered for the following 9 years with a NHB of $(1 \cdot 9) - 2 = 7$ QALYs (the health benefit less the investment cost). If $\lambda = \lambda_2$ then C would be maintained with NHB of 0 QALYs. Therefore, the expected NHB of withholding approval of D for one year would be $(0.5 \cdot 7) + (0.5 \cdot 0) = 3.5$ QALYs; which is greater than an immediate decision to use D.

Although the expected benefits of D would be delayed for one year, these opportunity costs of delay would be more than offset by the benefits of the purchaser retaining the option to abandon the coverage of the technology and therefore avoiding the investment costs if they choose not to cover the technology. In other words, there is an *option value* of delay which, in this example, is 3 QALYs (the difference in NHB between delaying for a year, 3.5 QALYs, and immediate coverage, 0.5 QALYs). This illustrates that investment or reversal costs, when combined with other sources of uncertainty, may appropriately lead to the delayed coverage of a technology that is expected to be valuable on the basis of current evidence, even when further research is not worthwhile. As previously stated, the option value of delay will also depend partly on the price of the technology since a lower price will make delay more costly. Also, future changes effect the value of evidence generated by future research (Phillips, Claxton, and Palmer 2008). Therefore, in addition to estimates of expected NHB, purchasers need to have some assessment of the scale of investment or reversal costs associated with the coverage of a technology, the sources of uncertainty and when these might be resolved.

What are the implications?

In our analysis, the value of early approval and coverage is not only determined by the expected benefits and costs of a technology but also the value of additional evidence, the effects of approval on the prospects of further research, and any investment or reversal costs associated with a range of sources of uncertainty. The purchaser should explicitly, or implicitly, weigh the expected benefits of early adoption against the possibility that the decision may be 'wrong' and needs to be reversed in the future, and the risk that early adoption will hinder or prevent the generation of valuable evidence. These concepts provide a set of guiding principles about the type of assessments that need to be made by purchasing bodies, whether or not they are informed by formal analysis. Of course, all these issues are affected in different ways by the price of the technology and should be considered jointly as they all impact in different ways on the health delivered by the resources available to the health care system. The essential features of the various schemes that have been proposed, and the choice between them, rest on these tradeoffs.

The technology’s characteristics and the purchaser’s range of authority

Before assessing the coverage options available to the purchaser, it is first worth considering the possible characteristics of the technology (the ‘technology space’) and the purchaser’s range of authority (the scope under their control) in order to determine what coverage options can and should be considered.

Figure 2 represents a stylized view of the technology space, encompassing six important scenarios (labelled A to F). The scenarios are based on whether an initial positive coverage decision about the technology is likely to be too costly to reverse if new evidence emerges suggesting it was incorrect, whether future research would be worthwhile and whether additional evidence could feasibly be generated following a decision to cover the technology. Future research is potentially worthwhile if the value of the uncertainty that could be reduced exceeds the costs of the research (this is a necessary condition in our framework for research being worthwhile), i.e., that the research is likely to be beneficial when taking account of both the benefits in terms of reduced uncertainty and the costs. Whether the evidence could be generated following coverage would depend upon the source of uncertainty (e.g., this may not be possible if the uncertainty relates to the relative effectiveness of a treatment for which a RCT may be the only suitable design to minimise potential bias but a RCT may be unable to recruit patients if the technology is already covered).

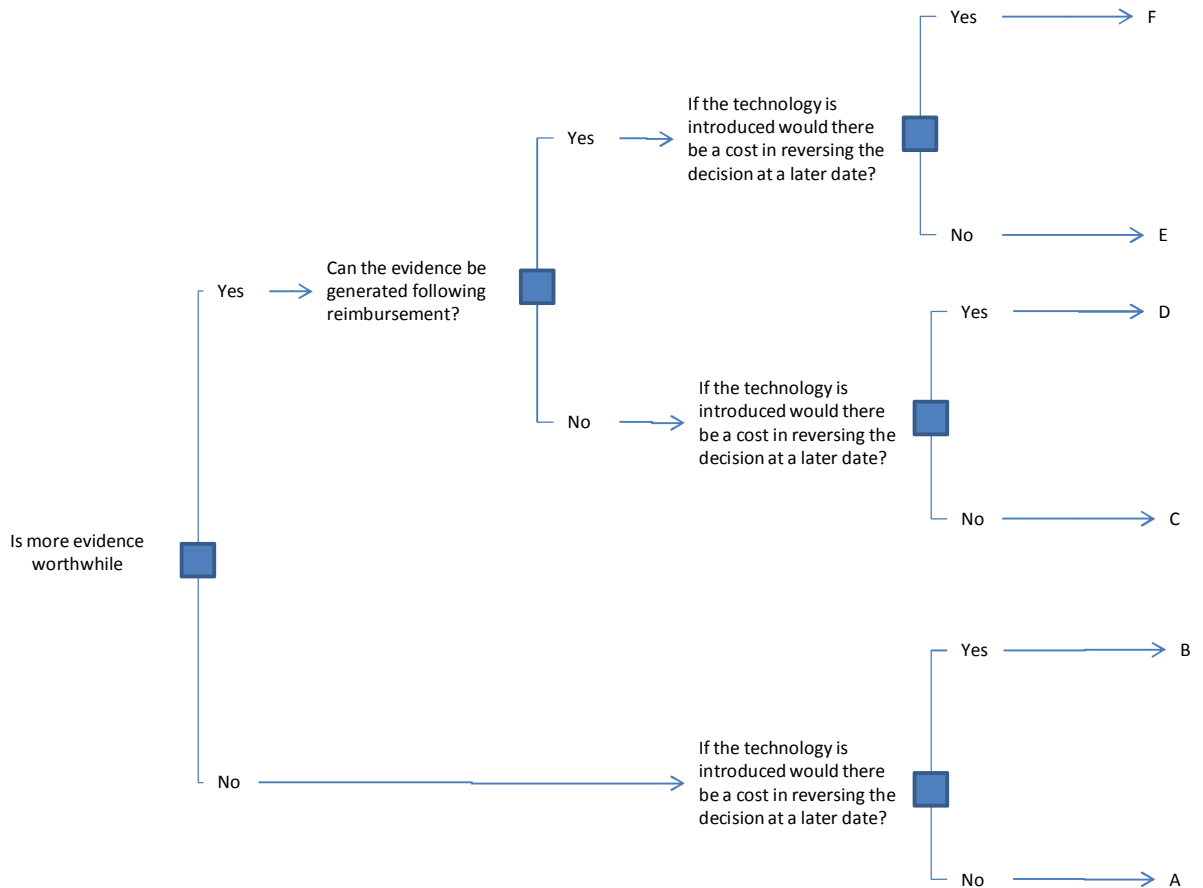


Figure 2: A pathway diagram showing the alternative characteristics of technologies (the ‘technology space’)

Figure 3 represents the possible ranges of authority that purchasers may have. Representing all possible nuances would be difficult. To simplify we present six key situations (labelled 1 to 6) based on a binary view of three dimensions of the purchaser's authority. These are, firstly, whether a purchaser can delay a decision or reverse it in the future in the face of new information. For example, some purchasers may be forced to make a decision on whether to cover the technology immediately, and that decision may be non reversible, whilst others may have the ability to delay the decision until more evidence is available or, if they initially decided to cover the technology, may be able to reverse this decision in the light of new evidence. Secondly, whether the purchaser has some influence over the effective acquisition price paid for a technology, either through price negotiation or some arrangements that reduce the effective price without changing the formal list price. The ability to influence price could be dependent on many factors, although here we assume for simplicity that the purchaser can only influence price if it can reject the technology grounds of lack of value, thereby giving the purchaser some bargaining power. Thirdly, whether the purchaser can contract for research or have some control of research budgets so as to ensure the research is conducted. It should be clear that the wider the range of authority of the purchaser the more extensive the options open to it when making coverage decisions on health care technologies.

