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Citation: Dimakou, S., Parkin, D., Devlin, N. & Appleby, J. (2006). The impact of government targets on waiting times for elective surgery: new insights from time-to-event analysis (06/05). London, UK: Department of Economics, City University London.

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The impact of government targets on waiting times for elective surgery: new insights from time-to-event analysis

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Discussion Paper Series

No. 06/05

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Introduction

Waiting lists for elective surgery - that is, routine, non-emergency clinical procedures - are a common feature of collectively-financed health care systems where coverage is universal and consumers face zero price at the point of demand. Waiting lists function, in part, as a non-price rationing device to reconcile differences between supply and demand. The number waiting at any point in time is determined by the rate at which people leave the list - by being admitted for surgery, self-deferring, being removed for clinical reasons or dying - relative to the rate at which people join the list, as a result of a decision by a consultant (Boyle and Appleby, 2001). Recent work by Gravelle, Smith and Xavier (2003) has demonstrated that waiting times and waiting list sizes act as signals that have an impact upon both supply and demand.

Long waiting lists and extensive waiting times for elective surgery have been a persistent source of policy, political and popular concern in the UK and other OECD countries for many years (Siciliani and Hurst, 2003). Notwithstanding the considerable resources and effort that have been directed to reducing waiting lists and waiting times in England since the NHS Plan (Department of Health, 2000), the number waiting for inpatient treatment is currently around 900,000, 9% of whom have waited six or more months; a further two million are waiting for an outpatient appointment (Department of Health, 2004).

Policies to reduce waiting lists can include both supply-side responses, for example extra funding for elective surgery, tackling supply bottlenecks, provider monitoring and management of waiting lists; and demand management, for example promulgating guidelines for appropriate referral and explicit methods for prioritising patients. Historically, NHS policy on waiting tended to reflect a view that waiting lists were a backlog of untreated patients, a problem that could be ameliorated by short-term increases in activity (Hamblin, Harrison and Boyle, 1998).

More recently, the emphasis of policy has shifted from waiting *lists* to waiting *times*, on the grounds that patients are more concerned about the speed with which the queue moves - and thus the time they spent on the list - rather than the number of people waiting in front of them.

While current policy combines a number of the supply- and demand-side strategies noted above, the main strategy has been to use waiting times targets (Harrison and Appleby, 2005). These take the form of maximum waiting times for elective surgery that providers should meet - with associated rewards and penalties for successful and unsuccessful performance. The waiting time target for inpatient elective surgery to be achieved by 2005 is 6 months, with intermediate targets of 18 months by March 2000; 15 months by March 2002, 12 months by March 2003 and 9 months by March 2004 (Appleby *et al*, 2005). Furthermore, the government has pledged that by 2008 there will be a maximum wait of only 18 weeks from initial referral of a patient by a general practitioner to inpatient treatment if required.

Although there has been some success in reducing waiting lists and times, such as the significant reductions in long waits since 2000, there are a number of concerns specifically relating to the use and impact of these targets. Principal among these is the extent to which targets distort clinical priorities, by changing the order, and thus speed, with which patients are treated. The National Audit Office reports that 20% of consultants surveyed in three specialties stated they had changed the order they had prioritised patients so as to meet the corresponding target (NAO, 2001).

In one respect, this changed order of priorities is not an unintended side effect, but indeed the entire point. Although there is evidence to suggest that the length of a patient's wait may have influenced clinical decisions to admit *even before* the introduction of targets (Appleby *et al*, 2005), presumably the targets reflect an explicit view that whatever clinical or social factors determined priority for treatment did not place sufficient weight on time waited, in particular, maximum time waited. However, if providers are meeting targets by substituting less urgent cases, with less ability to benefit, for more urgent cases, with higher ability to benefit, then this would be a potential cause for concern on both economic and ethical grounds.

The challenge in analysing the number, importance and effect of these changes in admission decisions arising from the targets is that the admission criteria without targets are neither clearly specified nor consistent. Individual clinicians assess patients' conditions according to their own personal judgements of clinical urgency. There are neither 'gold standard' admission criteria nor any systematic scoring system in widespread use in the UK to aid between-patient prioritisation. Culyer and Cullis (1976) advocated such an approach over 25 years ago, and there are examples of such systems from other countries (Siciliani and Hurst, 2003; Hadorn and Holmes, 1997).

More fundamentally, neither the way in which providers meet the targets nor what differentiates 'successful' from 'unsuccessful' trusts with respect to the targets are clear. For example, targets may be met principally by increasing surgical throughput - reducing waiting times for all patients - or by substituting the treatment of low wait patients with high wait patients, or a mix of both. The targets create incentives which might be expected to affect both manager and clinician behaviour.

Appleby *et al* (2005) provided some evidence on these issues for orthopaedic surgery. Waiting times distributions were compared before and after the introduction of targets, with differences in the distributions used to identify changes in admission patterns. The results appear to suggest that "...any reordering of cases had less to do with substituting very short wait (presumed urgent) cases with longer wait (presumed less urgent) cases but rather that the latter displaced some (less urgent) 'filler cases' - that is, those with short operating times which could be used to make best use of available theatre time". However, given the reliance on relatively crude before-and-after comparisons of waiting times distributions, this interpretation remains somewhat speculative.

This paper investigates these issues, using the techniques of time-to-event (survival) analysis. Our aim is to address the following questions: how have behavioural responses to the targets influenced the distribution of waiting times? How is the

probability of admission for any given waiting time affected by the targets? To what extent are clinical distortions evident in the pattern of admissions? Can variations between individuals' waiting times be explained by clinical, patient or provider-level characteristics? In addressing these questions, there is an additional aim, to assess whether or not time-to-event analysis provides additional insights and therefore whether or not it should be more widely applied to waiting time data.

Applying time-to-event analysis to waiting list data

Time-to-event analysis has been widely used in social and economic sciences, where it is also known as duration analysis, in biomedical sciences, where it is known as survival analysis, and in engineering, where it is known as failure-time analysis. It consists of a set of parametric and non-parametric methods for estimating survivor and hazard functions, which are explained below, allowing the comparison of the survival of different groups and estimating the impact of explanatory variables on survival. (See, for example, Cox and Oakes, 1984; Collett, 1994). In the context of this paper, 'survival' means remaining on a waiting list, the 'event' is admittance to a hospital and 'time' is that between being placed on a waiting list and being admitted.

Time-to-event analysis offers several advantages in analysing waiting time data. First, because the analysis is performed on individual record data, it may generate deeper insights than methods that focus on comparison of average waiting times. Secondly, survival analysis techniques are appropriate given that waiting times are not usually normally distributed. Thirdly, it addresses the problem of censored observations, which contain only partial information about waiting times, such as for patients who have been entered onto a waiting list but have not been admitted and patients who have been admitted, but for whom the date of entering the waiting list is not known. However, in this study the data set did not include censored observations.

Two key concepts in time-to-event analysis are *survival functions* and *hazard functions*. A survival function shows the conditional probability of a person surviving on the waiting list until a given time, which is a cumulative density function derived from the unconditional probability of survival. Such functions can be modelled parametrically, by assuming a particular distribution, but can also be estimated empirically usually through the application of the Kaplan-Meier or product limit estimator. In effect, this shows the rate at which people leave the waiting list and the variations in this rate over time. It also provides an estimate of the average waiting time as the integral of the survival function, though for data that are not censored this is merely an alternative calculation to a straightforward mean. An advantage of survival functions over means in this context is observing patterns of waiting list behaviour over time – the same average waiting time might be generated by very different means of managing lists over time. Survival functions can also be compared between different groups, defined for example by illness, treatment, doctor or patient characteristic, and differences can be tested statistically using the log-rank test. In addition, the impact of variables that affect waiting time patterns can be analysed.

The hazard function shows the probability of a person leaving the waiting list at a given time, conditioned by the probability that they remained on the waiting list until

that time. The survival and hazard functions are mathematically related, and any given hazard function generates a particular survival function and *vice versa*. For example, if the hazard function is constant – the instantaneous conditional probability remains the same at all times – this generates an exponential survival function of the form $S(t) = e^{-\lambda t}$, where λ is the hazard rate. The advantage of this is that it may reveal patterns of waiting list behaviour that would not otherwise be apparent – for example, if management effort in clearing waiting lists varies over time, so that patients have a varying probability of being admitted that is not related to the length of time that they have already waited.

In the context of analysing the impact of other variables on waiting list patterns, parametric models have two flavours, which depend on assumptions about the hazard rate. *Proportional hazard* (PH) models assume that there is a baseline hazard function that depends on time but not on the other variables and is therefore common to all individuals. The other variables, which are usually assumed to be time-invariant, essentially scale the hazard function for each individual. A valuable technique in estimating PH models is the semi-parametric Cox regression, which does not require any assumption about the hazard rate, simply the impact on it of the other variables. *Accelerated failure time* (AFT) models allow scaling to vary over time. Although these are therefore more flexible, they are entirely reliant on assumptions about the underlying hazard function; there is no equivalent of Cox regression.

Data and Methods

Hospital Episode Statistics (HES) data were used. HES data cover all episodes of care for UK NHS hospital patients provided in NHS hospitals or elsewhere. These are obtained from an annual snapshot of data submitted by NHS Trusts to the NHS-Wide Clearing Service (NWCS). The data are collected by financial year and in each year cover approximately 11 million admitted patient records. We analysed HES data for elective surgery in three specialties - general surgery, trauma and orthopaedics and ophthalmology - for the financial years 2001/2 and 2002/3. We chose these specialties because they constitute more than 50% of the overall proportion of patients waiting for elective treatment. The data include information on specialty, diagnosis, operation, Healthcare Resource Group (HRG), admitting hospital, type of admission and patient characteristics such as age, sex, ethnicity and residence⁵. However, because of confidentiality issues, we were not able to obtain data on individual referring GPs or admitting consultants, even at an anonymised level. Analysis of data that contained such information would obviously be very valuable, but our work only shows the potential of time-to-event analysis in such uses.

Analysis of waiting times included the following:

- Estimation of the survival and hazard functions of waiting times of patients admitted through waiting lists using the Kaplan-Meier estimator.

⁵ The data contained no items that would enable an individual patient to be identified.

- Exploring differences in survival and hazard functions according to specialty, operation, HRG, provider level and admission type, by both graphical methods and by the log-rank test.
- Adjustment of survival and hazard rate functions for the effects of covariates which may impact on waiting times, using parametric Proportional Hazard and Accelerated Failure Time models under different distributional assumptions and Cox regression.

Results

Identifying variations in waiting times for elective surgery in different specialties

Figure 1 shows the survival curves for all patients admitted in the three specialties during 2001/2. The results for 2002/3 are not shown but are very similar. Initial analysis revealed that there are some patients in the HES data who appear to wait for a very long time (~ 3500 days), but as this is most likely the result of coding problems, all patients waiting more than three years - 1095 days - (around 0.1% of all patients) were excluded from the analysis. The curves show the proportion of patients on the waiting lists at each time. At time 0, all patients are on the list and the curve falls as they leave the list by being admitted. At around 600 days the proportion of people remaining on each list is very small. This figure is truncated at 730 days to reveal more informative patterns. Future drafts of this paper will follow the same display.

The shortest waiting times are for general surgery, followed by ophthalmology with orthopaedics having the longest times. The log-rank test for equality of the survivor functions revealed that the differences between waiting times for the three specialties are statistically significant for both financial years. The shapes of the curves are quite different, and the curves for general surgery and ophthalmology cross, so that after the point of intersection (at around 456 days) ophthalmology patients have the shortest times.

Figure 1: Kaplan-Meier survival curves for three specialties - 2001/2

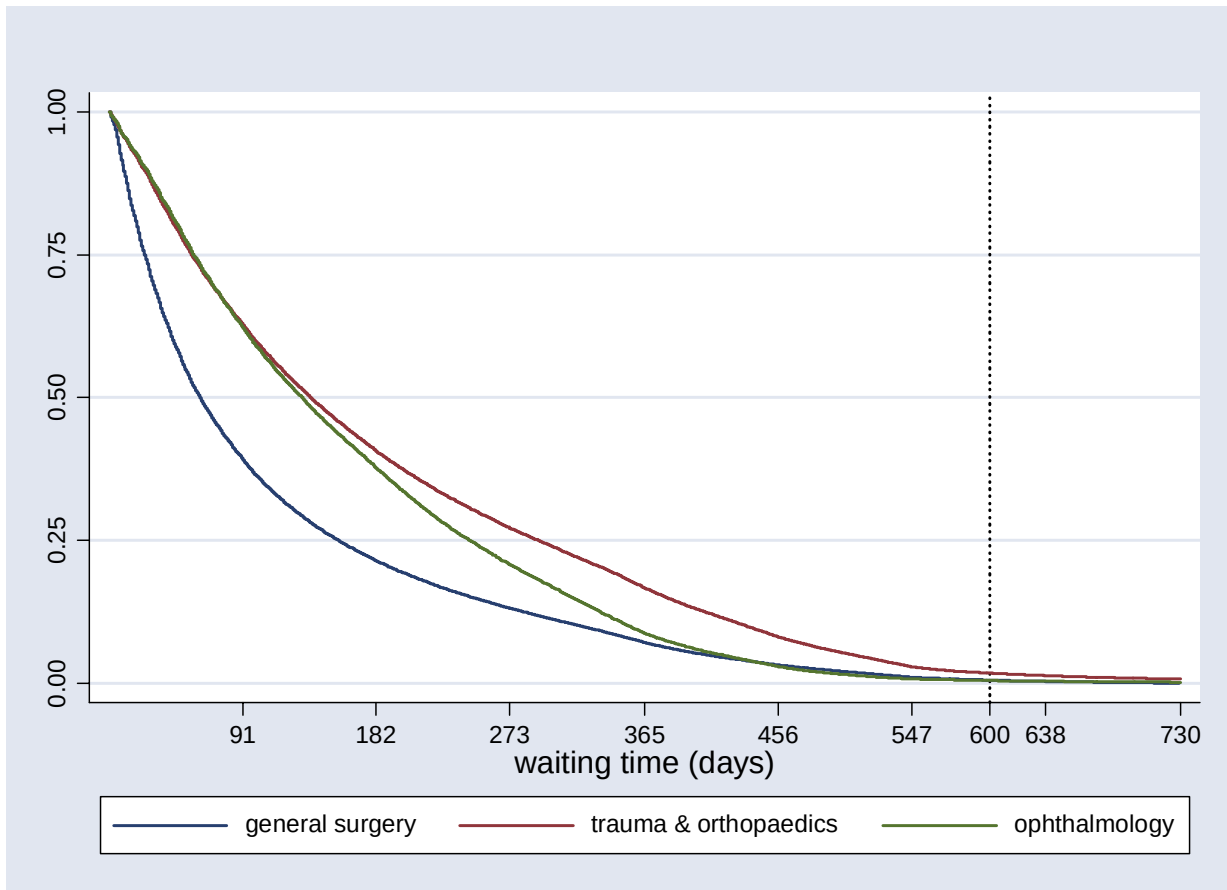


Figure 2: Hazard curves for three specialties - 2001/2



Figure 2 shows the estimated hazard functions for 2001/2. In that year, the national maximum waiting time target was 15 months. The hazard functions show the effects of waiting times targets clearly: the probability of admission for those whose wait approaches a target increases markedly and falls when their wait exceeds the target. The impact is, however, different in the different specialties. For general surgery, increased waiting list activity is observed as a peak in the curve for people waiting 4 and 15 months; for ophthalmology, at exactly 15 months; for orthopaedics between 15 and 18 months.