The figure characterises six different types of purchases in terms of their range of authority. It is possible to locate some decision makers into these categories. For example, NICE in the UK probably most closely resembles Purchaser 3 in that it has the ability to delay and review decisions, has limited scope to ensure that research is conducted and has little influence on price. Other systems, such as HAS in France, have more influence over price.

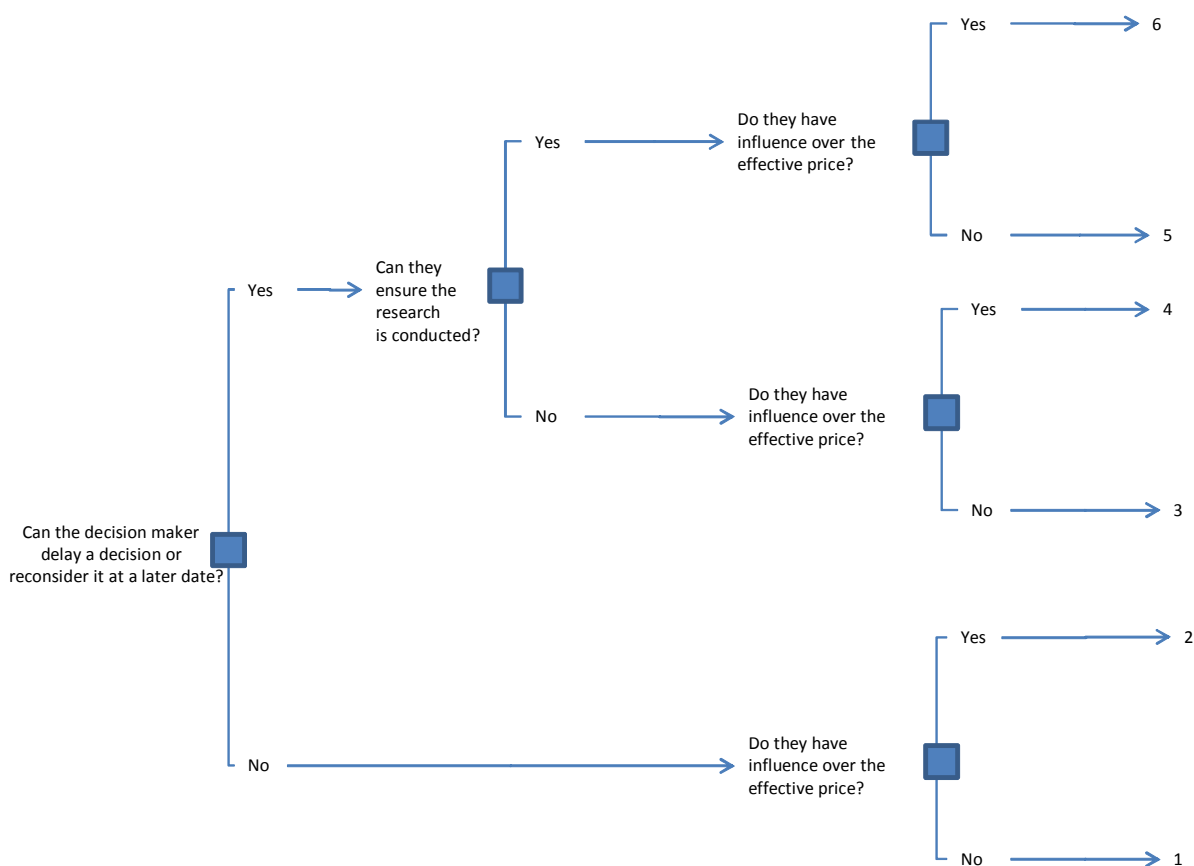


Figure 3: The purchaser's range of authority

Choosing among coverage options

The previous section has characterised the technology space and the purchaser's range of authority each in terms of six possible states (A to F and 1 to 6, respectively). Combining these creates 36 possible combinations, out of which falls a set of feasible coverage options which we consider in detail in Tables 3 and 4. These are as follows:

- *Reject*- The technology is not paid for on the grounds that it is not considered beneficial (i.e. of value) to the health system. This may be able to be reversed at a later date.
- *Accept*- The technology is paid for on the basis that it is considered beneficial to the health system. This may be able to be reversed at a later date.
- *Influence over effective price*- The purchaser has some influence over the effective price paid for a technology. It is assumed that influence will only be possible if the purchaser can reject the coverage of the treatment, otherwise it would have no bargaining power to ensure a reduction in the effective price.
- *Only in research (OIR)*- Coverage of a technology is available only to patients involved in research. This option may involve the purchaser paying for the research, which would require the purchaser to have some influence over research decisions (i.e., being able to contract for the research to be conducted). Alternatively, it may involve the purchaser rejecting the technology and simply recommending research, with the research being paid for by another party (e.g. the manufacturer or another stakeholder), which would not require the purchaser to be able to ensure the research was conducted.
- *Only with research (OWR)*- A positive coverage decision is conditioned upon the collection of additional evidence to support continued, expanded or withdrawal of coverage (Carlson et al. 2010). So the technology is paid for all but further research is also required. This research may be funded by the purchaser, the manufacturer or another stakeholder, but such a decision would require that the purchaser was able to enforce that the research is actually conducted, so will be treated here as an available option only when the purchaser can ensure the research is conducted.

Table 3 shows the coverage options from which the purchaser should choose in each scenario when the technology is expected to be of value (i.e. assessed explicitly or implicitly) on average based on existing evidence. The options available when the technology is not expected to be of value based on current evidence are shown in Table 4. How these choices should be made is then discussed with reference to the conceptual framework presented earlier. The analysis in these tables is based on the introduction of a new technology. When considering the future coverage of an already funded technology similar judgements must be made, although they are not specifically considered here. Our analysis also only considers uncertainties which will be resolved through research and not those uncertainties which will be resolved through some other method (e.g., waiting to observe the generic price of a technology).

Whatever the purchaser's range of authority, when there is no cost of reversal and collection of evidence is not worthwhile then the decision should be based solely on whether the treatment is expected to be of value or not. Therefore in column A of Table 3 the treatment is universally accepted, whilst in column A of Table 4 the treatment is rejected unless the purchaser has influence over the price paid, in which case, if the price can be lowered to a level where the treatment is expected to be of value, it should be paid for.