Although the hazard curves for 2002/3 are not shown, they suggest that as the targets waits become shorter, the peaks changed, in each case occurring at a lower waiting time. For general surgery, the main peak was at 12 months, which coincides with the waiting time target for that year; for orthopaedics between 12 and 15 months; for ophthalmology, around 14 months.

Identifying variations of waiting times for different operative procedures

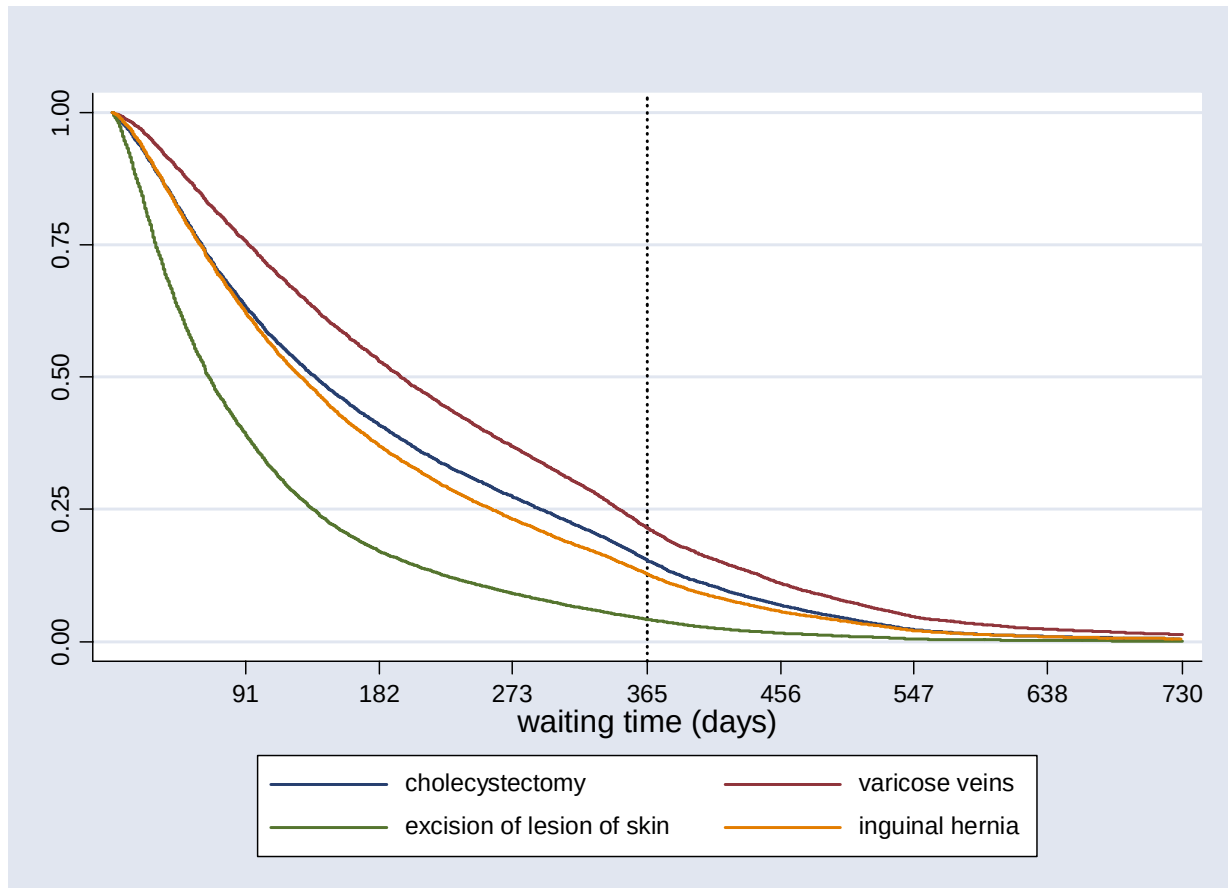
Given that waiting lists may be managed for particular operations, rather than specialties as a whole, waiting time patterns between operations may differ. Waiting times were therefore analysed for the most four frequently occurring operative procedures within each specialty:

<i>General surgery</i>	
Excision of gall bladder (total cholecystectomy) Ligation of varicose veins of leg Excision of lesion of skin Primary repair of inguinal hernia	These 4 operative procedures account for 26% of all general surgeries
<i>Trauma & Orthopaedics</i>	
Release of entrapment of peripheral nerve at wrist Total prosthetic replacement of hip joint using cement Total prosthetic replacement of knee joint using cement Endoscopic operations on semilunar cartilage	These 4 operative procedures account for 28% of all orthopaedic surgeries
<i>Ophthalmology</i>	
Extirpation of lesion of eyelid Incision of capsule of lens Prosthesis of lens Cauterisation of lesion of retina	These 4 operative procedures account for 71% of all ophthalmologic surgeries, with lens prosthesis being the prevalent one (62% of all ophthalmologic procedures)

In each case, analysis at the level of was also performed; this gave similar results to that at the operation level and is therefore not reported.

Figure 3 shows survival curves for procedures within general surgery for 2001/2. The shortest waiting times were for excision of lesion of skin, followed by inguinal hernia repair, gall bladder excision and varicose vein ligation. The log-rank test for equality of the survivor functions also revealed that these differences were statistically significant. They may be due to the urgency of the operation type, its difficulty and whether or not it can be performed as a day case.

Figure 3: Survival curves for four operative procedures in general surgery – 2001/2

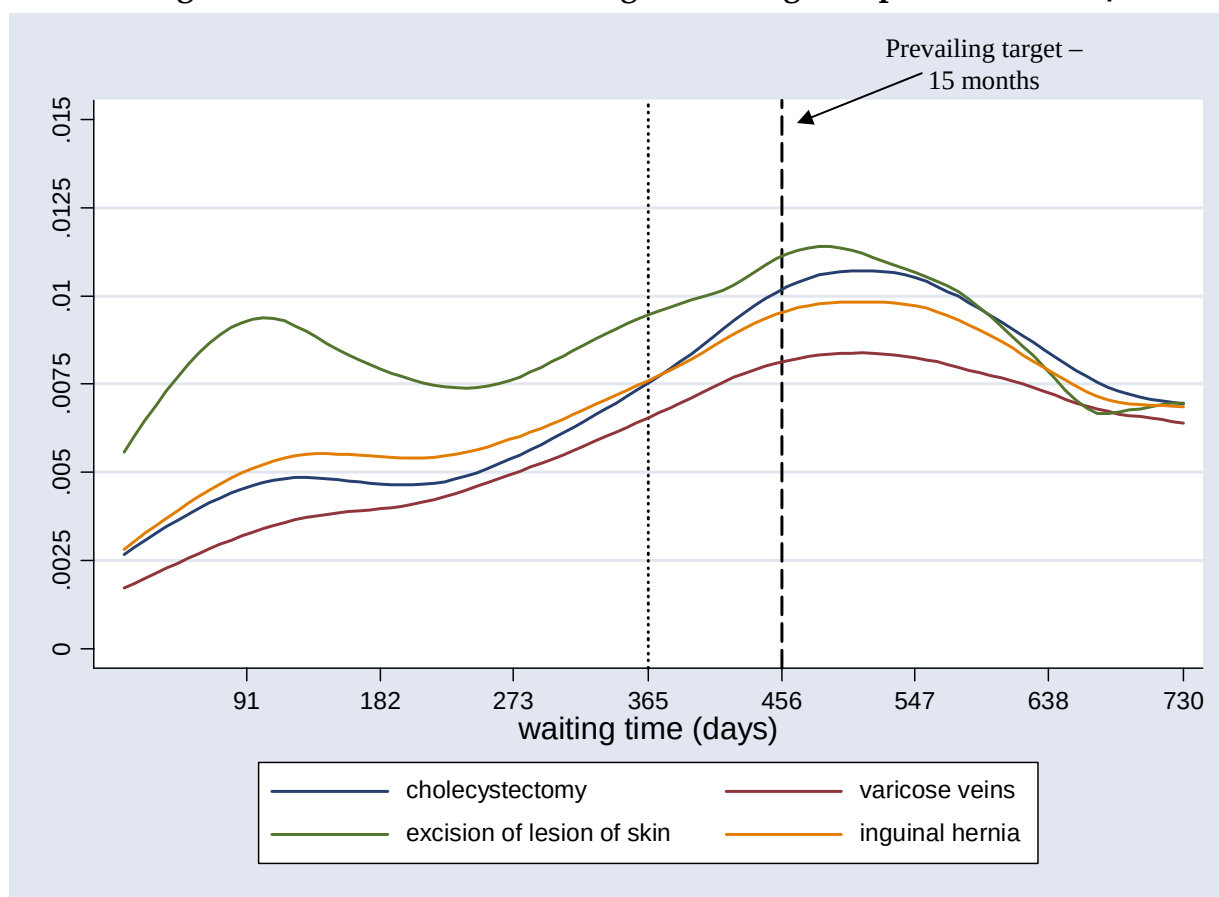


The survival curves for orthopaedic and ophthalmology procedures are not shown, but again there were significant differences between them. In orthopaedics, the order from shortest to longest waiting time was release of entrapment of peripheral nerve at wrist, endoscopic operations on semilunar cartilage, hip replacement and knee replacement. There is a particularly large difference between the curves of the first two procedures compared with the last two. Within ophthalmology, the order was cauterisation of lesion of retina, incision of capsule of lens, extirpation of lesion of eyelid and prosthesis of lens.

Figure 4 shows the hazard curves for general surgical operations for 2001/2. Observed peaks in hazard rates for excision of lesion of skin are around 3 months and 15 months. For 2002/3, the first peak is unchanged, but the second peak reduces to almost 12 months; suggesting that increased activity due to shorter target times affects patient waits that are at or longer than the target, but not those that are shorter than the target. For the other procedures, peaks are observed at just after 15 months for 2001/2 and

between 12 and 15 months for 2002/03, again suggesting a response to the shorter maximum waiting times targets over the two years.

Figure 4: Hazard curves for four general surgical operations – 2001/2



Hazard rates for orthopaedic and ophthalmologic procedures are not shown, but also show differences between procedures and responses to shorter targets. Within orthopaedic procedures, the main peaks for endoscopic operations on semilunar cartilage, hip replacement and knee replacement all reduced from 15 months in 2001/2 to between 12 and 15 months for 2002/3. However, the main peak for release of entrapment of peripheral nerve at wrist increased from between 6 and 12 months in 2001/2 to 12 months in 2002/3 – both of which are shorter than the prevailing target.

Within ophthalmologic procedures, the main peak for prosthesis of lens reduced from 15 months in 2001/2 to between 12-15 months for 2002/3. The other three procedures had far earlier peaks, at around 3 months, which were the same in both years. There were other peaks in 2001/2, between 12 and 15 months for extirpation of lesion of eyelid, a little less than 12 months and around 18 months for incision of capsule of lens, and a little more than 15 months for cauterisation of lesion of retina. In 2002/3, these secondary peaks were between 12-15 months for all three. Again, this suggests activity to achieve the targets was focussed on tackling longer waits.

Identifying variations in waiting times for elective surgery between providers

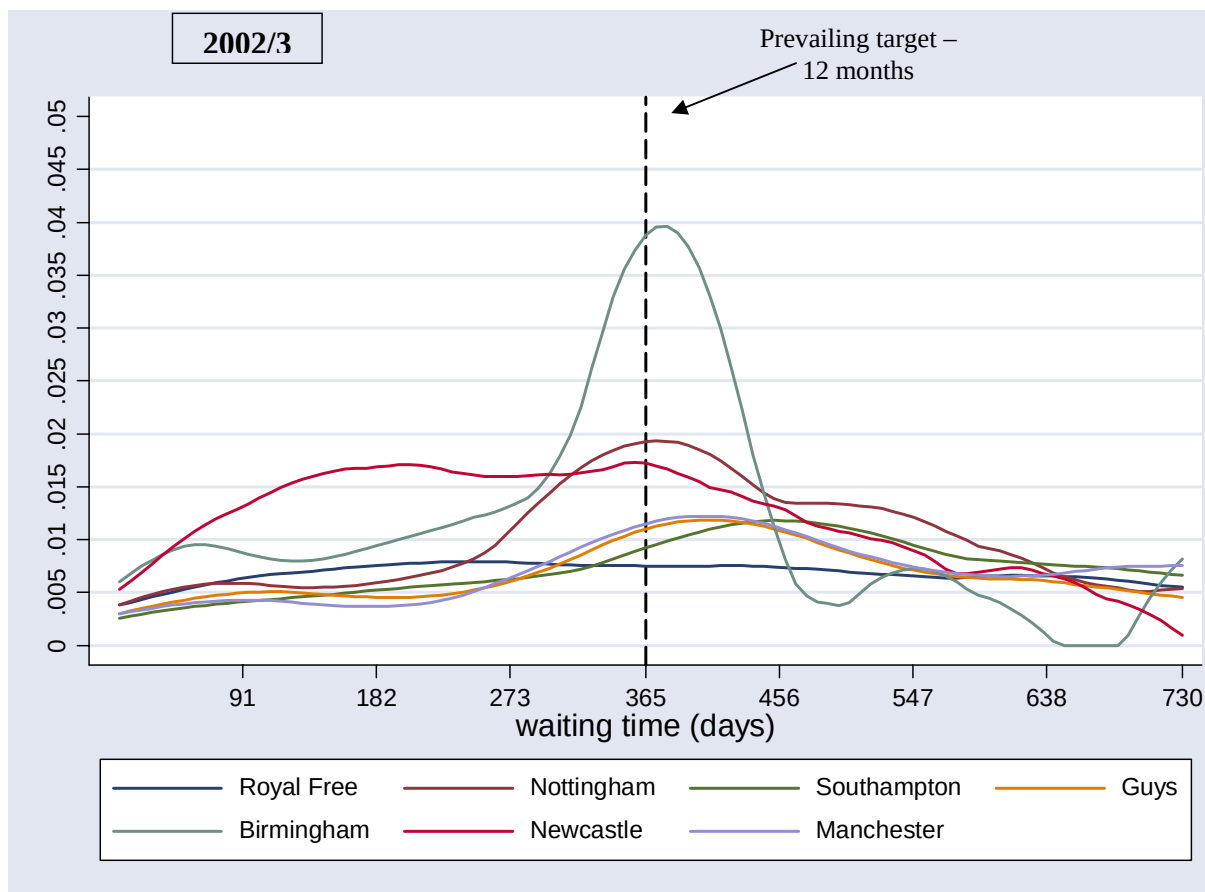
Seven Trusts were chosen for analysis at the provider level:

- The Royal Free, Hampstead
- Queen's Medical Centre, Nottingham University Hospital
- Southampton University Hospitals
- Guy's & St.Thomas
- Birmingham Heartlands & Solihull
- The Newcastle Upon Tyne Hospitals
- Central Manchester and Manchester Children's University Hospitals

We chose these providers because first they represent different geographical entities from all over England and secondly due to the different pattern of admissions they exhibit. According to the geographical area they belong to, trusts are part of distinct Health Authorities and broader Regional Offices. Manchester University Hospital belongs to North West Regional Office, Nottingham University Hospital to Trent Regional Office, Southampton Hospital to South East Regional Office, Newcastle Upon Tyne Hospitals to Northern and Yorkshire Regional Office, Birmingham Heartlands & Solihull to West Midlands and finally the Royal Free and Guy's & St.Thomas Hospitals to North Central and South East London respectively.

Figure 5 shows hazard curves for these providers for both years. The patterns and peaks differ greatly between providers. Some do not have notable peaks; for these providers, the probability of admission did not vary much over waiting time, for example the hazard curve for the Royal Free Hospital is almost horizontal in both years. All of the peaks, for those providers that have them, change over time, although the extent of this differs. For example Newcastle Upon Tyne Hospitals has a main peak of between 6 and 12 months for both years, with only a small difference between them. Other providers' changes were much larger, for example, the main peak for Birmingham Heartlands & Solihull reduced from a little more than 15 months in 2001/2 to 12 months in 2002/3.

Figure 5: Hazard rates for seven Trusts in 2001/2 and 2002/3



Identifying variations in probability of admission according to admission source

Patients can be admitted to hospital from a variety of sources. Three major routes are waiting lists, where there is no exact date of admission, booked admissions, where there is an exact date for admission and planned, where there is an exact date of admission for a course of treatment over time or a second operation. Figure 6 shows survival curves for 2001/2 by admission method and Figure 7 shows corresponding hazard curves for both 2001/2 and 2002/3.

Booked admission patients are usually admitted fairly quickly, and survival curves show that the proportion of patients waiting for booked admission decreases more quickly over time than for planned or waiting list patients. Below 450 days, survival on the waiting list for planned admissions is below that for waiting list admissions; after that the reverse occurs. The main peaks in the hazard curves for booked admissions are 3 months for both years and also 15 months for 2001/2 and 12 months for 2002/3. The patterns for the hazard rates do not change for planned admissions over the two years, but the main peak for waiting list admissions reduces from around 15 months in 2001/2 to between 12-15 months in 2002/3.

Figure 6: Survival curves by type of admission: 2001/2

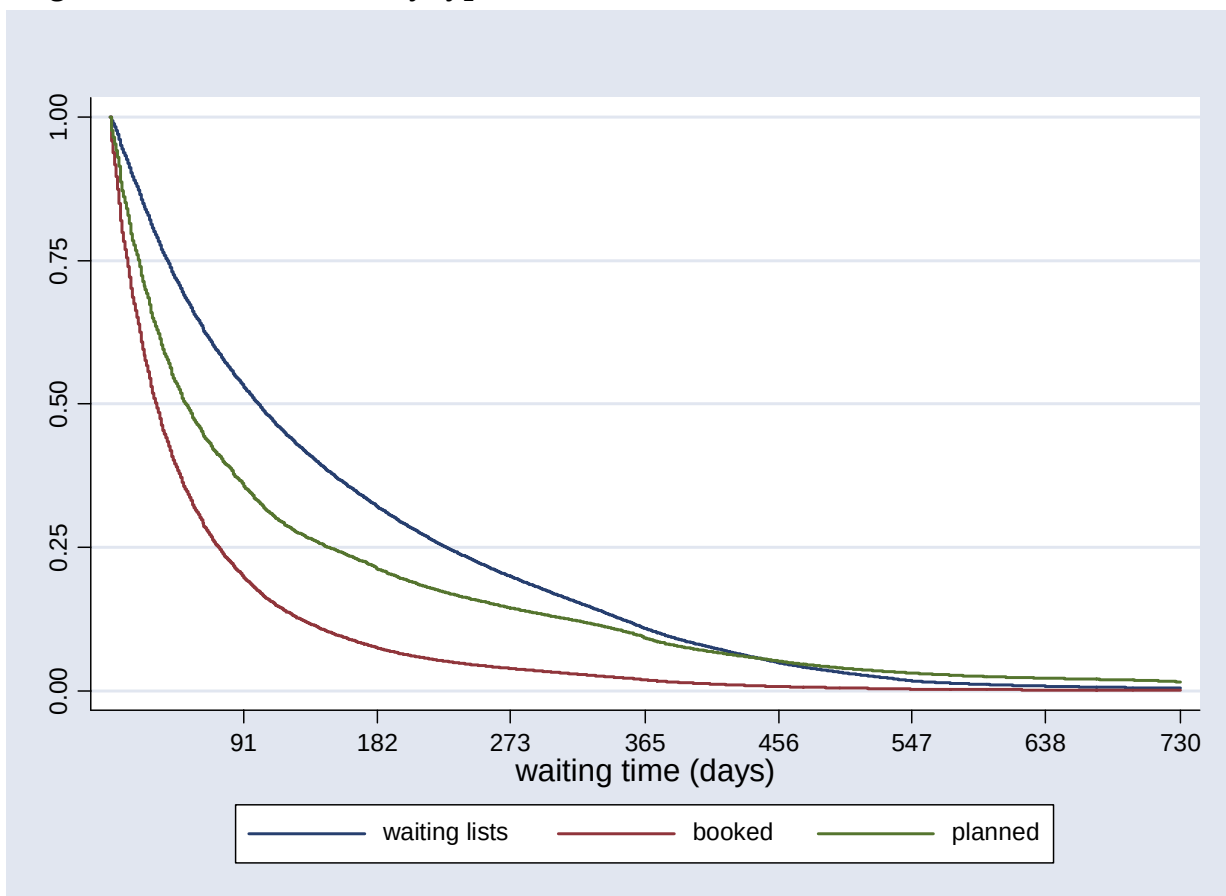
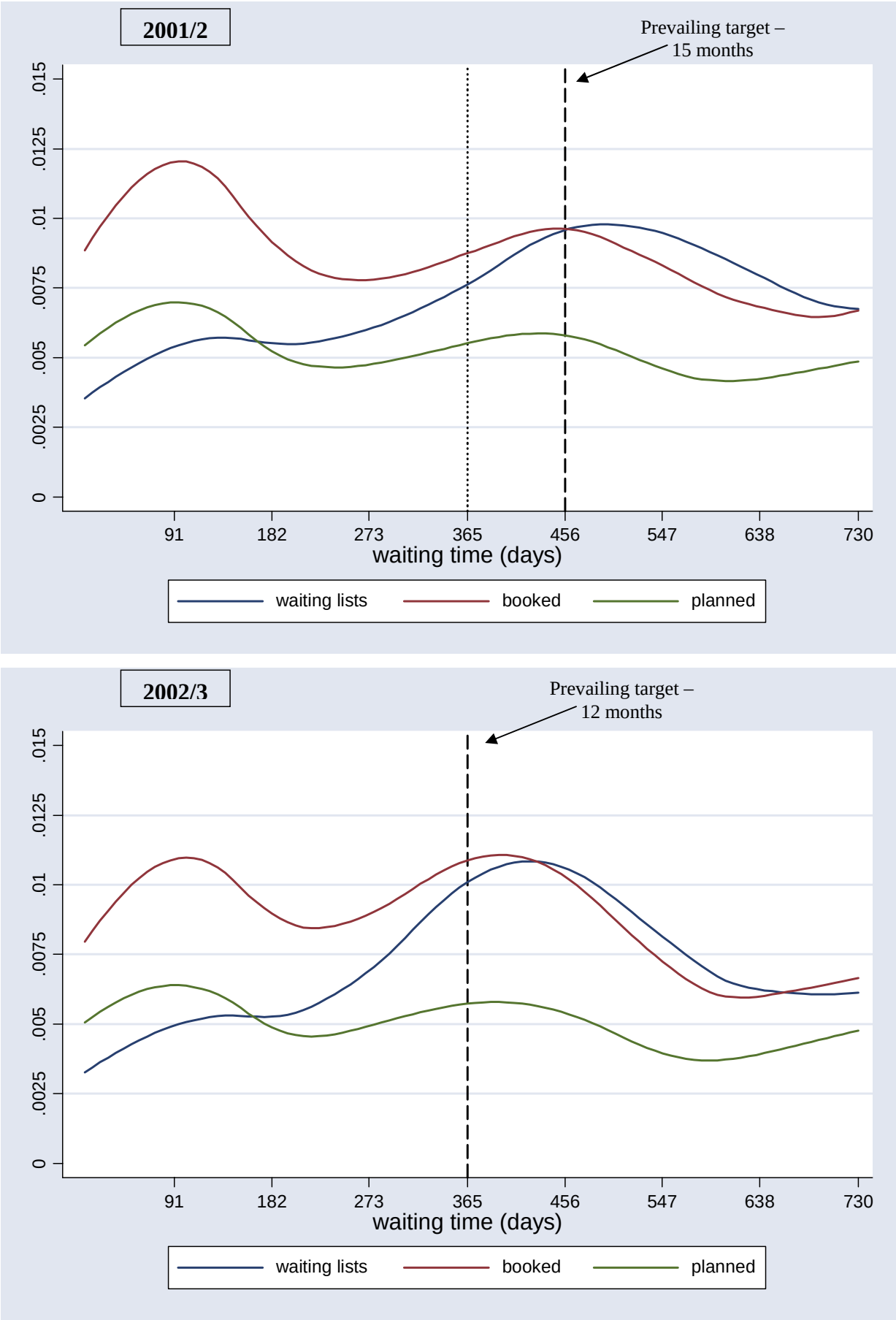


Figure 7: Hazard rates by type of admission



Applying time-to-event analysis at a less aggregated level

The analysis undertaken so far has clearly established that patterns of admissions for elective surgery varies between different specialties, operative procedures, HRGs, particular providers and different admission methods. Yet, all previous graphs exploited data at an aggregated level; the first part of the paper concentrated on the differences between specialties as a whole, that is in all NHS trusts of England during financial years 2001/2 and 2002/3, the second part revealed waiting times variations according to type of operation undertaken in all UK hospitals, the third part explored differences between providers with respect to the entire set of waiting lists of each provider treating the different heterogeneous lists as a homogeneous unique waiting list and finally the last part used all available data of the two financial years to compare different admission methods in terms of differences of the time period a patient has to wait for treatment.

This section of the paper will address the same questions using less aggregated data. It aims to expose behavioural responses to targets by managers and clinicians and their effect on waiting times distributions. Evidence for possible clinical distortions will be also examined. In order to demonstrate the potential of time-to-event analysis at exploiting less aggregated data we show the successive levels of analysis – aggregate towards less aggregate – using the seven trusts mentioned previously. The cases of Birmingham Heartlands & Solihull and Royal Free, Hampstead, which as illustrated in figure 5 behave quite differently, are analysed in more detail in the main text. Selected survival and hazard curves of the rest trusts are presented at appendix I.

Figures 8 and 9 show the survival and hazard curves of overall waiting times for elective surgery for Birmingham Heartlands & Solihull. For 2001/2, at time 0, all patients are on the list, at around 2 months (57 days) 50% of them have been moved from the list to be treated and at around 4.5 months (143 days) the proportion increases to 75%. The same pattern characterises the waiting time distribution of elective patients for 2002/3. Hazard curves reveal notable peaks at a little more than 15 months for 2001/2 and 12 months for 2002/3. It is obvious that this trust attempts to adjust to national targets by changing the probability of moving patients from lists for admission. What is not obvious, though, is whether it maintains the same behaviour for all specialties and different operative procedures.