Table 3: Coverage options when the treatment is expected to be (i.e. on average) effective (or cost-effective) given existing evidence

E(NHB)>0		A	B	C	D	E	F
		No cost of reversal Evidence not worthwhile	Cost of reversal Evidence not worthwhile	No cost of reversal Evidence worthwhile Cannot get evidence with approval	Cost of reversal Evidence worthwhile Cannot get evidence with approval	No cost of reversal Evidence worthwhile Can get evidence with approval	Cost of reversal Evidence worthwhile Can get evidence with approval
1	Can't delay/reconsider No influence over effective price Can't ensure research is conducted	Accept	Accept	Accept	Accept	Accept	Accept
2	Can't delay/reconsider Influence over effective price Can't ensure research is conducted	Accept	Accept	Accept	Accept	Accept	Accept
3	Can delay/reconsider No influence over effective price Can't contract for research	Accept	Accept	Accept OIR	Accept OIR	Accept OIR	Accept OIR
4	Can delay/reconsider Influence over effective price Can ensure research is conducted	Accept	Accept Price influence	Accept Price influence OIR	Accept Price influence OIR	Accept Price influence OIR	Accept Price influence OIR
5	Can delay/reconsider No influence over effective price Can ensure research is conducted	Accept	Accept	Accept OIR	Accept OIR	OIR	OIR OWR
6	Can delay/reconsider Influence over effective price Can ensure research is conducted	Accept	Accept Price influence	Accept Price influence OIR	Accept Price influence OIR	Price influence OIR OWR	Price influence OIR OWR

Table 4: Coverage options when the treatment is not expected to be (i.e. on average) effective (or cost-effective) given existing evidence

E(NHB)<0		A	B	C	D	E	F
		No cost of reversal Evidence not worthwhile	Cost of reversal Evidence not worthwhile	No cost of reversal Evidence worthwhile Cannot get evidence with approval	Cost of reversal Evidence worthwhile Cannot get evidence with approval	No cost of reversal Evidence worthwhile Can get evidence with approval	Cost of reversal Evidence worthwhile Can get evidence with approval
1	Can't delay/reconsider No influence over effective price Can ensure research is conducted	Reject	Reject	Reject	Reject	Reject	Reject
2	Can't delay/reconsider Influence over effective price Can ensure research is conducted	Reject Price influence	Reject Price influence	Reject Price influence	Reject Price influence	Reject Price influence	Reject Price influence
3	Can delay/reconsider No influence over effective price Can ensure research is conducted	Reject	Reject	Reject OIR	Reject OIR	Reject OIR	Reject OIR
4	Can delay/reconsider Influence over effective price Can ensure research is conducted	Reject Price influence	Reject Price influence	Reject Price influence OIR	Reject Price influence OIR	Reject Price influence OIR	Reject Price influence OIR
5	Can delay/reconsider No influence over effective price Can ensure research is conducted	Reject	Reject	Reject OIR	Reject OIR	Reject OIR	Reject OIR
6	Can delay/reconsider Influence over effective price Can ensure research is conducted	Reject Price influence	Reject Price influence	Reject Price influence OIR	Reject Price influence OIR	Reject Price influence OIR OWR	Reject Price influence OIR OWR

Below we consider in detail how each type of purchaser should choose between the coverage options available in the other scenarios.

Purchaser 1 who cannot delay or reconsider a decision and has no influence over price

In this context, the decision regarding whether or not to pay for the use of the technology should be based solely on whether or not the technology is expected to be of value. Future research is not of value as the decision cannot be delayed or reconsidered at a later date when evidence becomes available and, similarly, any costs of reversal do not need to be taken into account as the decision cannot be reversed subsequently. Therefore, when the treatment is expected to be of value the technology should always be covered (see row 1 Table 3), and when the technology is not expected to be of value it should always be rejected (see row 1 Table 4).

Purchaser 2 who cannot delay or reconsider a decision and has influence over the price

Purchaser 2 is in a similar position to Purchaser 1 although it has influence over the price. Similarly to Purchaser 1, the value of evidence and any reversal costs do not need to be considered as it cannot delay or reconsider their decision. When the technology is of value on average, given existing evidence, it should always be accepted (see row 2 Table 3). Even though the purchaser has influence over the price, it has no bargaining power in the case when the treatment is expected to be of value as the manufacturer knows that the treatment will be paid for at the current price, and therefore has no incentive to reduce it. When the treatment is not expected to be of value it should not be paid for unless a reduction in the effective price can be secured to make the technology valuable on average (see row 2 Table 4).

Purchaser 3 who can delay or reconsider a decision and has no influence over price or scope to ensure the research is conducted

This purchaser has three coverage options available, either to accept or reject immediately, or to reject for now but recommend research be carried out to reduce any uncertainty (OIR recommendation).

In a situation where the technology is expected to have positive NHB based on current evidence (Table 3), if additional evidence collection is not worthwhile the treatment should always be accepted, as the evidence will never be generated to show that the decision was incorrect (3A and 3B Table 3). When gathering additional evidence is worthwhile then the purchaser should decide whether the value of immediate acceptance (the expected NHB as a result of patients receiving the treatment during the period it would take to conduct the research) outweighs the value of the evidence that will be forgone. If not then an OIR decision should potentially be recommended instead of accepting the technology immediately. As the purchaser is unable to ensure the research is conducted they cannot enforce that evidence will be generated following acceptance, so it does not matter whether the evidence can be generated with acceptance or not. Costs of reversal can be ignored as the evidence which would change the decision will not be generated if the technology is accepted. Therefore, the choice remains between accept and OIR (see 3C, 3D, 3E and 3F).

When the technology is not expected to be of value given existing evidence (Table 4), the treatment should be rejected unless the research is expected to be worthwhile, in which case an OIR decision may be appropriate.

Purchaser 4 who can delay or reconsider a decision and has influence over price but cannot ensure the research is conducted

Purchaser 4 is in a similar situation to Purchaser 3 but also has influence over the price paid. Therefore, the choice between coverage options is very similar in the two situations, although now a manufacturer may be able to alter an OIR decision to an accept decision by lowering the price and thus improving the technology's value. In this case the value of immediate acceptance would increase, and it may also reduce the value of additional evidence (making OIR less favourable) (see 4C, 4D, 4E, 4F Table 3 and Table 4). Even when the evidence is not worthwhile (i.e., the value of the reduction in uncertainty does not exceed the cost), when the treatment is not expected to be of value on the basis of current evidence the manufacturer can lower the price to a level so the treatment is expected to be of value to secure the coverage of their treatment (see 4A and 4B Table 4).

Purchaser 5 who can delay or reconsider a decision, can ensure the research is conducted but has no influence over price

Here the purchaser has more coverage options available as OWR and OIR options are included where the purchaser actively ensures the research is conducted rather than recommending research (and hoping it takes place) through an OIR decision, as has been assumed with the previous purchasers.

Let us first consider where the treatment is expected to be of value given current evidence (Table 3). When the collection of additional evidence is not worthwhile or research cannot be undertaken once the technology has been paid for (for example, when relative effectiveness evidence is required but coverage makes recruitment difficult), then the purchaser faces the same coverage options as Purchaser 3. Hence they should weigh up the same factors when choosing between the available options although they can now ensure that the research is conducted through an OIR decision (5A, 5B, 5C, 5D Table 3). However, when it is considered that the evidence can be generated once the treatment has been paid for, then a OWR option may now be worthwhile. When there is no cost of reversal, OWR becomes the dominant coverage option as more patients will receive the treatment that is expected to be of value therefore providing the same expected benefit to patients as immediate acceptance and the valuable additional evidence will also be collected (5E Table 3).

When there is a cost of reversal the choice is now between OWR and OIR (OWR still dominates outright acceptance as the uncertainty will also be resolved). The purchaser should now weigh up the expected NHB as a result of patients receiving the treatment in the time it would take to conduct the research against the probability that the treatment will not be found to be of value by the research and the cost of reversing the decision is incurred in the future. If the expected benefits outweigh the probability and costs of reversal then a OWR decision should be made; however, if they do not, an OIR decision should be considered (5F Table 3).

When the treatment is not of value, given current evidence, OWR should never be considered and the purchaser faces the same choices as those of Purchaser 3, although they should weigh up whether they should fund the research themselves, or simply recommend that the research is conducted.

Purchaser 6 who can delay or reconsider a decision, has influence over price and can ensure the research is conducted

Purchaser 6 faces the same coverage options as Purchaser 5 but also has influence over the effective price of the product. The purchaser should base their choice of coverage option on the same

discussion as for 5, but now should note that, by agreeing effective reductions in price, they can change the balance between the various coverage options. For example, whilst previously the probability and costs of reversal may have made an OIR decision favourable in 5F Table 3, by securing a reduction in price the NHB to patients increases, and may make a OWR decision worthwhile (6F Table 3).

When the treatment is not expected to be of value given existing evidence (Table 4), OWR now becomes a coverage option when the evidence can be generated with coverage, as long as the price can be reduced to a level where the treatment is expected to be of value (6E and 6F). In the case of 6E, if the treatment is now expected to be of value, OWR will dominate OIR (the patients all receive the treatment with expected NHB and the evidence will also be generated). In the case of 6F, the purchaser should now weigh up the expected NHB as a result of patients receiving the treatment during the research period, against the probability that the treatment will be found by the research not to be of value, hence incurring the cost of reversing the decision in the future. If the expected benefits outweigh the probability and costs of reversal then an OWR decision should be made; however, if they do not, an OIR decision should be considered.

What assessments are needed when choosing between coverage options?

The above has set out how purchasers should choose between the set of coverage options under various scenarios. There are a number of general insights which come from these analyses. When the purchaser is unable to delay a decision or to reconsider it at a later date their options are heavily limited: they should accept or reject, and the coverage decision should be based solely on the expected value of the treatment (whether assessed explicitly or implicitly). For technologies which are expected not to be of value given existing evidence, the purchaser is generally limited to rejecting the technology or trying to ensure a reduction in the price.

More interesting are the choices available when a technology is expected to be of value on average, given existing evidence. As the purchaser's range of authority expands so also do the coverage options available to them, and the choice should be based on more than the expected value for money. When evidence is worthwhile, but cannot be generated following a decision to pay for use, the value of the evidence should be weighed against the expected benefits to patients of immediate coverage. When there are large reversal costs the purchaser should weigh the expected benefits to patients against the probability that the decision will be wrong and should be reversed and the reversal costs incurred. What should be apparent is that the choice between coverage options always revolves around the trade-offs between the expected value for money of the treatment, the amount of uncertainty (and how much of it can be resolved through research) and the costs of reversing a decision.

A new taxonomy of coverage options

Conventionally, coverage decisions have been considered to be binary in nature: technologies were either paid for or not. However, the range of options available to purchasers can be expanded to include many more options which should be considered. A plethora of coverage options has developed although, as yet, there is no common terminology for those observed in practice. To address this, a taxonomy of coverage options is presented which attempts to draw out the key features of observed coverage decisions on the basis of the analyses presented previously. To this end the coverage options are split into two broad areas: those associated with reductions in the effective price (i.e. influence over pricing decisions from the previous section); and those associated with evidence generation (i.e. OIR or OWR decisions from the previous section). Differences between the coverage options are often a result of variation in how the allocation of risk between the manufacturer and the purchaser is altered by the agreement. Carlson et al. previously suggested a taxonomy which, although it has influenced the approach here, differs in that their key distinction was between whether the coverage options are outcome or non outcome based rather than whether the focus is on altering the expected price or reducing the uncertainty surrounding value for money through the generation of additional evidence (Carlson et al. 2010).

The taxonomy here is intended to assist an understanding of the coverage decisions observed in practice. The aim is to make a clear distinction between those coverage decisions which alter the expected value of a treatment with existing evidence through changing its effective price, and those which provide additional evidence so as to reduce the uncertainty surrounding value. Further differences are down to the allocation of risk between the purchaser and the manufacturer and to what is actually specified in the agreement, e.g. is the price explicitly linked to the results of research? In Appendix A various examples of these coverage decisions have been described and allocated to the appropriate section of the taxonomy.

Reductions in the expected effective price

Decisions which result in reductions in the effective price make treatments more desirable by reducing their expected costs to the health care system. The previous section grouped such agreements as having influence over the price paid for a technology; however, there are many nuances to the coverage decisions observed in practice which, whilst all affecting the effective price, do so in different ways. Figure 4 distinguishes reductions in effective price as being one of two types: outcome and non-outcome based.

Outcome based coverage decisions

Outcome based coverage decisions link the effective price paid for a technology to some measure of clinical outcome and, therefore, operate at the level of the individual patient (although these can be aggregated so payments are effected at the population level). Three types of outcome based agreements can be distinguished: money back guarantees, conditional treatment continuation and price linked to outcome.

i) Money back guarantees

Money back guarantee schemes involve the health service being refunded if a patient does not achieve a specified target. The level of refund is not necessarily 100% of the acquisition cost, and the refund may be financial or through another method, for example in replacement stock of the treatment. These schemes can be considered risk shifts, where the risk of a patient not achieving the outcome, and therefore having a negative NHB if the list price was paid, is shifted away from the health care system to the manufacturer who faces the risk of having to pay a refund. A notable