Figure 8: Survival curves of overall waiting times for Birmingham

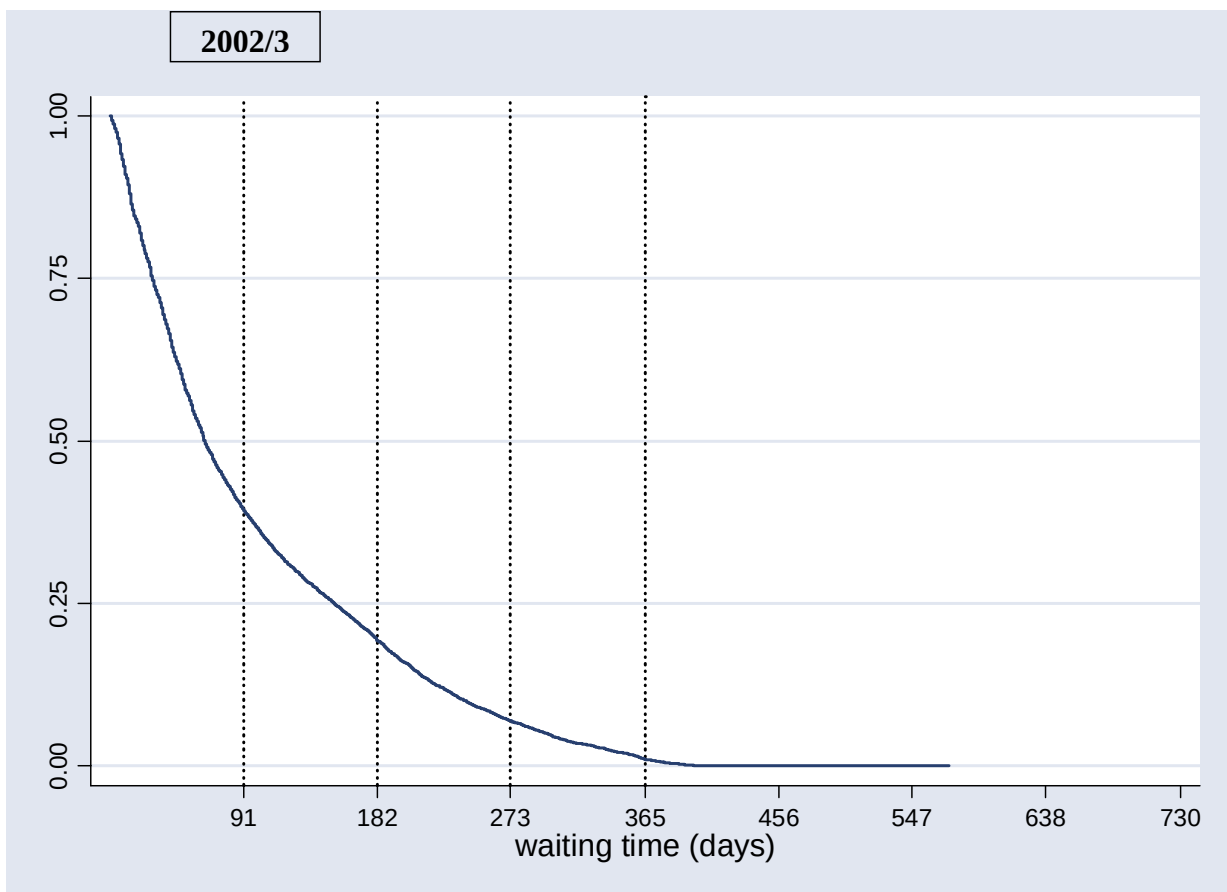
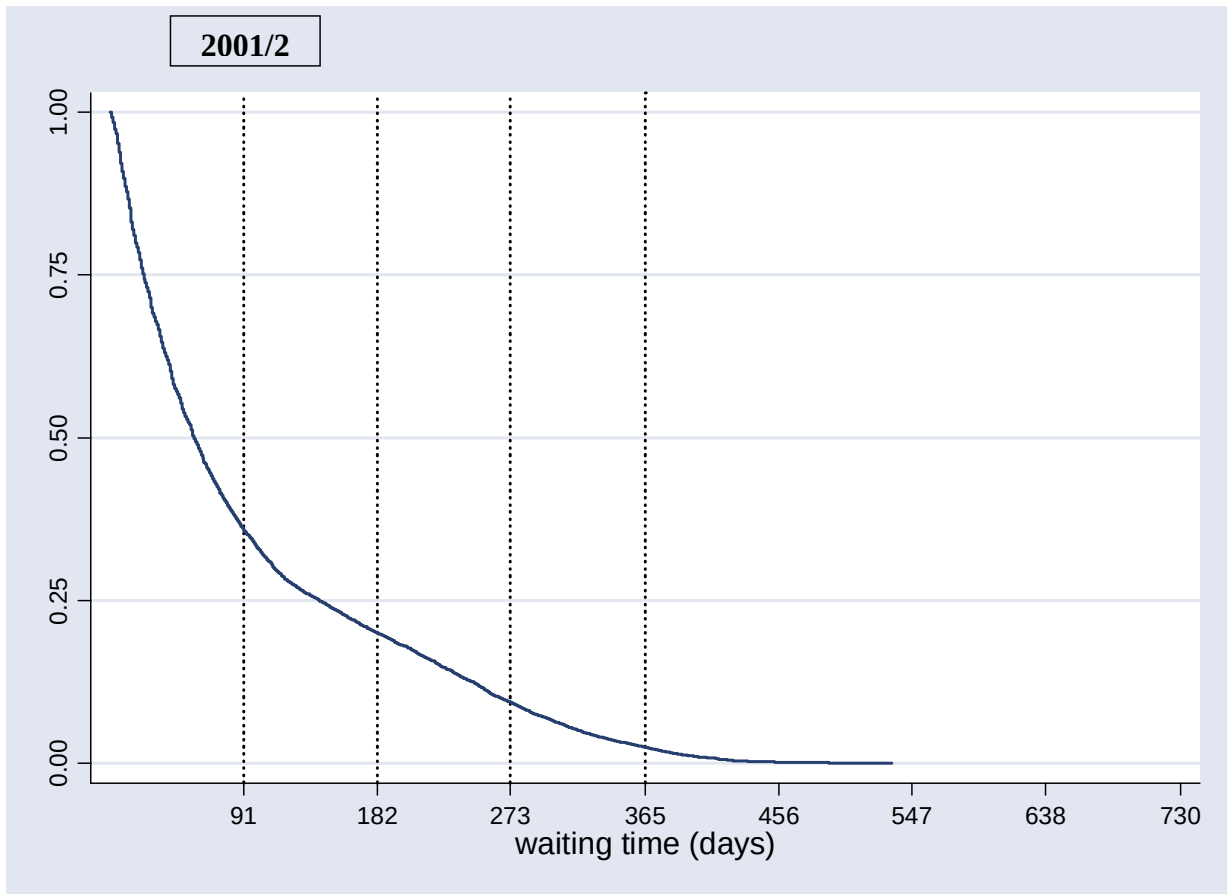
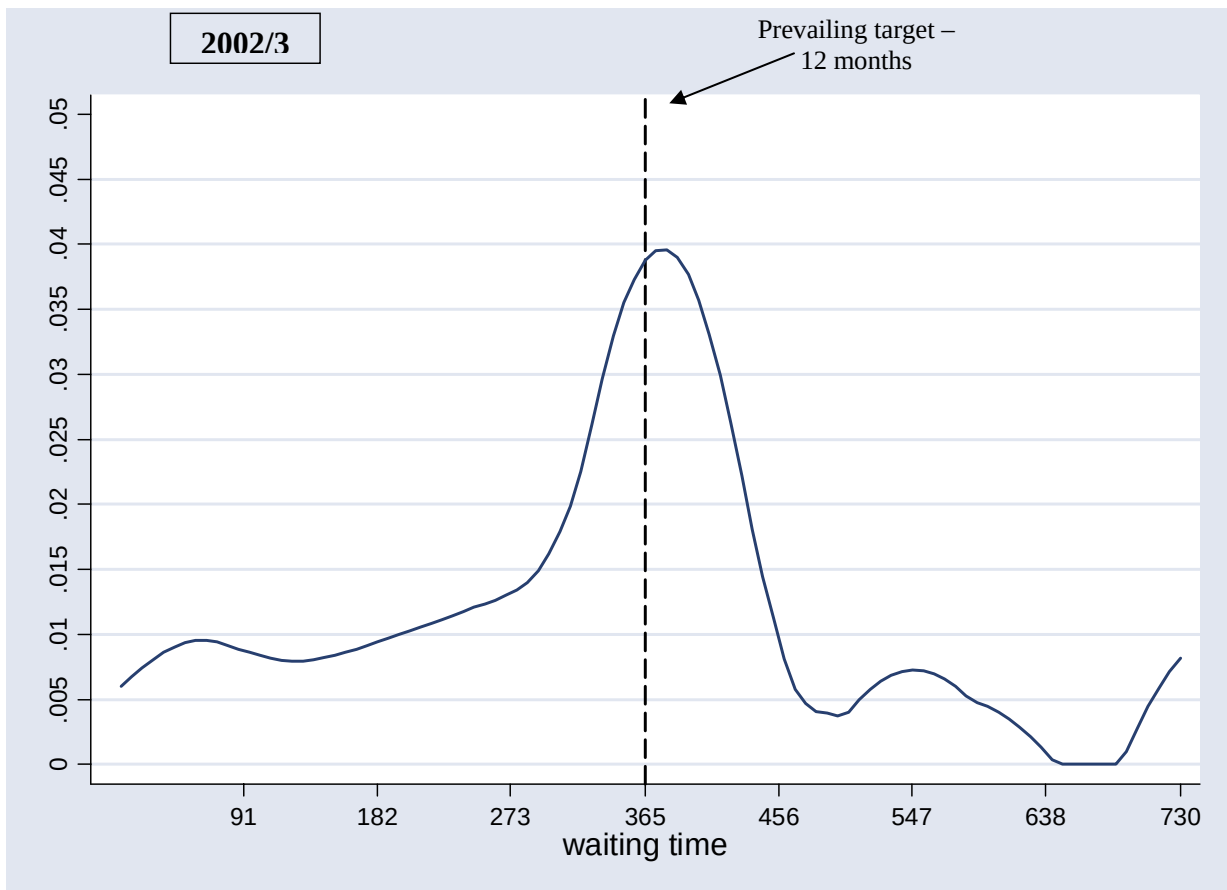
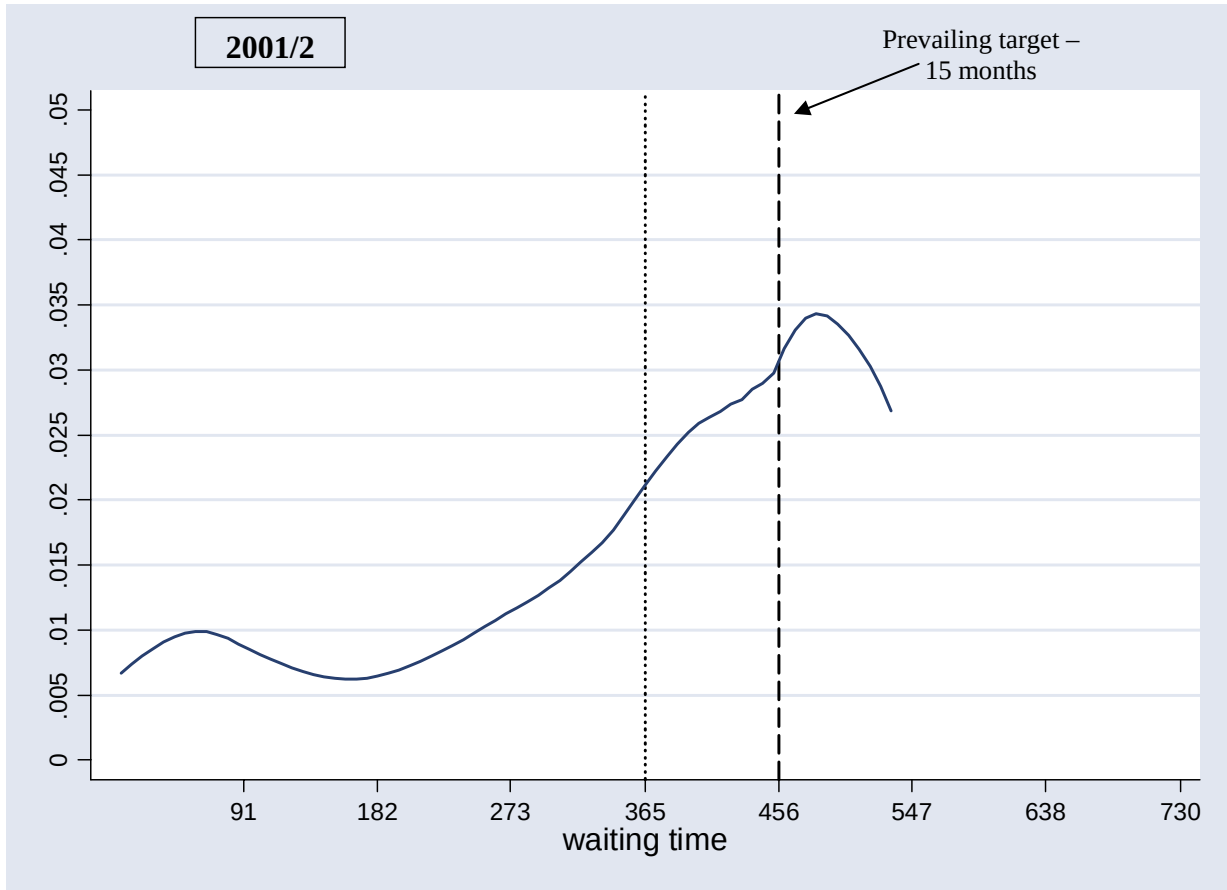


Figure 9: Hazard rates of overall waiting times for Birmingham



Analysis at the level of specialty and type of operation illustrates variability in trust responses (figures 10 and 11). Figure 10 demonstrates the estimated survival and hazard functions by different specialties for both financial years. In particular, patients waiting for general surgery tend to wait shorter periods than patients scheduled for orthopaedic or ophthalmologic procedures. According to 2001/2 survival curve, 50% of patients are moved off the general surgery list at around 1 month (34 days), the orthopaedic list at 4 months (111 days) and the ophthalmologic list at 6.5 months (194 days). At around 8 months the survival curves of the last two specialties intersect. Similar results are observed for the subsequent year, yet the difference between orthopaedic and ophthalmologic specialties diminishes.

In addition, the probability of admission does not remain constant and exhibits different patterns for each specialty. For general surgery, increased waiting list activity is observed as peaks in the curve for people waiting 3, 8 and 11 months for 2001/2 and 2, 6, and 12 months for 2002/3; for orthopaedic surgeries, between 12 to 15 months for 2001/2 and at 12 months for 2002/3; for ophthalmology around 15 months for 2001/2 and a little less 12 months for 2002/3. Two insights can be drawn from this figure. Firstly, it is clear that the trust does not adopt the same behaviour in managing different surgical lists; general surgery waiting lists follow a very different pattern than the other two specialties. Secondly, it shows the effects of waiting times targets; the probability of admission for those whose wait approaches a target increases and falls when their wait exceeds the target.

Figure 11 shows the estimated survival and hazard functions by different operative procedures for Birmingham Heartlands & Solihull during 2001/2 and 2002/3. The shorter waiting times are for cholecystectomy, inguinal hernia and varicose vein procedures and the longest for lens prosthesis and hip replacement. Furthermore, it is worth emphasising that the three general surgical waiting lists do not consist of patients waiting more than 1 year, while lens prosthesis and hip replacement do. The hazard curve for 2001/2 reveals the following patterns: for cholecystectomy, peaks are at 2 and 6 months, for inguinal hernia at 2 and 8 months, for varicose vein at 2 and 7 months, for lens prosthesis at almost 15 months and for hip replacement at between 12 and 15 months. The hazard curve for 2002/3 demonstrates that as the targets become tougher, the peaks change towards the lower waiting times. A peak at exactly 12 months, which is the target of that year, is observed for hip replacements and a little less than 12 months characterises lens prosthesis.