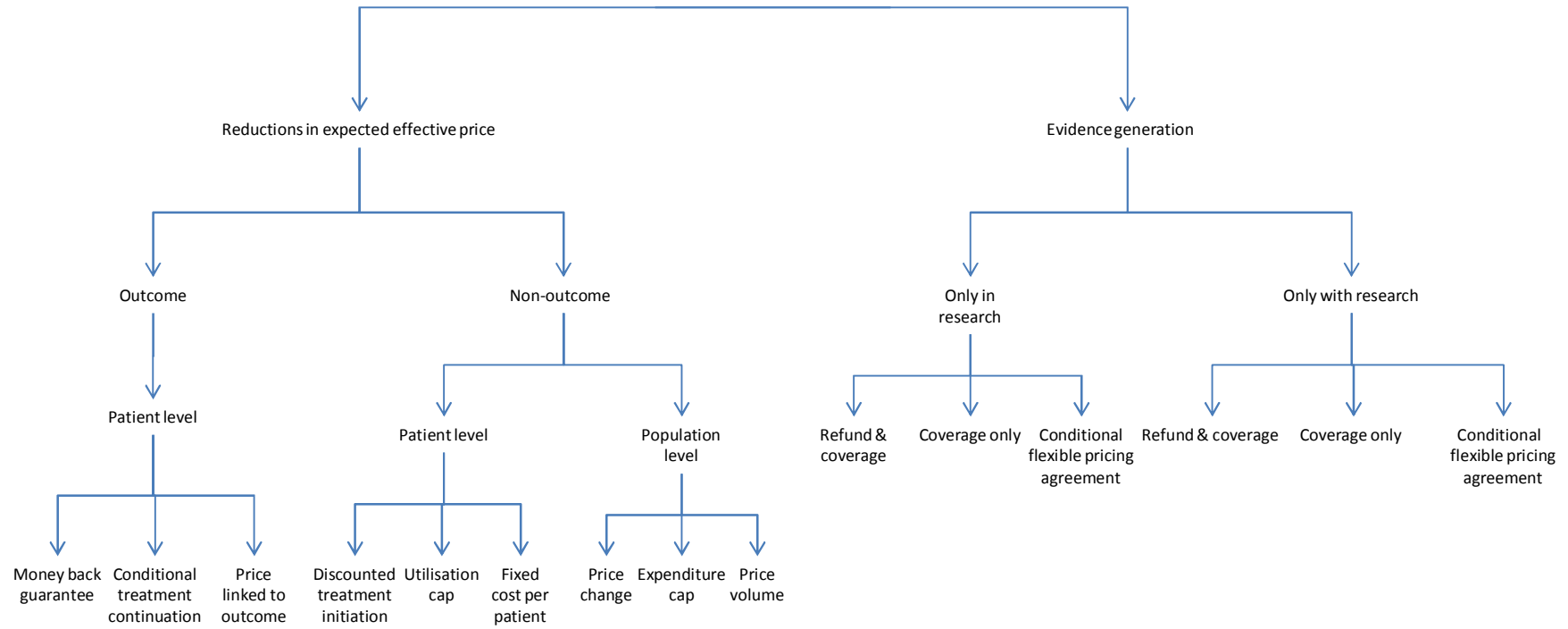


Figure 4: A new taxonomy of coverage options

example of such a scheme is the agreement between Park Davis and North Staffordshire Health Authority for the use of atorvastatin in patients with high cholesterol. The manufacturer agreed to rebate the health authority if patients did not achieve a low density lipoprotein cholesterol concentration target of less than 3 mmol/l after treatment with atorvastatin (Chapman et al. 2003).

ii) Conditional treatment continuation

Conditional treatment continuation involves payment for the continued use of a technology only in patients who have achieved a target clinical effect (Carlson et al. 2010). Such schemes can also involve refunds, either full or partial, for those patients who do not achieve the targeted effect. An example of such an agreement is that between Johnson and Johnson and the UK's NHS for the use of bortezomib for multiple myeloma. Johnson and Johnson agreed to repay the NHS in either cash or product for patients who fail to respond after 4 cycles of treatment with bortezomib. Those who respond would receive an additional 4 cycles of the treatment (Green et al. 2006).

iii) Price linked to outcome schemes

Price linked to outcome schemes involve the price being directly linked to a specified outcome for each patient (either clinical or other; for example, a biomarker). These schemes share similarities with money back guarantees, although the risk is still shared as the better a patient's outcome the higher the price. No examples of such schemes have been identified in practice, perhaps as a result of their high management costs or the possibility for gaming by either the manufacturer or the purchaser to artificially raise or lower price. However, with the advancement in health care information systems it may be possible in the future to have schemes where the price paid for the treatment of a given patient is directly related to their outcome.

Non outcome based coverage decisions

With non outcome based coverage decisions effective prices can be determined at an individual patient level or at a population level (i.e. across all patients).

a) Patient level non outcome coverage decisions

Patient level non outcome coverage decisions are characterised by different effective prices for a given technology for different patients, but this is not achieved by linking prices to measures of outcome; rather, prices are linked to other factors associated with treatment. Three types of such agreements can be identified: discounted treatment initiation, utilisation caps and fixed cost per patient.

i) Discounted treatment initiation

Discounted treatment initiation schemes involve patients receiving a technology for a price which is different to the list price at the initiation of treatment. The price then reverts to the list price if the patient remains on the treatment after a set number of courses or period of time. Unlike with conditional treatment initiation schemes, there is no agreement that that patients will remain on treatment should they achieve a target effect. In the UK, the first cycle of sunitinib for the treatment of renal cell carcinoma is provided free for all patients (NICE 2009).

ii) Utilisation caps

Utilisation caps, or individual volume agreements, involve the cost of treatment of patients being reduced (often falling to zero) following an agreed length of treatment if the patient is judged still to require further treatment. These arrangements shift the risk of the health system incurring the cost of the patient undergoing more treatments than a specified maximum from the payer to the manufacturer. Novartis agreed a utilisation cap in the UK for the use of ranibizumab for the treatment of wet acute macular degeneration (Economist Intelligence Unit 2008). The NHS covers

the cost of 14 injections in the treated eye with any subsequent injections being covered by Novartis.

iii) Fixed cost per patient

A coverage decision of a fixed cost per patient involves a set price for an entire course of a patient's treatment regardless of the number of treatments actually received. Such a decision still involves a risk share between the manufacturer and the purchaser with the purchaser paying more for patients who require very few treatments but less for those who require many. In the UK such a scheme has been agreed for the use of gefitinib for the treatment of non-small cell lung cancer (NICE 2010).

b) Population level non outcome coverage decisions

Population level non outcome coverage decisions are characterised by the effective price being determined at the level of the health care system rather than the individual patient. Three forms of such decisions can be distinguished: price changes, expenditure caps and price volume agreements.

i) Price changes

Price changes involve the negotiation of a price per unit of the technology between the manufacturer and the purchaser which differs from the list price. Examples of such agreements are limited, however, as a result of global reference pricing, where pricing of a drug in one country is linked to the price in another. As a result of this most such agreements would not be published as the manufacturer would fear price erosion of their technology in other markets as well.

ii) Expenditure caps

Expenditure caps limit the total expenditure by a health system on a treatment without limiting the total quantity of the treatment available, effectively linking the price of a treatment directly to how much of it is used. These agreements also involve a risk shift, with the manufacturer facing the risk of treatment levels being above those consistent with the expenditure cap. In Australia an agreement was reached that the Australian Health Authority would cover \$100 million (Australian) a year of the cost of etanercept for the treatment of rheumatoid arthritis, with the manufacturer meeting any costs in excess of this (Lu et al. 2004). It was estimated that the treatment would cost \$140 million a year.

iii) Price volume agreements

Price volume agreements link the price paid per unit for a health care technology to the total number of units purchased. An example of such a decision is non linear pricing schemes where the price paid for a unit differs dependent on the total number of units purchased, perhaps being reduced once a threshold number of treatments purchased is reached.

Evidence generation

Evidence generation schemes allow access to technologies whilst also providing evidence so as to reduce the uncertainty surrounding the value of the technologies. These schemes fall into one of two groups: OIR, where the therapy is only paid for in patients involved in the research; and OWR where all patients are given access to the technology but new evidence is also generated. This distinction is the same as that considered earlier, but further distinctions can be made in how the evidence generated is used as discussed below.

Only in research

Evidence generated under OIR can be used to inform coverage decisions in one of three ways:

i) Coverage only

These schemes involve collecting additional evidence to inform a future decision on wider coverage of the technology, perhaps at a set date or when the evidence base is considered sufficiently mature to revise the decision. In the UK, NICE has recommended many treatments only be used in the context of randomised trials or studies. The use of laparoscopic surgery for colorectal cancer was initially rejected by NICE and recommended only in the context of a clinical trial which was ongoing. When the treatment was reconsidered at a later date it received positive guidance on the basis of evidence provided by the clinical trial (Chalkidou 2006).

ii) Refund and coverage

In addition to the arrangements used outlined for coverage only, this category of coverage decision also involves some form of refund or rebate from the manufacturer to the health care system if the additional research suggests that the price paid for the technology for those patients participating in research resulted in the technology not being of value.

iii) Conditional pricing agreement

Conditional pricing agreements involve an agreement between manufacturer and purchaser that the price of the technology will be determined directly by the results of the research. Such schemes may also include a refund linked to the cost of treatment for the research if the results show the treatment is not of value.

Only with research

Only with research schemes allow all patients access to the technology, but also provide for the generation of evidence to reduce uncertainty. Such schemes can be used to inform future coverage decisions in the same ways as those described under OIR.

An example of a coverage only scheme is the Swedish agreement on the use of rosuvastatin for high cholesterol, whereby the manufacturer agreed to provide additional data on the use of the drug in clinical practice and the long term effects of the drug on morbidity and mortality (Anell and Persson 2005). Another is the French authorities' agreement to a refund and coverage scheme to cover risperidone at list price for patients with schizophrenia whilst the manufacturer performed further studies to evaluate whether the treatment helped patients stay on their medication. If the studies showed this not to be the case the manufacturer would refund a portion of the money spent on the treatment (Whalen 2007).

The UK's NHS agreed a conditional pricing arrangement regarding the use of interferon beta or glatiramer acetate for the treatment of multiple sclerosis. The treatments were funded on the condition that their effect on disease progression in a cohort of patients was monitored for 10 years. Potential price adjustments were to be made every 2 years to ensure an agreed cost per QALY gained of the therapy was no more than £36,000 (Pickin et al. 2009).

Discussion

The concepts described in this paper are intended to provide a set of guiding principles for purchasers when considering whether to pay for new health care technologies. Even if these concepts are not implemented using formal analysis they should still be in the forefront of the purchaser's thoughts when making such decisions.

The paper also sets out a stylized version of the technology space and the purchaser's range of authority and shows the coverage decisions which should be considered under the resulting 36 possible combinations. The greater the purchaser's range of authority, the greater the coverage options available and, therefore, the increased scope to optimise decisions. In many countries the range of authority described above is split between several decision making bodies - for example, with a purchaser having control of payment for use decisions and another body having control of research funding. The conceptual framework, however, demonstrates that these are intrinsically linked and both have real effects on health outcomes. The separation of these areas of authority, or the failure of purchasers and other decision makers to act together, may, therefore, result in inefficient decisions by limiting the coverage options available.

It should be noted that only uncertainty which can be reduced through research has been considered here. As previously mentioned, there are other sources of uncertainty which cannot be resolved by further research but might be resolved over time (e.g., the future price of a technology or its comparators). In the presence of such uncertainty it may be optimal to reject a technology, which is expected to be value for money given existing evidence, and to wait until the uncertainty has been resolved.

The taxonomy provided here aims to assist the understanding of the coverage decisions observed in practice. Table A in Appendix A provides details of coverage decisions from several countries and allocates the decision in accordance with the taxonomy. In terms of schemes involving reductions in expected effective price, many were observed in practice on the basis of information available in the public domain. Seven examples of money back guarantee schemes are presented, of which four were observed in the UK, two in Germany and one in the USA. Ten versions of conditional treatment continuation schemes have been identified in the UK, the USA, Australia, Italy and Canada. No examples of decisions where the price was directly linked to outcome were found, this may be as a result of the large transaction and information costs associated with such schemes. Only one example of each of a discounted treatment initiation, utilisation cap and fixed cost per patient were observed, all of which were in the UK. No examples were found where there was a change in the list price agreed between the purchaser and the manufacturer, although it is likely that such agreements would remain private as a result of the impact they could have on prices in other countries through global reference pricing. Examples of expenditure caps were observed in Australia and Germany.

Many examples of schemes involving evidence generation were observed. Most involved OIR decisions where the evidence was to be used to inform future coverage decisions (with such approaches observed in the UK and USA). The UK also had one example of an OIR scheme where the evidence was directly linked to the price (an OIR conditional flexible pricing schemes). OWR schemes were less commonly observed in practice, with the exception of Sweden where such decisions are common. Given the size of the USA there have been surprisingly few examples there so far of these new coverage decisions. A key factor may be the unwillingness of the private sector in the USA to adopt such schemes, particularly with regards to those coverage decisions involving evidence generation, where the benefit is not fully internalised by the purchaser as a result of the public good properties of information.

Another set of schemes which have emerged but which are not dealt with explicitly in our taxonomy are cost-sharing schemes whereby patients cover some or all of the cost of treatments that are not deemed to meet acceptable value for money standards (Austvoll-Dahlgren et al. 2008; Jackson 2010). These have not been dealt with explicitly here as they involve more complex issues around patient choices and preferences particularly when considered across different types of health systems.

Our conceptual framework was developed around the concept of the expected value of a new technology on the basis of existing evidence. This is described principally in terms of cost effectiveness as defined by organisations such as NICE in the UK - that is, whether a new technology generates more health gain to relevant patients than is forgone to others as a result of services being displaced to fund the new intervention. It is recognised that some jurisdictions do not use formal cost-effectiveness analysis to support coverage decisions, but it can be argued that such systems (e.g. Medicare in the USA and the French health system) conduct an implicit comparison of the magnitude of gain in effectiveness offered by a new technology, against relevant comparators, and its acquisition cost. Any assessment of value for money needs to consider which consequences of new technologies are valued. Our conceptual framework simplifies this to a concern only with aggregate health gain in a population. In reality other factors may be important to purchasers including a range of equity considerations and the need to encourage innovation amongst technology manufacturers. Although these wider considerations complicate the technical implementation of cost effectiveness, the value of the conceptual framework remains in showing the types of assessments purchasers need to make in selecting appropriate coverage options.

Conclusions

Several key policy implications fall out of this paper. Firstly, coverage decisions should no longer be based solely on the expected value given available evidence. Instead the purchaser should weigh the expected value, the value of further evidence, the effect on evidence generation of immediate coverage and the reversal costs for uncertain decisions. Secondly, this paper demonstrates the importance of a purchaser's range of authority: given a certain range of authority, there is a set of coverage decisions potentially available to purchasers. However, in many cases, there is a separation of the areas of authority, for example in many countries a purchaser decides on payment for use decisions whilst another decision making body is in control of research. However, these are intrinsically linked and both have real effects on the health gain produced by health care systems. The separation of these areas of authority, or the failure of purchasers and other decision makers to act together, may result in inefficient decisions.