Figure 10a: Survival curves for three specialties in Birmingham

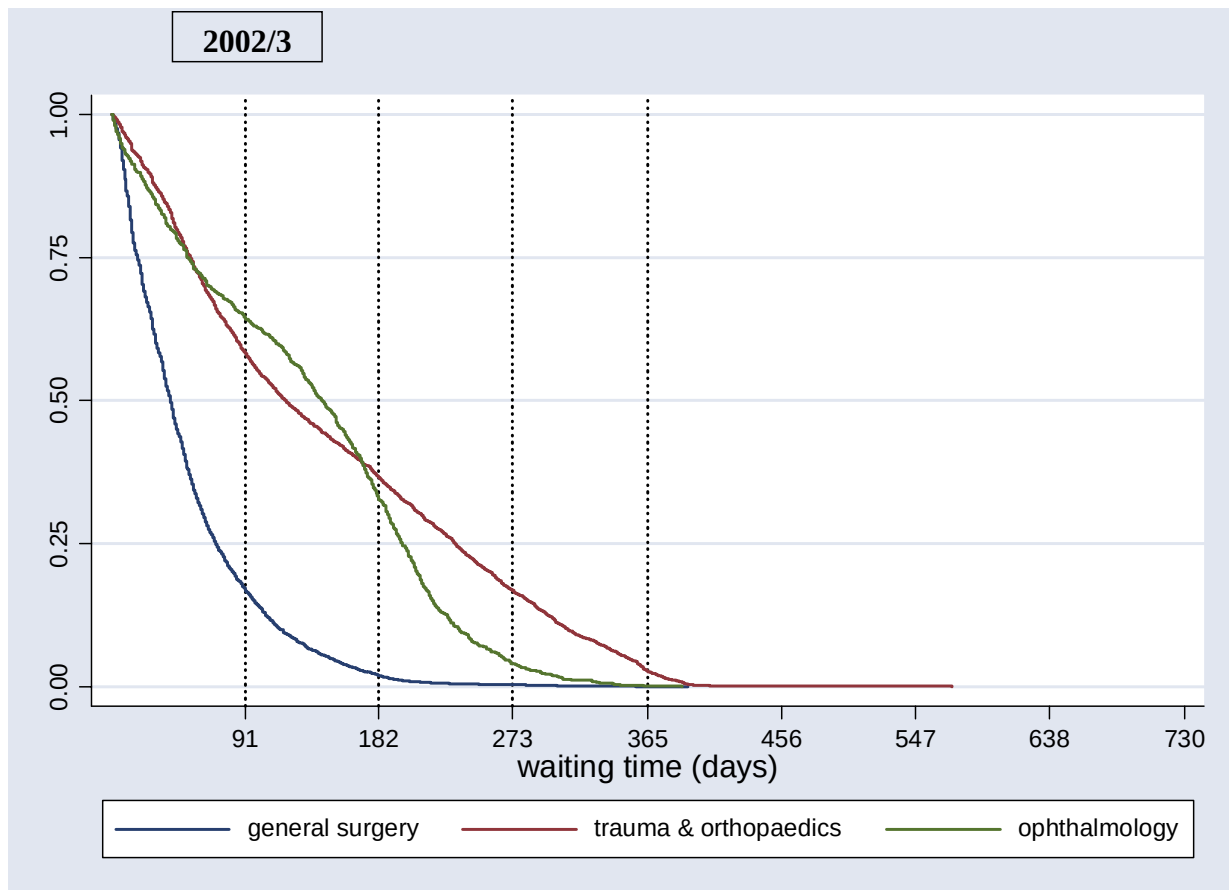
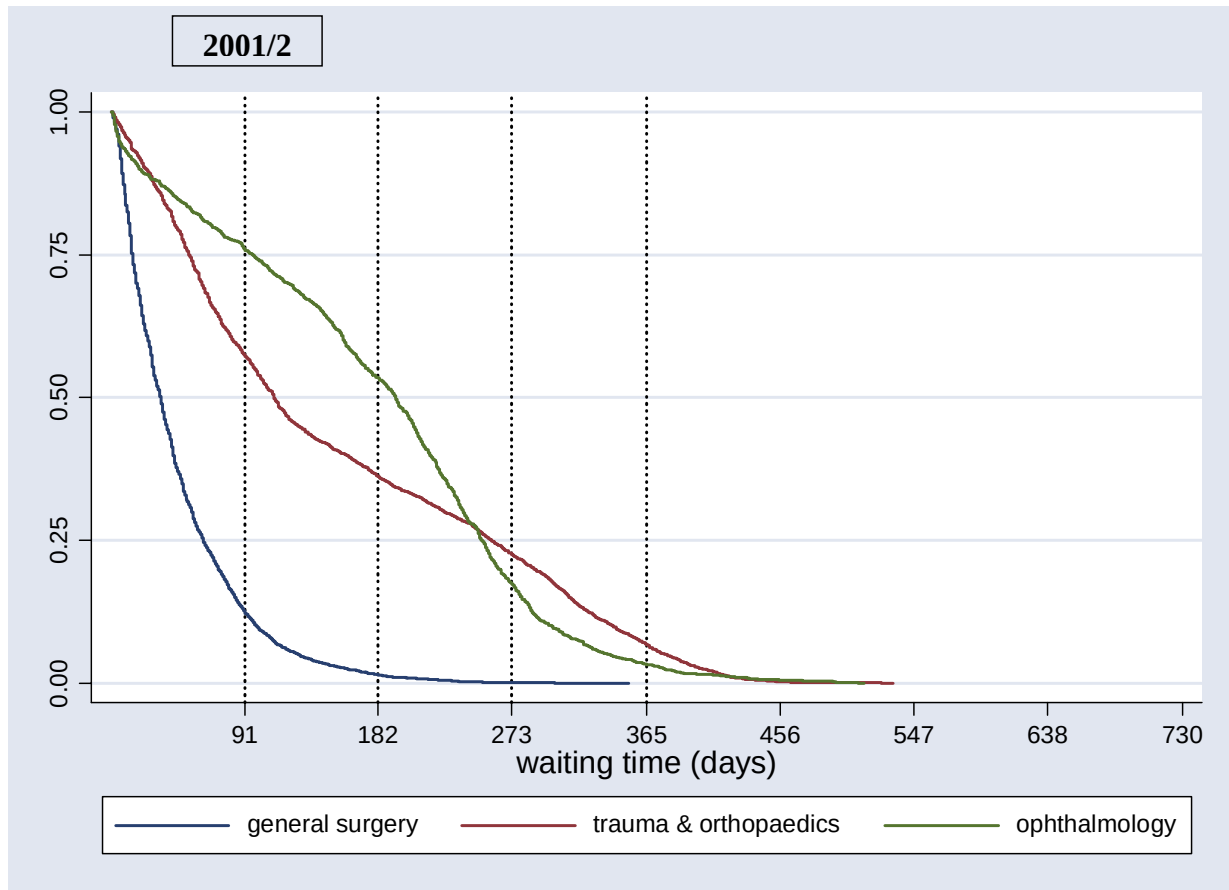


Figure 10b: Hazard rates for three specialties in Birmingham

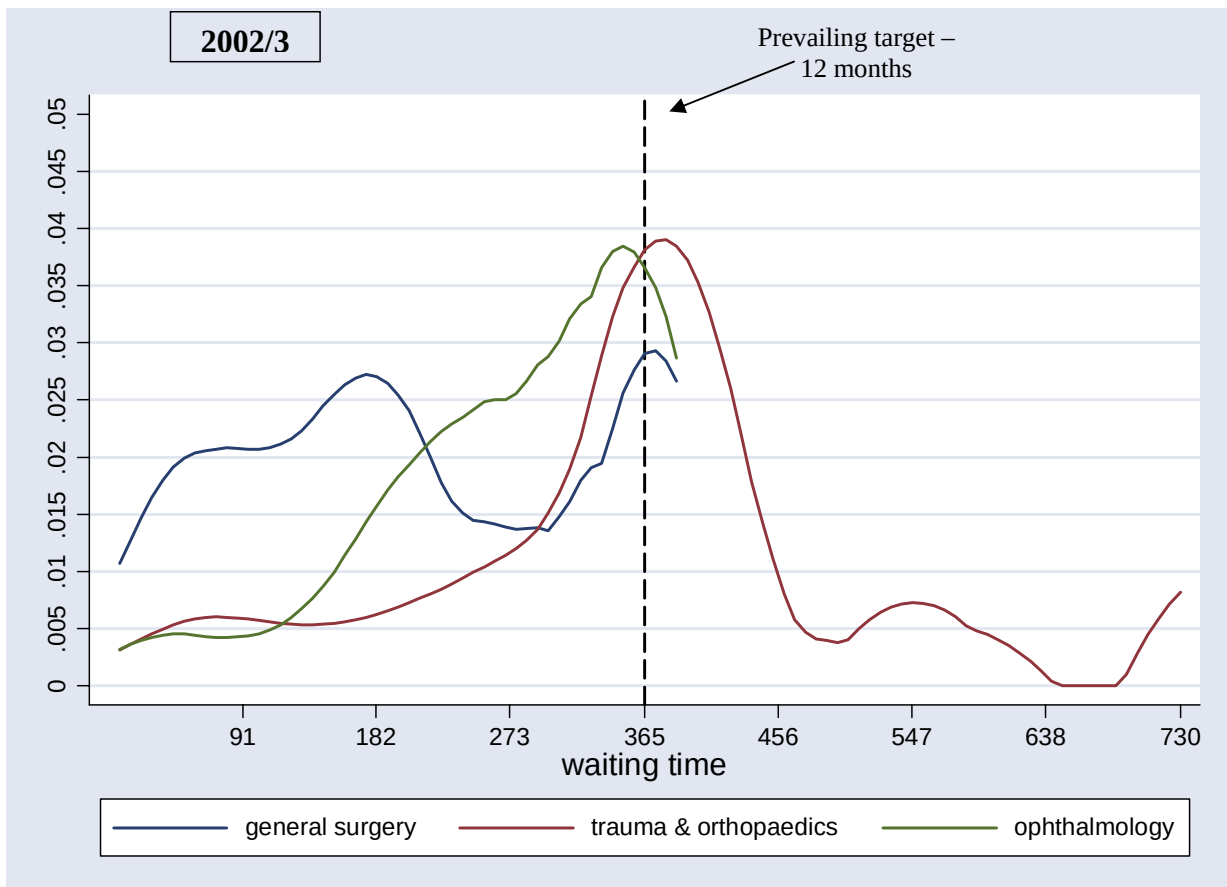
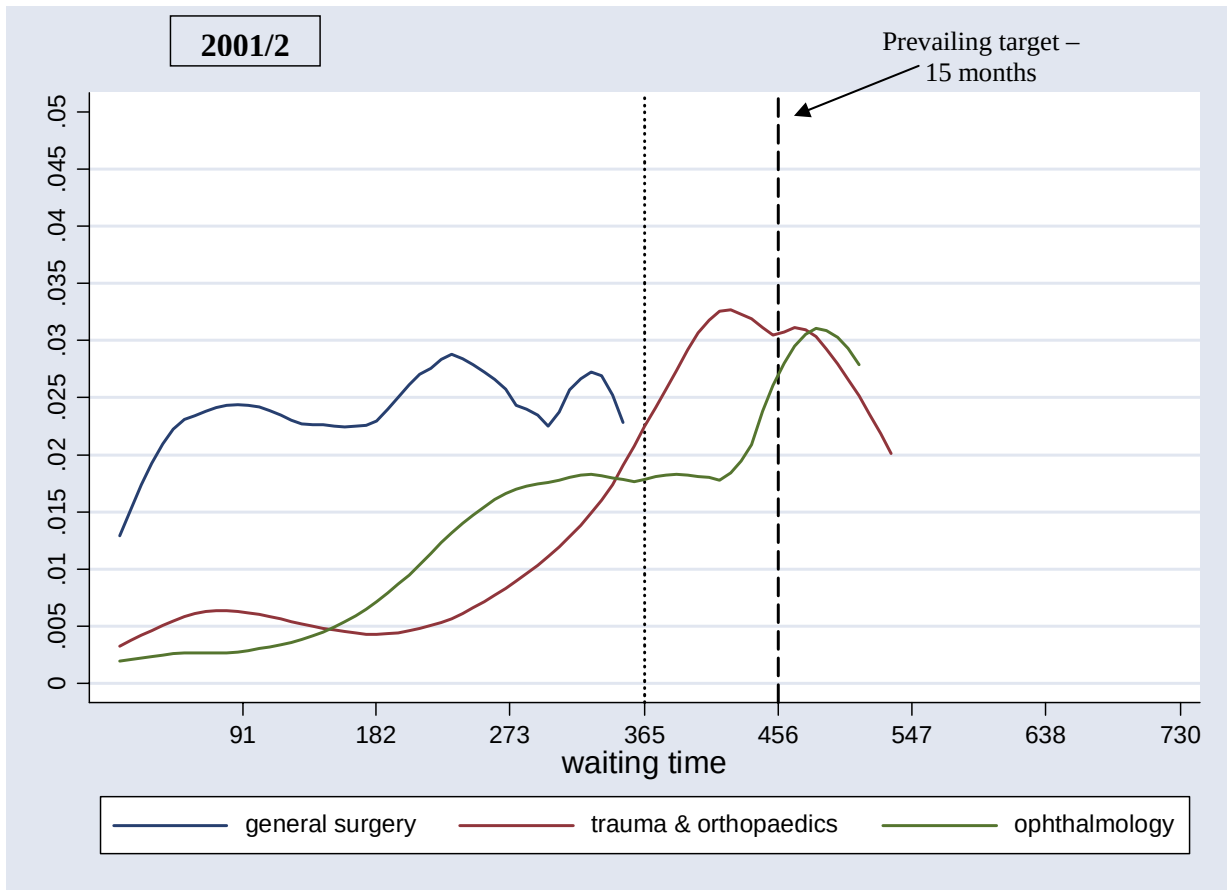


Figure 11a: Survival curves for five operations in Birmingham

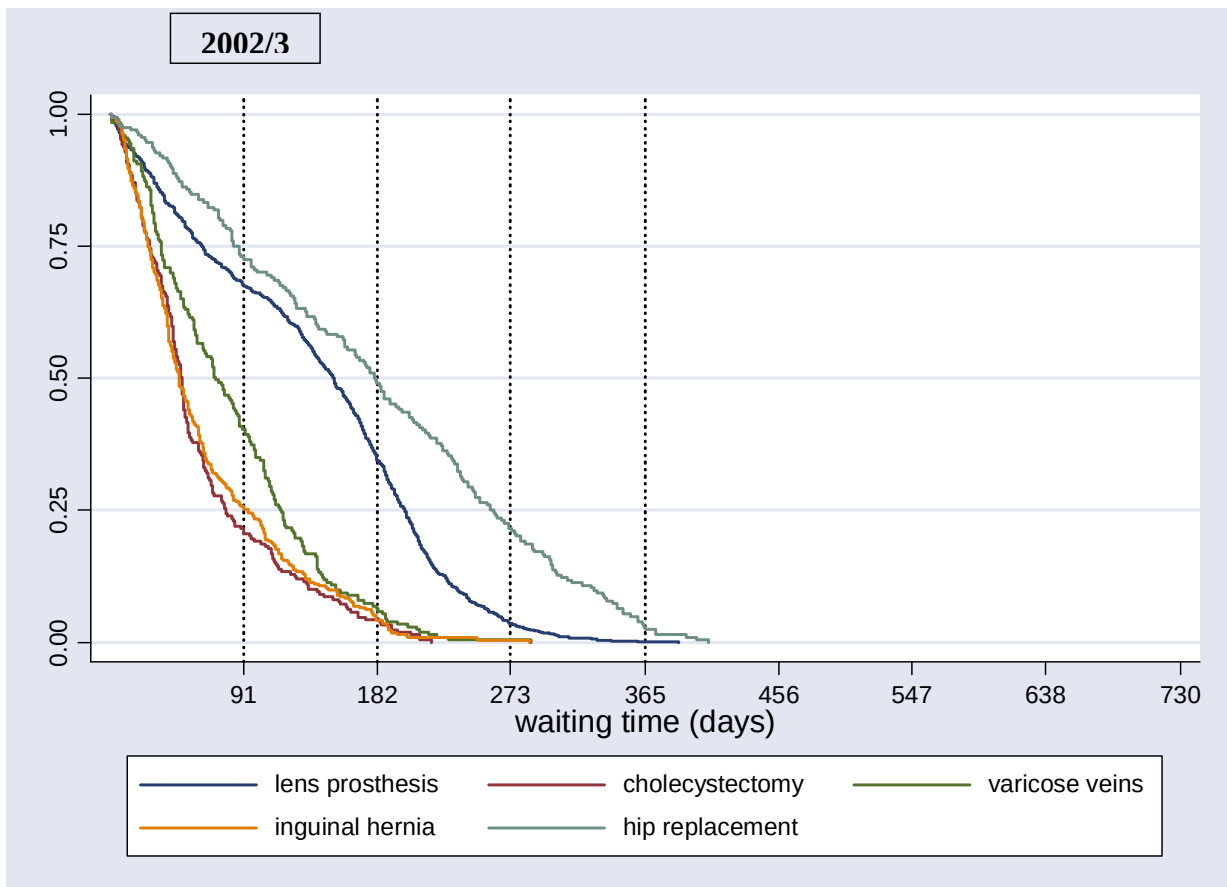
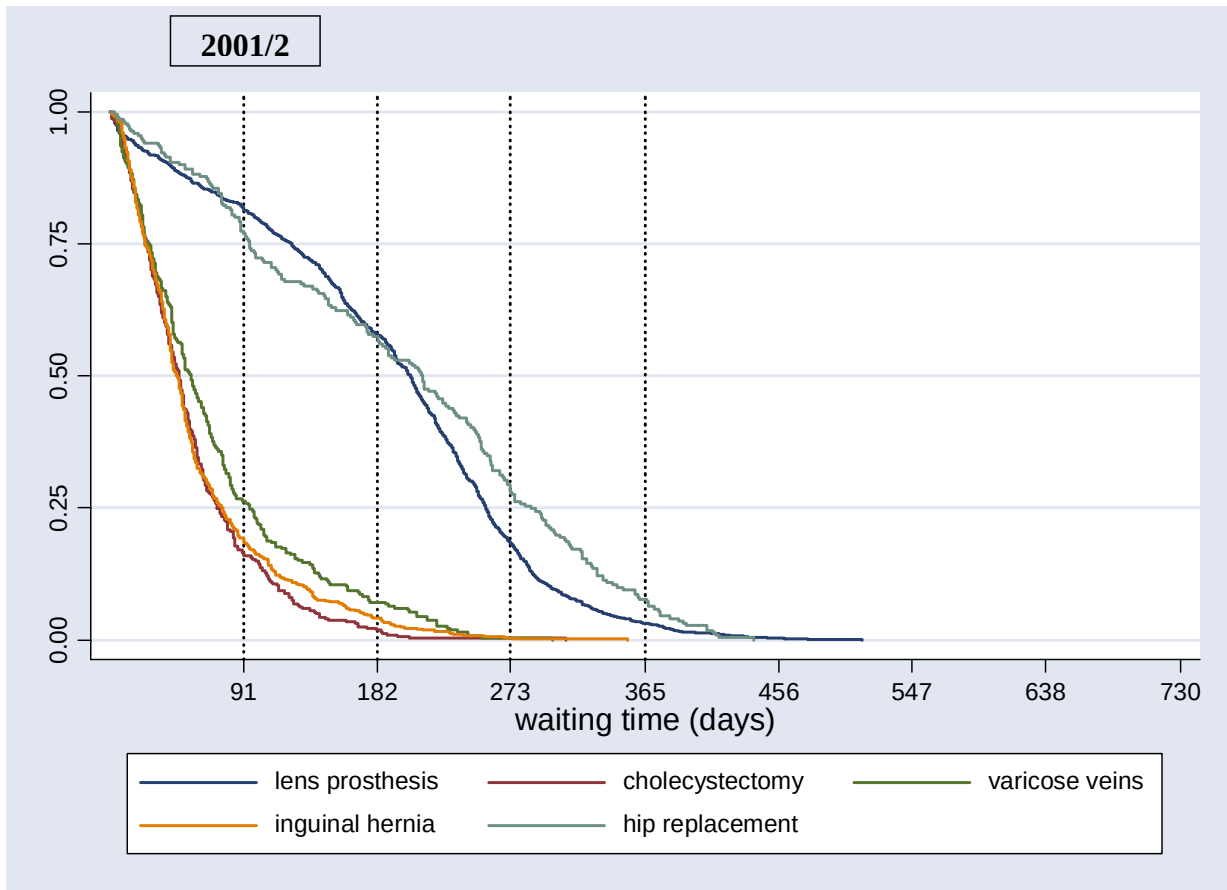
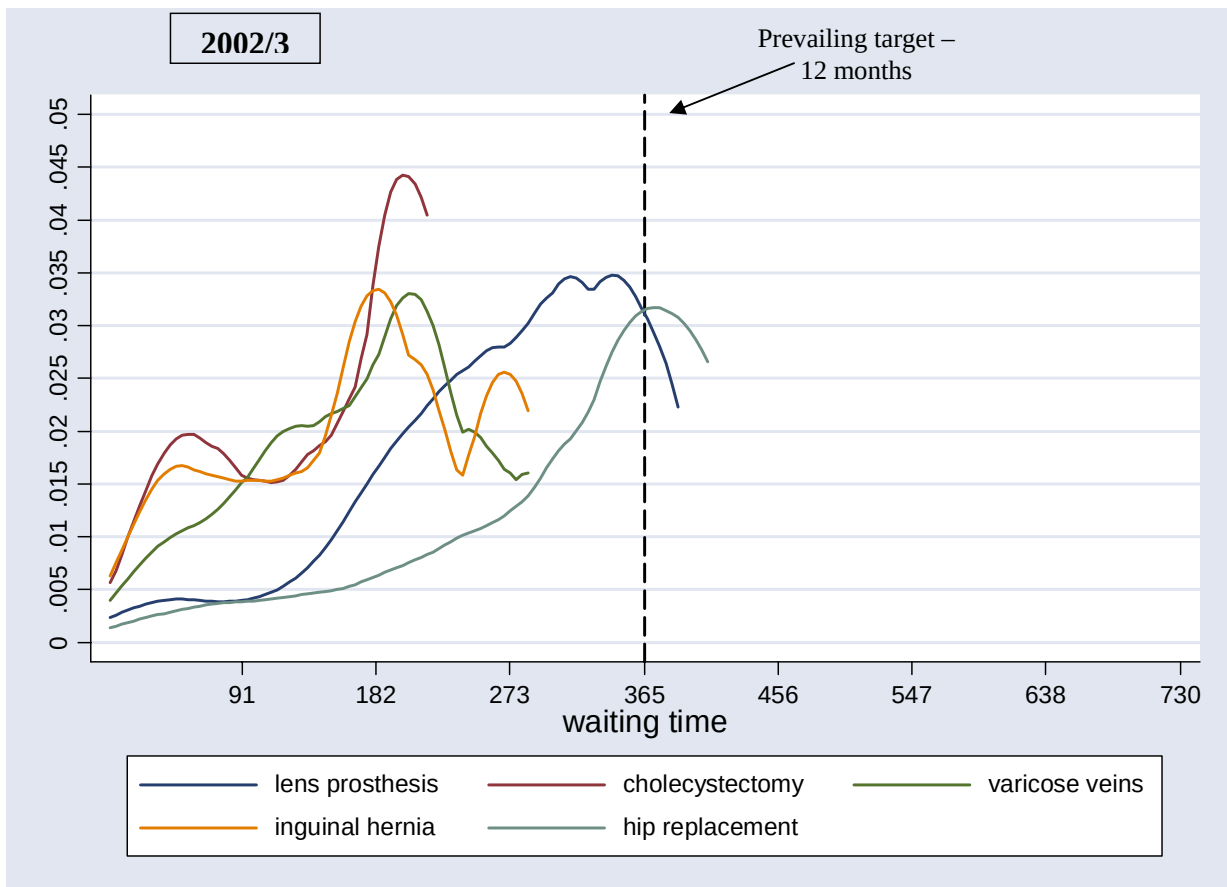
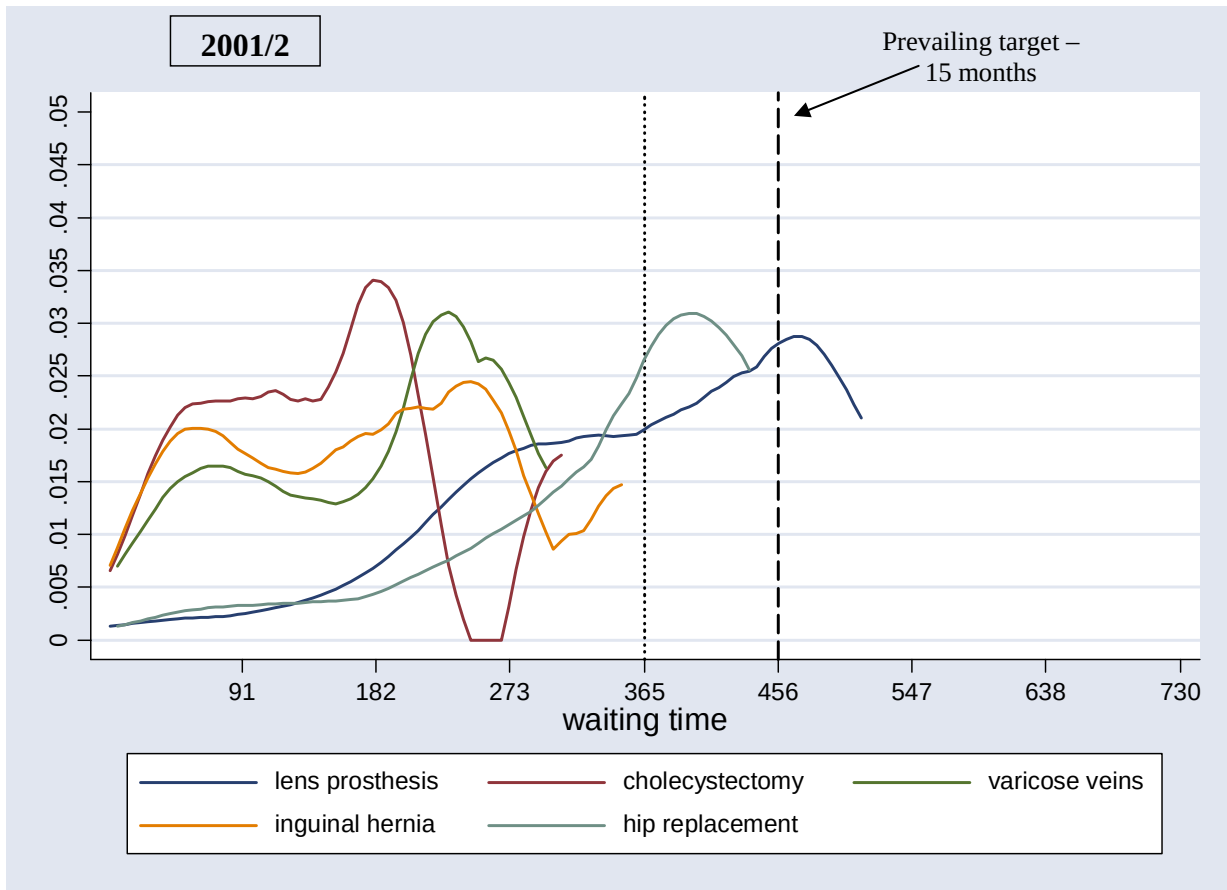


Figure 11b: Hazard rates for five operations in Birmingham



Birmingham Heartlands & Solihull exhibits great management activity to tackle excessive waiting times for its patients. Quite different behavioural responses are observed by the Royal Free Hampstead trust in London. Analysis of the overall waiting times of the trust reveals that Royal Free patients have to wait a little longer for treatment as the rate of admission over time changes slower (figure 12). But, more importantly, the hazard rates do not change substantially, for 2001/2 there is only a very small peak at around 17 months while for the following year there is no peak at all. The probability of admission for elective surgery is constant, independent of the time patients spent on waiting lists.

According to official returns of waiting lists for elective surgery (Department of Health Hospital Waiting Times/List Statistics) Birmingham Heartlands & Solihull had achieved the national waiting time targets for 2001/02, while Royal Free had not. One limitation of the official waiting list statistics of that year was that broad time intervals of waiting were used (eg. patients waiting for 12-17 months). Thus, we cannot calculate exactly the number of people waiting more than 15 months. However, none of their patients had to wait more than 12 months at the end of 2003, thus they both achieved the waiting time target for the next year. Conversely, based on NHS performance ratings, both trusts had no inpatients waiting longer than the standards for both years tested (Department of Health, *NHS Performance Ratings 2001/02, 2002/03*).

Yet, less aggregated analysis at the level of three specialties suggests that probability of admission is constant only for general surgeries for 2001/2 and orthopaedic procedures for 2002/3 (figure 13). The other specialties follow different patterns with the presence of specific peaks. At the less aggregated level of waiting lists for different operations, the image is much clearer (figure 14). For 2001/2, varicose vein procedures have constant hazard rates, cholecystectomy is characterized by a very wide peak at 15 months, the hip replacement hazard curve is positively sloped but with a very small slope; on the contrary, lens prosthesis exhibits increased admission activity for people waiting 15 months and inguinal hernia for 18 months. For 2002/3, with the exception of hip replacement and inguinal hernia (peaks at around 15 months) the hazard rates of all other procedures are represented by almost flat lines.

Figure 12a: Survival curves of overall waiting times for Royal Free

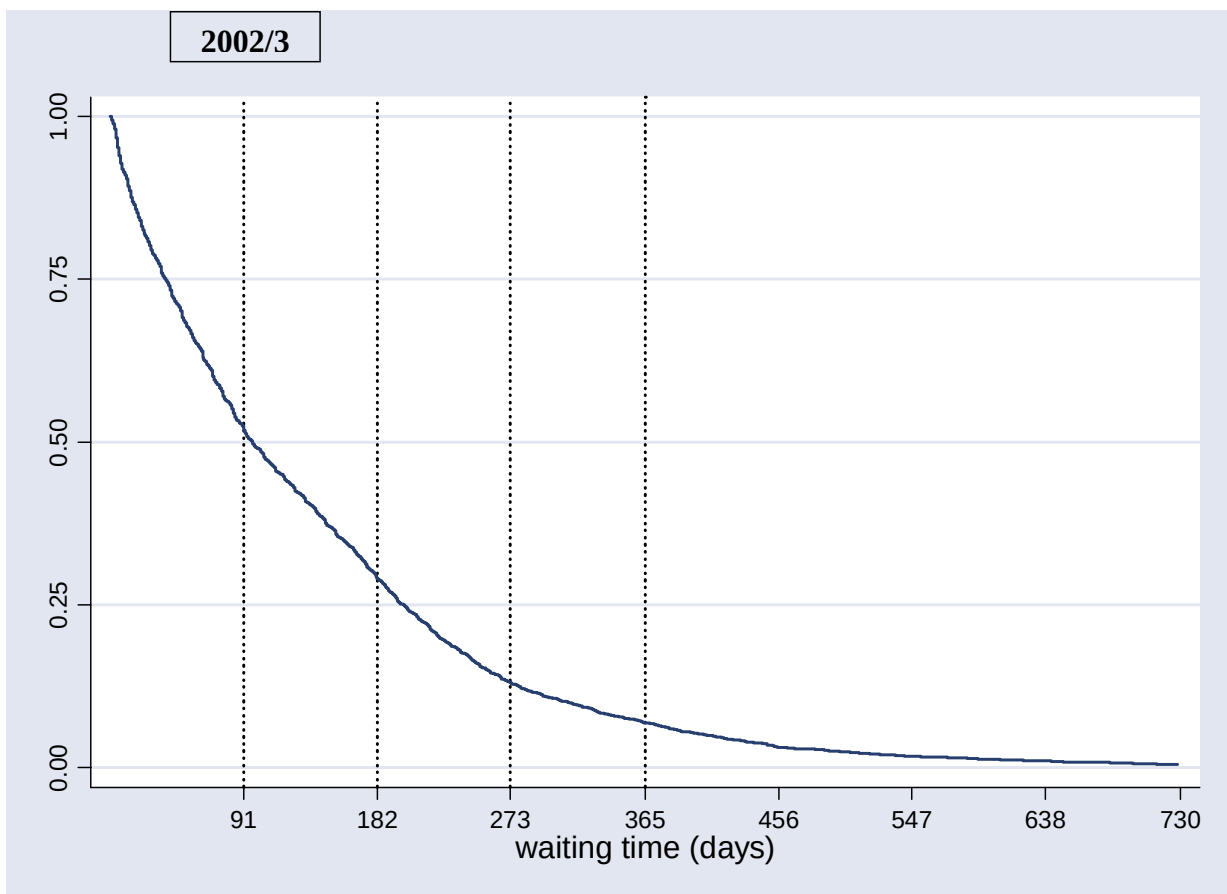
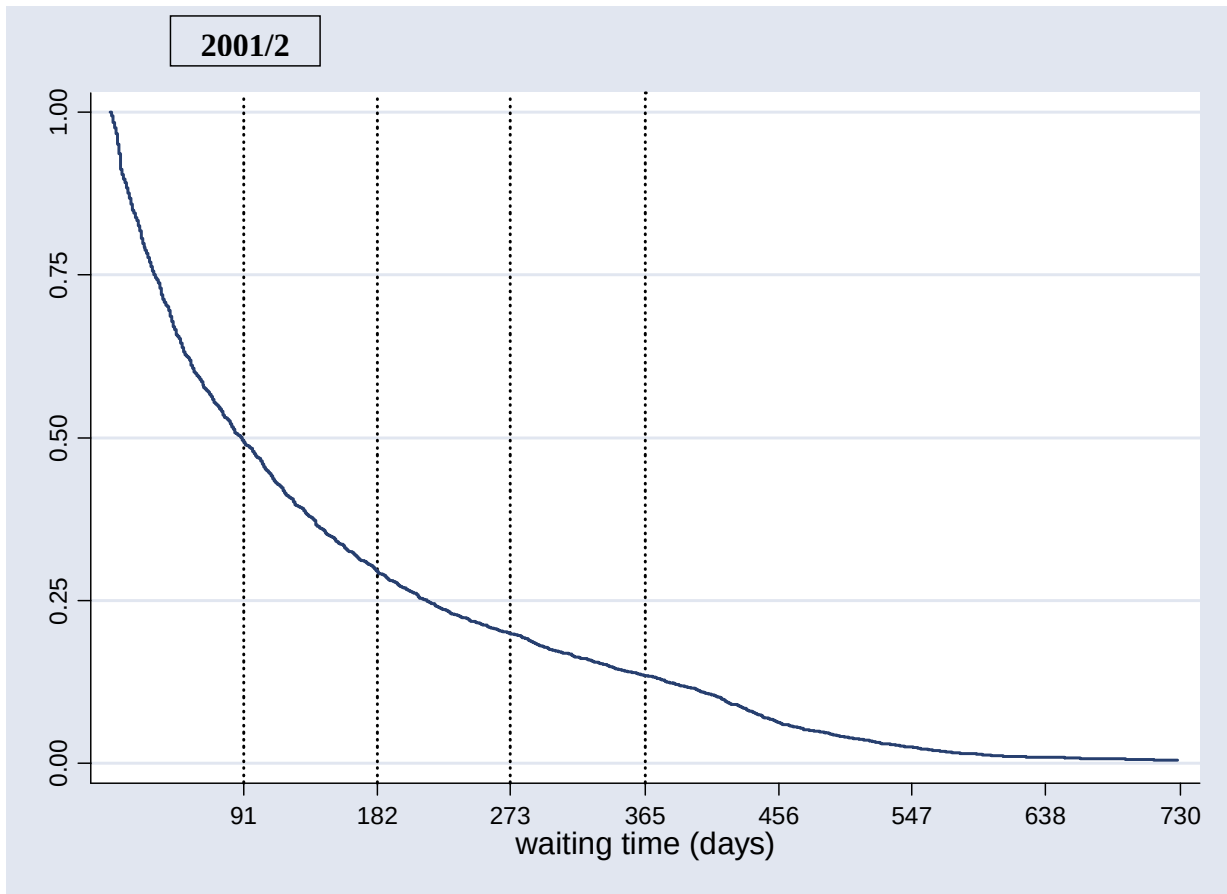