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Appendix A

Disease area	Product	Agreement	Type of scheme
UK			
High cholesterol	Statins	Manufacturer agreed to rebate the North Staffordshire health authority if patients did not achieve a low density lipoprotein cholesterol concentration target of < 3 mmol/l after treatment with statins (Chapman et al. 2004).	Money back guarantee
Asthma	Omaluzimab	Manufacturer offers replacement products for appropriately diagnosed patients who fail to achieve an agreed target clinical response (Sparrowhawk 2007).	Money back guarantee
Colorectal cancer	Cetuximab	Rebate direct to primary care trust on the cost of any vials of cetuximab used for patients who do not achieve a pre-agreed clinical outcome ('nonresponders') at up to 6 weeks (with a maximum rebate of 3200 milligrams) (Thomson 2008).	Money back guarantee
Multiple myeloma	Lenalidomide	The manufacturer has agreed to meet the drug cost of lenalidomide for all patients who have received 2 or more previous therapies and who remain on therapy for more than 26 weeks (Gajraj et al. 2009).	Money back guarantee
Multiple myeloma	Bortezomib	J & J agreed to reimburse the NHS in either cash or product for patients who do not respond after 4 cycles of treatment with bortezomib. Responding patients receive an additional 4 cycles (Green et al. 2006).	Conditional treatment continuation
Renal cell carcinoma	sunitinib	The first cycle of sunitib for the treatment of renal cell carcinoma is free to all patients (NICE 2009).	Discounted treatment initiation
Macular degeneration	Ranibizumab	The NHS will pay for up to a maximum of 14 injections with any further treatments being reimbursed by the manufacturer (Economist Intelligence Unit 2008).	Utilisation cap
Non-small cell lung cancer	Gefitinib	A fixed cost per patient scheme was agreed for the use of gefitinib for the treatment of non small cell lung cancer (NICE 2010).	Fixed cost per patient
Chronic myelogenous leukemia (CML)	Gleevec (Imatinib Mesylate)	If a person has been taking imatinib for CML while in the chronic phase, but still goes on to the accelerated or blast crisis phase, NICE has recommended that imatinib treatment is continued only as part of a research study (NICE 2002).	OIR- Coverage only
Brain cancer	Temozolomide	Temozolomide is only recommended as the initial chemotherapy treatment for patients with brain cancer when they are taking part in a clinical trial (NICE 2001).	OIR- Coverage only
Age-related macular degeneration	Photodynamic therapy (PDT)	PDT is not recommended for people who have wet ARMD with mostly classic subfoveal CNV (that is, at least half is classic but there is also some occult CNV). The exception is where the person is treated as part of a clinical study designed to provide useful information on the effectiveness of the treatment (NICE 2003a).	OIR- Coverage only
Breast cancer	Taxanes	The use of taxanes for adjuvant treatment of early breast cancer should be limited to randomized clinical trials. Docetaxel was eventually recommended for breast cancer whilst paclitaxel was not (Chalkidou 2006).	OIR- Coverage only

Colorectal cancer	Laparoscopic surgery	The use of laparoscopic surgery for colorectal cancer should be limited to randomized clinical trials. The treatment was eventually recommended.(Chalkidou 2006)	OIR- Coverage only
Colorectal cancer	Oxaliplatin and irinotecan	Neither treatment was recommended for first line treatment of advanced colorectal cancer except as part of a clinical trial. Both treatments were eventually recommended (Chalkidou 2006).	OIR- Coverage only
Alzheimer's disease	Memantine	Memantine is not recommended as a treatment option for patients with moderately severe to severe Alzheimer's disease except as part of well designed clinical studies (Chalkidou 2006).	OIR- Coverage only
Non-Hodgkin's lymphoma	Rituximab	NICE has said that rituximab should only be used in people with diffuse large-B-cell lymphoma in stage 1 if they are being treated as part of a clinical study (NICE 2003b).	OIR- Coverage only
Diabetes	Metformin	Metformin in combination with insulin should only be used within research studies because the effectiveness of this combined treatment in improving glycaemic control is uncertain (NICE 2004).	OIR- Coverage only
Lung cancer	Post-operative chemoradiotherapy	Patients who are pathologically staged as II and III non-small cell lung cancer following resection should not receive post-operative chemoradiotherapy unless it is within a clinical trial (NICE 2005).	OIR- Coverage only
Multiple Sclerosis	Interferon beta's or glatiramer acetate	Drugs were funded on the condition that their effect on disease progression in a cohort of patients was monitored for 10 years. Potential adjustments in price would be made every 2 years to ensure an agreed cost per QALY of no more than £36,000 (Pickin et al. 2009).	OIR- Conditional flexible pricing agreement
		PPRS- Financially based patient access schemes- the company does not alter list price but offers effective discounts or rebates linked to, for example: i) Number of patients treated (price/volume agreement) ii) Number of doses required iii) Response of patients treated (Department of Health 2008).	i) Price volume ii) Utilisation cap iii) Money back guarantee / Price linked to outcome
		PPRS- Outcome based patient access schemes- 3 types: i) Proven value price increase- company seeks agreement for later increases in price subject to a re-review of the drug in light of additional evidence collected ii) Expected value rebate- Company seeks agreement for a price subject to the collection of additional evidence. The drug will be subject to a rebated and a subsequent reduction in list price in the event of the additional evidence not supporting the original price iii) Risk sharing- outcomes are measured and prices adjusted &/or cash transferred in one or both directions (Department of Health 2008).	i) OWR- Conditional flexible pricing agreement ii) OWR- Conditional flexible pricing agreement & Refund and coverage iii) OWR- Conditional flexible pricing agreement & Refund and coverage

USA			
High cholesterol	Simvastatin	Merck promised to refund up to six months of prescription costs if simvastatin plus diet did not help lower LDL cholesterol to target concentrations identified by their doctors (Moldrup 2005).	Money back guarantee
Hemodialysis	Erythropoiesis-stimulating agents	CMS will reimburse erythropoiesis-stimulating agents until the patient achieves a hemoglobin level of 10g per dl (Berns et al. 2005).	Conditional treatment continuation
Type 2 diabetes	Sitagliptin	Manufacturer has agreed to link the price paid for sitagliptin (and sitagliptin plus metformin) to how well patients are able to control their blood sugar (Pollack 2009).	
Osteoporosis	Risedronate sodium	Manufacturers have agreed to reimburse insurer for the costs of treating related fractures (Pollack 2009).	
Cognitive impairment	FDG-Positron emission tomography (PET) scan	An FDG-PET scan is covered in patients with mild cognitive impairment or early dementia in the context of a clinical trial (Centers for Medicare and Medicaid Services).	OIR- Coverage only
Hearing loss	Cochlear implant	CMS may cover cochlear implantation for treatment of hearing loss in the context of an approved clinical trial (Centers for Medicare and Medicaid Services).	OIR- Coverage only
Oncology	FDG-PET scan	An FDG-PET scan is covered in patients with brain, ovarian, pancreatic, small cell lung, testicular cancers, and certain indications for cervical cancer in the context of an approved clinical registry (Centers for Medicare and Medicaid Services).	OIR- Coverage only
Tachyarrhythmia's	Implantable Cardioverter Defibrillator	Implantable Cardioverter Defibrillators are covered in the context of an approved clinical trial or registry (Centers for Medicare and Medicaid Services).	OIR- Coverage only
Chronic hypoxemia	Home use of oxygen	The home use of oxygen is covered for those beneficiaries with arterial oxygen partial pressure measurements from 56 to 65 mmHg or oxygen saturation at or above 89% who are enrolled subjects in clinical trials approved by CMS and sponsored by the National Heart, Lung & Blood Institute (NHLBI) (Centers for Medicare and Medicaid Services).	OIR- Coverage only
Atherosclerotic disease	Angioplasty and Stenting	CMS covers Percutaneous Transluminal Angioplasty and Stenting of intracranial arteries for the treatment of cerebral artery stenosis $\geq 50\%$ in patients with intracranial atherosclerotic disease when furnished in an approved clinical trial (Carlson et al. 2010).	OIR- Coverage only
Colorectal cancer	Oxaliplatin, irinotecan, cetuximab, bevacizumab	Oxaliplatin, irinotecan, cetuximab, or bevacizumab for the treatment of colorectal cancer are covered in the context of an approved clinical trial sponsored by the National Cancer Institute (Carino et al. 2006; Centers for Medicare and Medicaid Services).	OIR- Coverage only
Emphysema	Lung volume reduction surgery	Medicare would fund the treatment only as part of a clinical trial (Carino, Sheingold, and Tunis 2004).	OIR- Coverage only
Breast Cancer	OncotypeDx	United Healthcare agreed to reimburse the OncotypeDx test for 18 months while it and Genomic Health monitor the results. If the number of women receiving chemotherapy exceeds an agreed upon threshold, even if the test suggests they do not need it, the insurer will negotiate a lower price (Pollack 2007).	OWR-

Sweden			
High cholesterol	Rosuvastatin and ezetimibe	The companies had to provide additional data on the use of the drug in clinical practice and the long-terms effect of the drug on morbidity and mortality (Anell and Persson 2005).	OWR- Coverage only
Diabetes Mellitus	Insulin detemir	The company must provide additional data on the frequency of hypoglycaemic events (Anell and Persson 2005).	OWR- Coverage only
Diabetes Mellitus	Insulin glargine	The company had to provide additional data on cost-effectiveness for use among patients with type 2 diabetes (Anell and Persson 2005).	OWR- Coverage only
Diabetes Mellitus	Inhalable insulin	The company shall provide additional data to support the economic value of inhalable insulin in a Swedish clinical day-to-day setting (Carlson et al. 2010).	OWR- Coverage only
Obesity	Orlistat and sibutramine	The companies had to provide data of actual use in the Swedish health care system (Anell and Persson 2005).	OWR- Coverage only
Testosterone therapy	Testogel	The company had to provide data of actual use in the Swedish health care system (Anell and Persson 2005).	OWR- Coverage only
Eczema.	Pimecrolimus	The company shall provide additional data on the effect of pimecrolimus for patients who are resistant to steroid-treatment and its use in the Swedish clinical day-to-day setting (Carlson et al. 2010).	OWR- Coverage only
Type 2 diabetes and Obesity	Rimonabant	The company shall provide additional data on the long-term effects of rimonabant and its economic value in a Swedish clinical day-to-day setting (Carlson et al. 2010).	OWR- Coverage only
Parkinson's disease	Rasagiline	The company shall provide additional data on the cost-effectiveness of rasagiline versus entakapon and selegilin (Carlson et al. 2010).	OWR- Coverage only
Smoking cessation	Varenicline	The company shall provide additional data on the long-term effects of varenicline (Carlson et al. 2010).	OWR- Coverage only
Cervical cancer	Human papillomavirus quadrivalent	The company shall provide additional data on ongoing and planned studies in order to determine the cost-effectiveness from a long-term perspective (Carlson et al. 2010).	OWR- Coverage only
Grass pollen allergy	Lyophilisate	The company shall provide additional data on the long-term effects of lyophilisate and a new health-economic evaluation based on costs and medical effects of the drug in clinical practice (Carlson et al. 2010).	OWR- Coverage only
Parkinson's disease	Rotigotine	The company shall provide additional data the effect of rotigotine in the Swedish clinical day-to-day setting (Carlson et al. 2010).	OWR- Coverage only
Psoriasis	Efalizumab	The company shall provide additional data on quality of life and effectiveness from use in real clinical treatment (Anell and Persson 2005).	OWR- Coverage only

Schizophrenia	Risperidone	The company shall provide data on hospitalisation and quality of life from use in real clinical treatment (Anell and Persson 2005).	OWR- Coverage only
France			
Schizophrenia	Risperidone	France's health care authority agreed to cover risperidone at list price if the manufacturer performed further studies to evaluate whether treatment helps patients stay on their medications. If the studies show otherwise, the manufacturer will partially reimburse France for spending on the treatment (Whalen 2007).	OWR- Refund and Coverage
Australia			
Rheumatoid arthritis	Etanercept, infliximab, adalimumab and anakinra	Australian health authority will condition coverage of these biological on an assessment of effectiveness at 12 weeks. Coverage will continue for patients in whom it is effective (Lu, Williams, and Day 2007).	Conditional treatment continuation
Ankylosing spondylitis	Infliximab and etanercept	Coverage for infliximab and etanercept for ankylosing spondylitis is conditioned on an assessment of short-term effectiveness. Coverage will continue for patients in whom it is effective (Carlson et al. 2010).	Conditional treatment continuation
Chronic myelogenous leukaemia	Imatinib mesylate	Coverage is conditioned on an assessment of short-term effectiveness evaluated at 18 months. Coverage will continue only for those patients in whom it is effective (Carlson et al. 2010).	Conditional treatment continuation
Rheumatoid arthritis	Etanercept (enbrel)	The manufacturer will cover all expenditure over \$100 million (Lu et al. 2004).	Expenditure cap
Pulmonary arterial hypertension	Bosentan	The price of bosentan for pulmonary arterial hypertension is linked to the survival of patients followed in an observational study (Włodarczyk et al. 2006).	OWR- Conditional flexible pricing agreement
Italy			
Renal cell carcinoma	Sunitinib and sorafenib	A hospital discount of 50% applies to the first 2/3 months of treatment with sorafenib and sunitinib. For responding patients, the treatment is then reimbursed and the discount dropped (Carlson et al. 2010; IMS 2007).	Conditional treatment continuation
Alzheimer's disease	Alzheimer's disease drugs	During first 3 months, patients starting Alzheimer's disease drugs are assessed for short-term effectiveness with the drug provided free by manufacturer during this period. If treatment goals are met after 3 months, treatment is continued for a max of 2 years (Sparrowhawk 2007).	Conditional treatment continuation
Small cell lung cancer	Erlotinib	Price is only 50% for first 2 months on the basis that 50% of patients are expected to progress in this period (Rutten, Uyl-de Groot, and Vulto 2009).	Conditional treatment continuation
Chronic myelogenous leukemia	Nilotinib	Cost of treatment will be refunded for all patients who do not achieve an agreed response after one month (Wagstaff 2009).	Conditional treatment continuation

Canada			
Alzheimer's disease	Donepezil, rivastigmine and galantamine	Patients using donepezil, rivastigmine, or galantamine will be reimbursed for a period of up to 3 months after which further coverage will be made available to those patients whose disease has not progressed/deteriorated while on this drug (Carlson et al. 2010).	Conditional treatment continuation
Germany			
Kidney transplantation	Cyclosporine, mycophenol acid or everolimus)	Manufacturer has agreed to refund money for cyclosporin, mycophenol acid or everolimus if a patient loses his/her donor kidney (Carlson et al. 2010).	Money back guarantee
Osteoporosis	Zoledronic acid	DAK and Barmer (a German insurance company) have a money back guarantee for zoledronic acid if an osteoporosis related fracture occurs (Carlson et al. 2010).	Money back guarantee
Wet aged macular degeneration	Ranibizunab	The payer will not have to pay more than 315 million euro a year (Rutten, Uyl-de Groot, and Vulto 2009).	Expenditure cap