Figure 12b: Hazard rates of overall waiting times for Royal Free

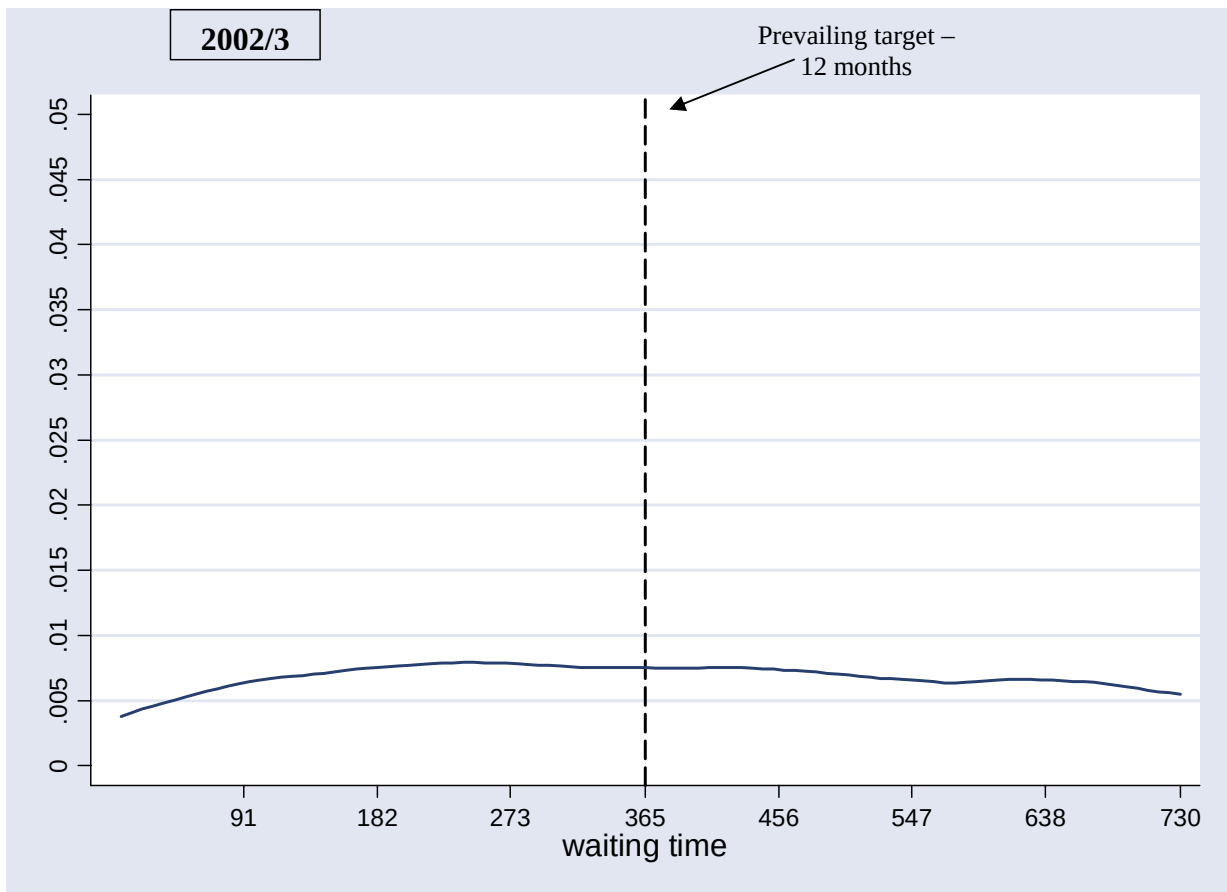
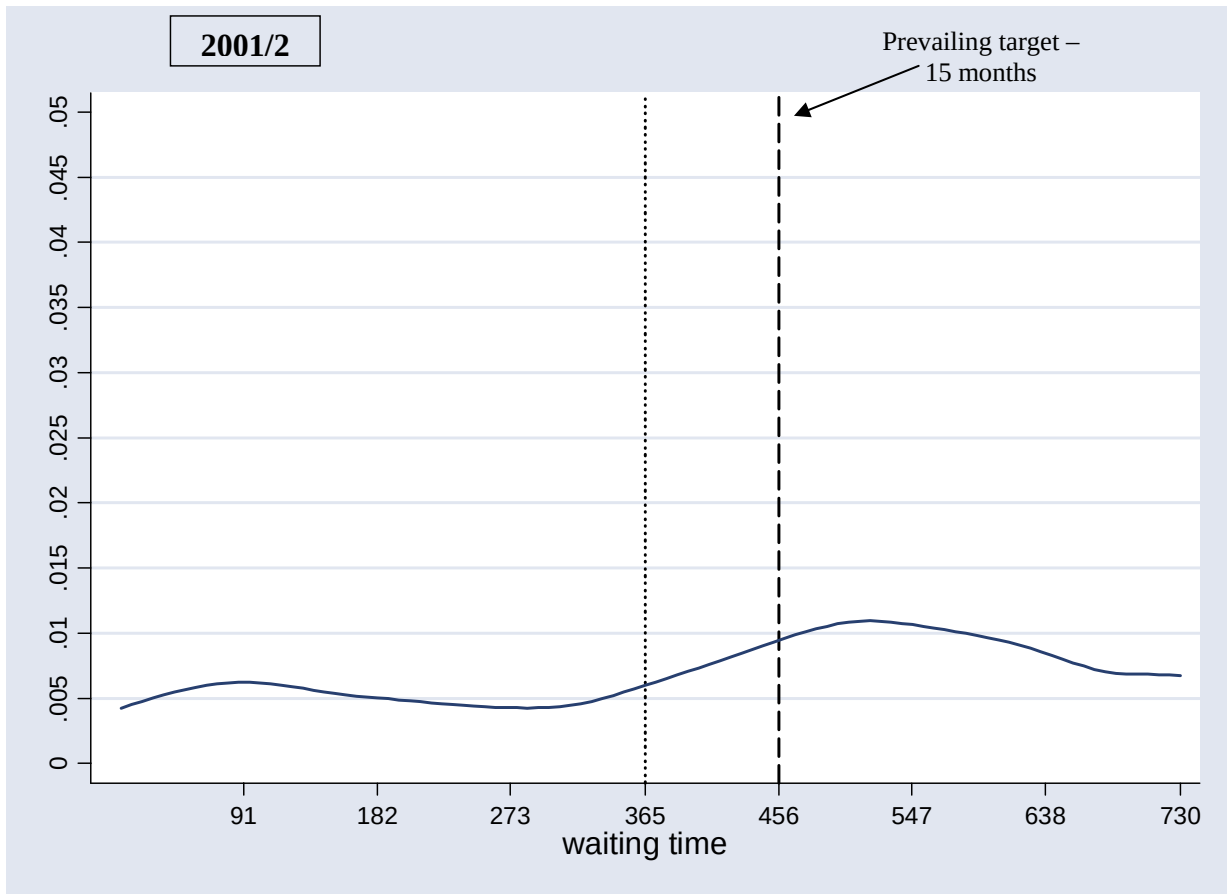


Figure 13a: Survival curves for three specialties in Royal Free

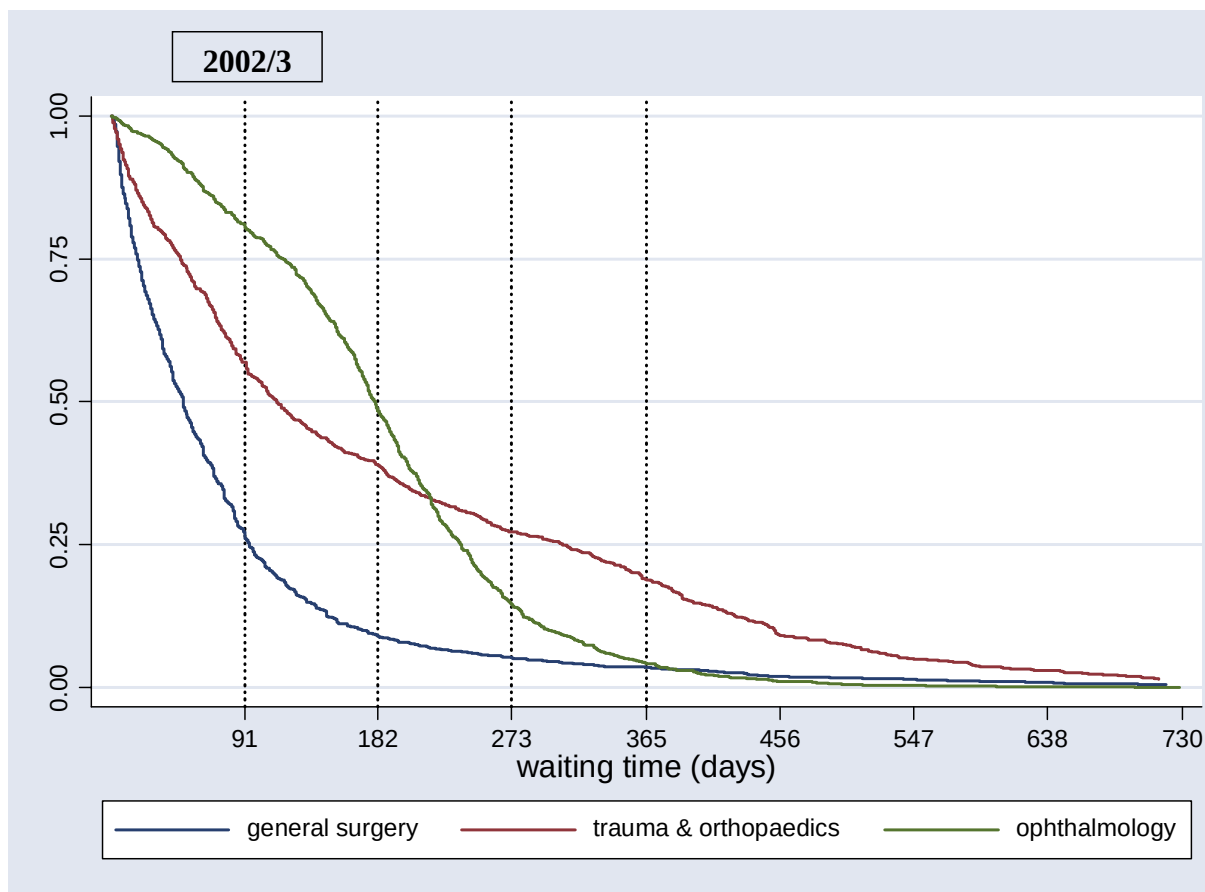
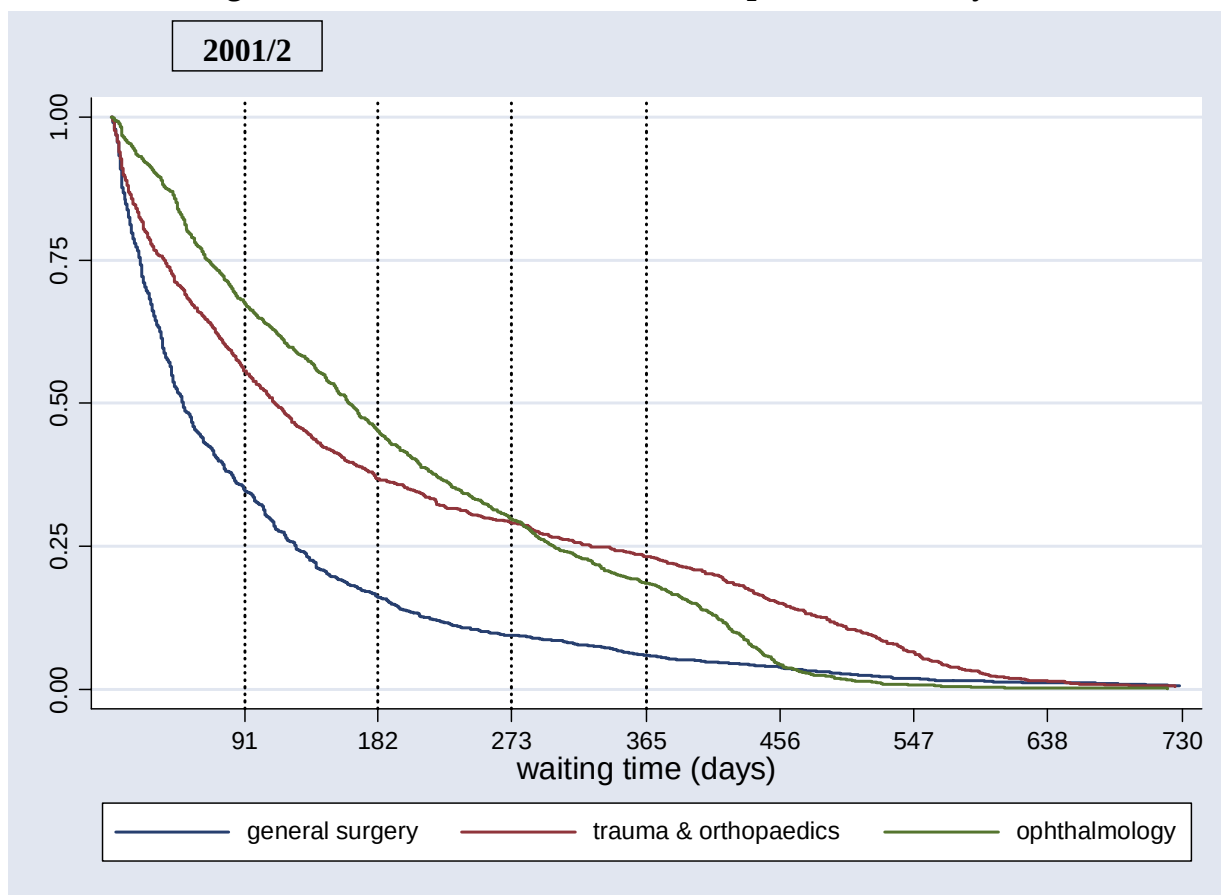


Figure 13b: Hazard rates for three specialties in Royal Free

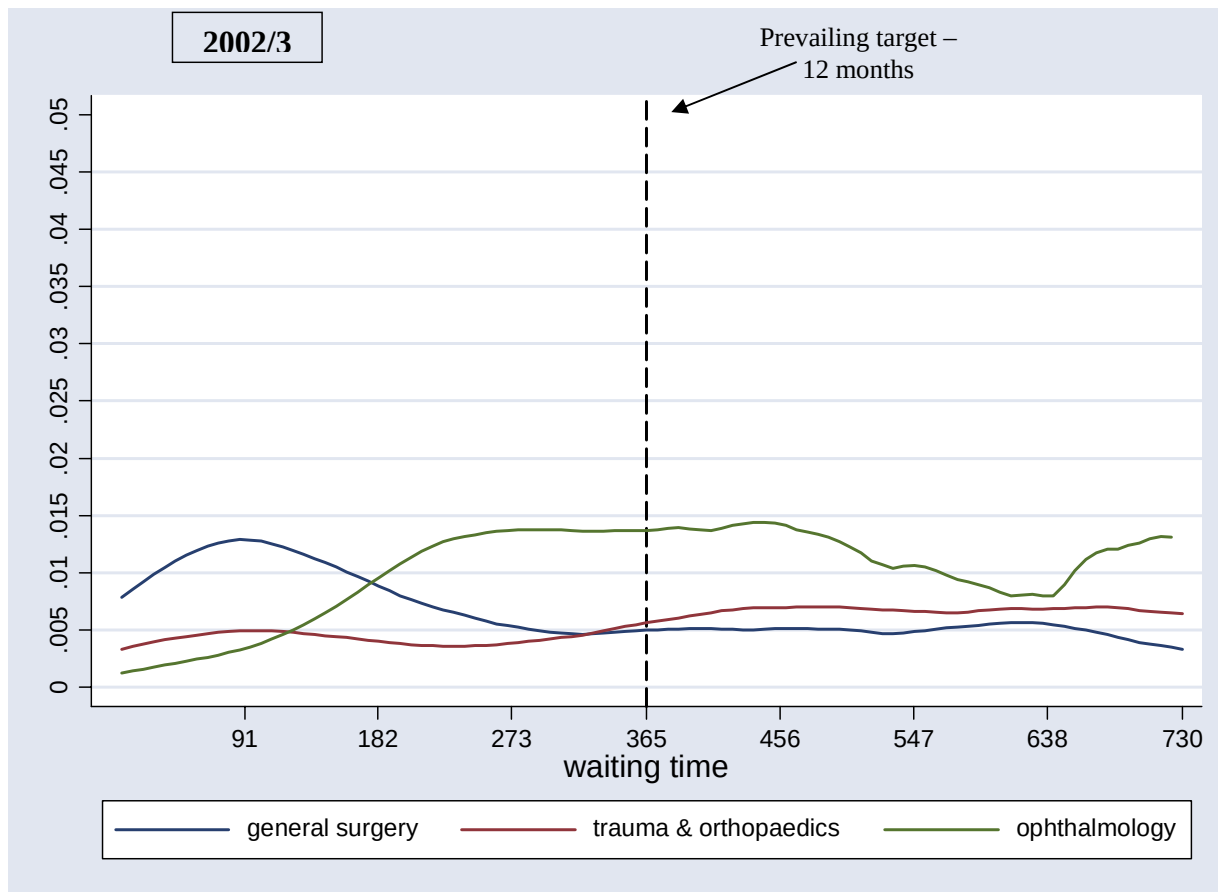
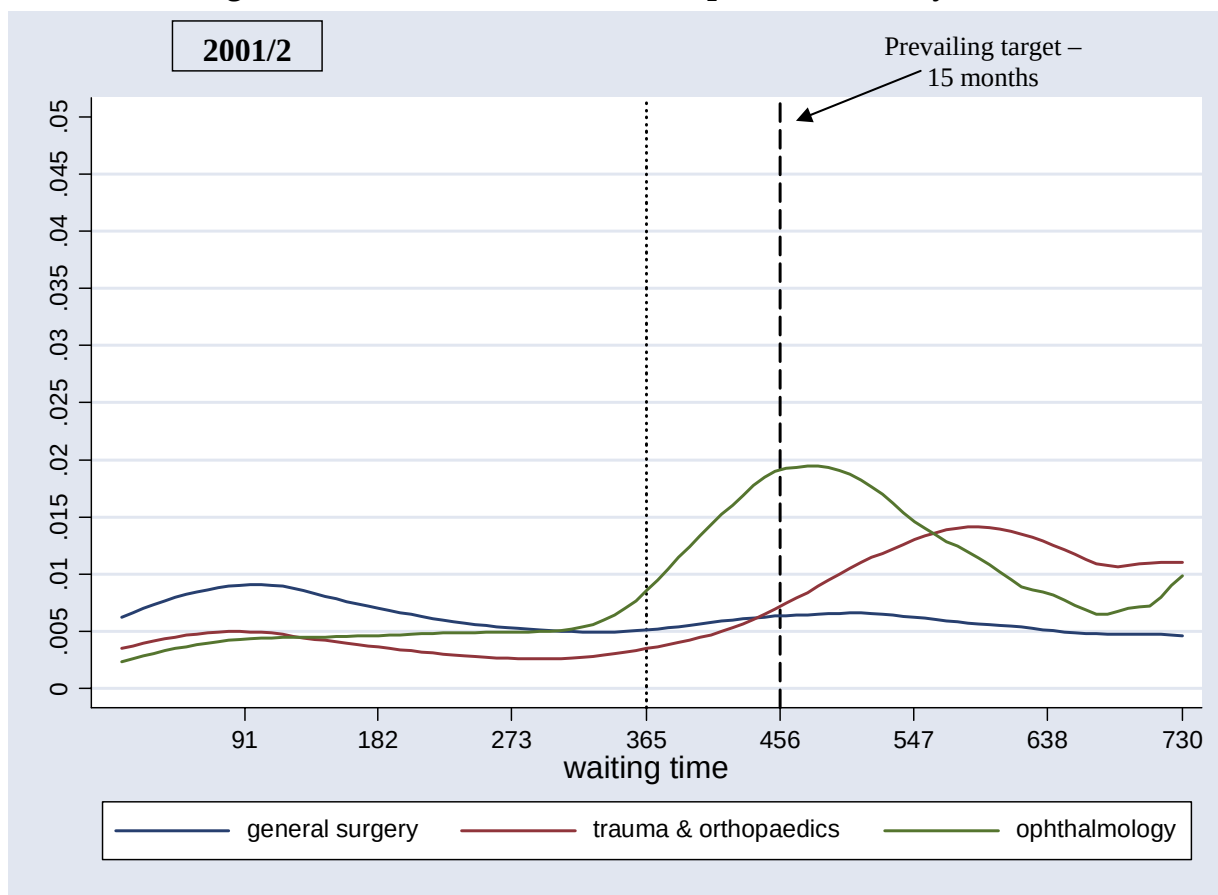


Figure 14a: Survival curves for five operations in Royal Free

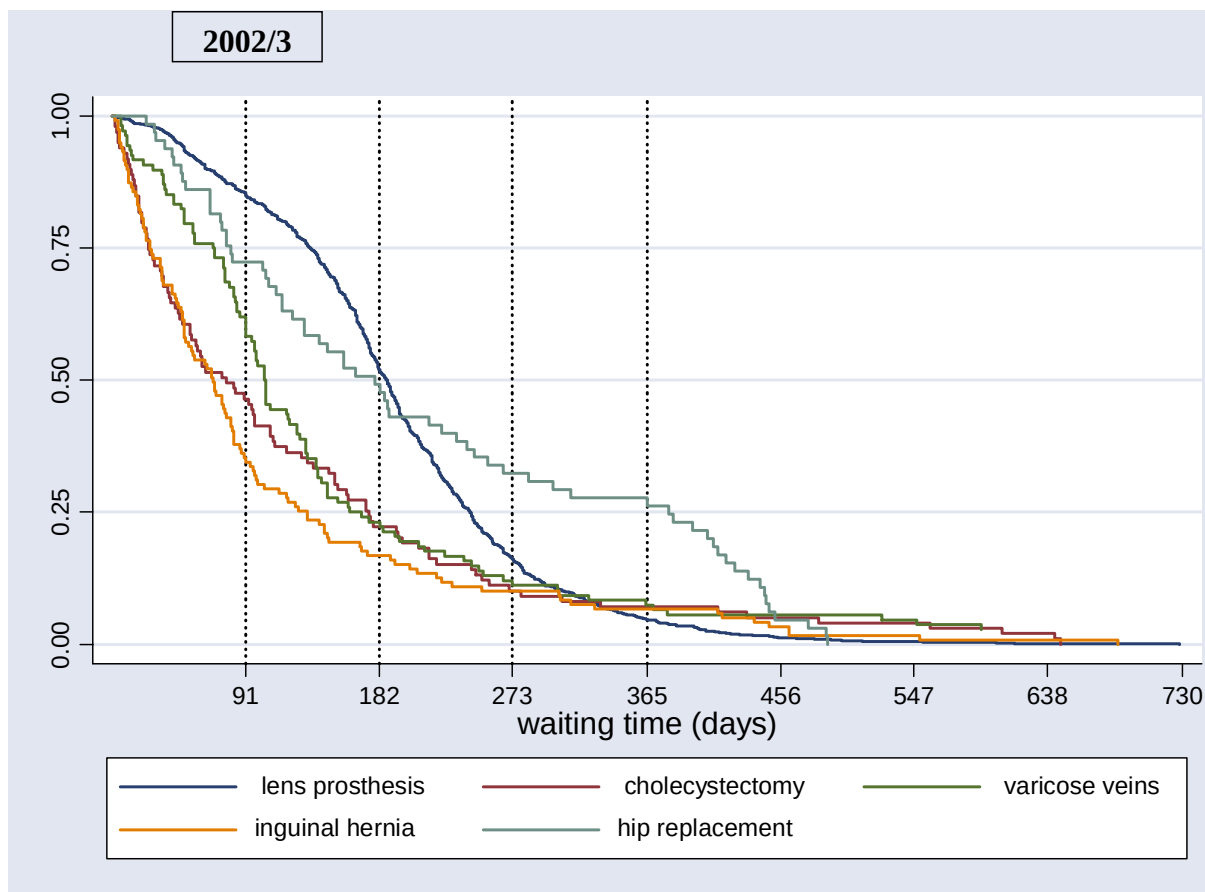
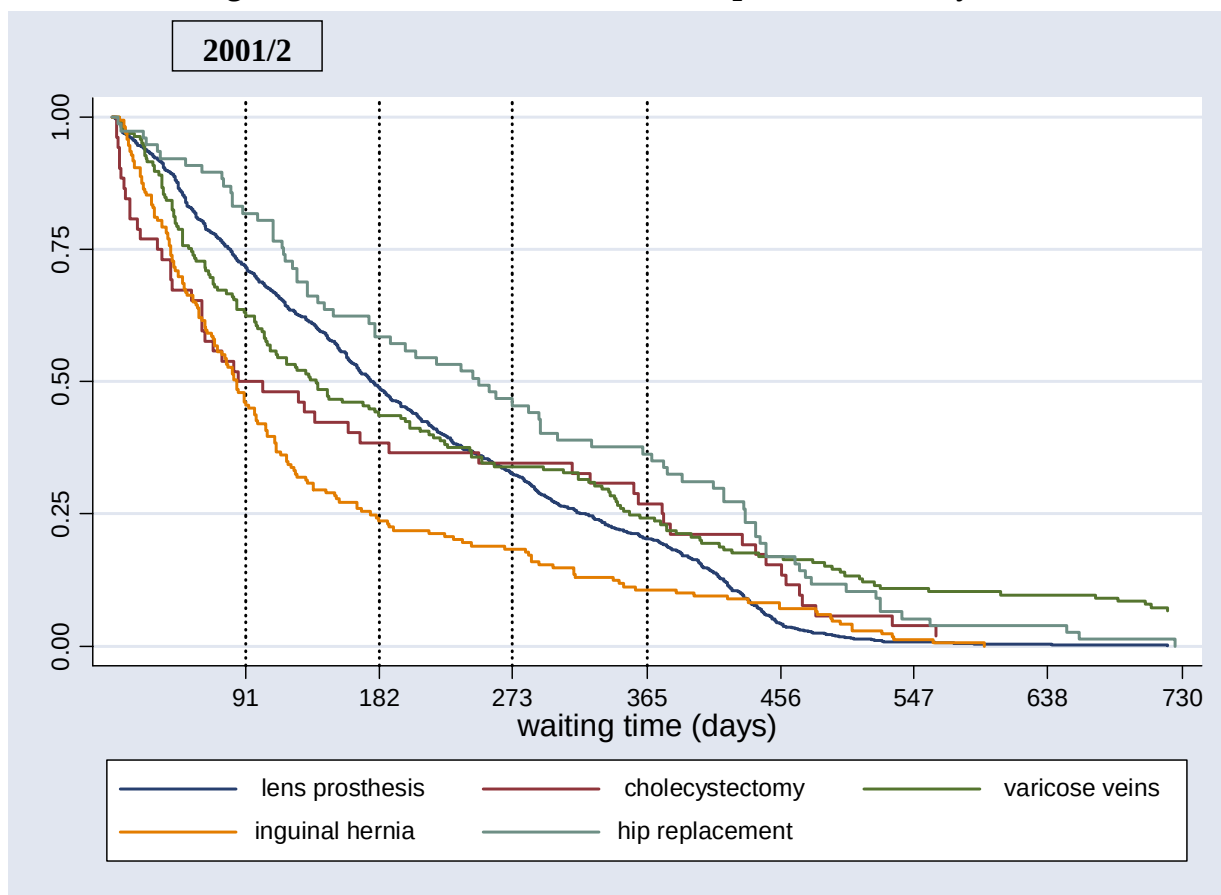
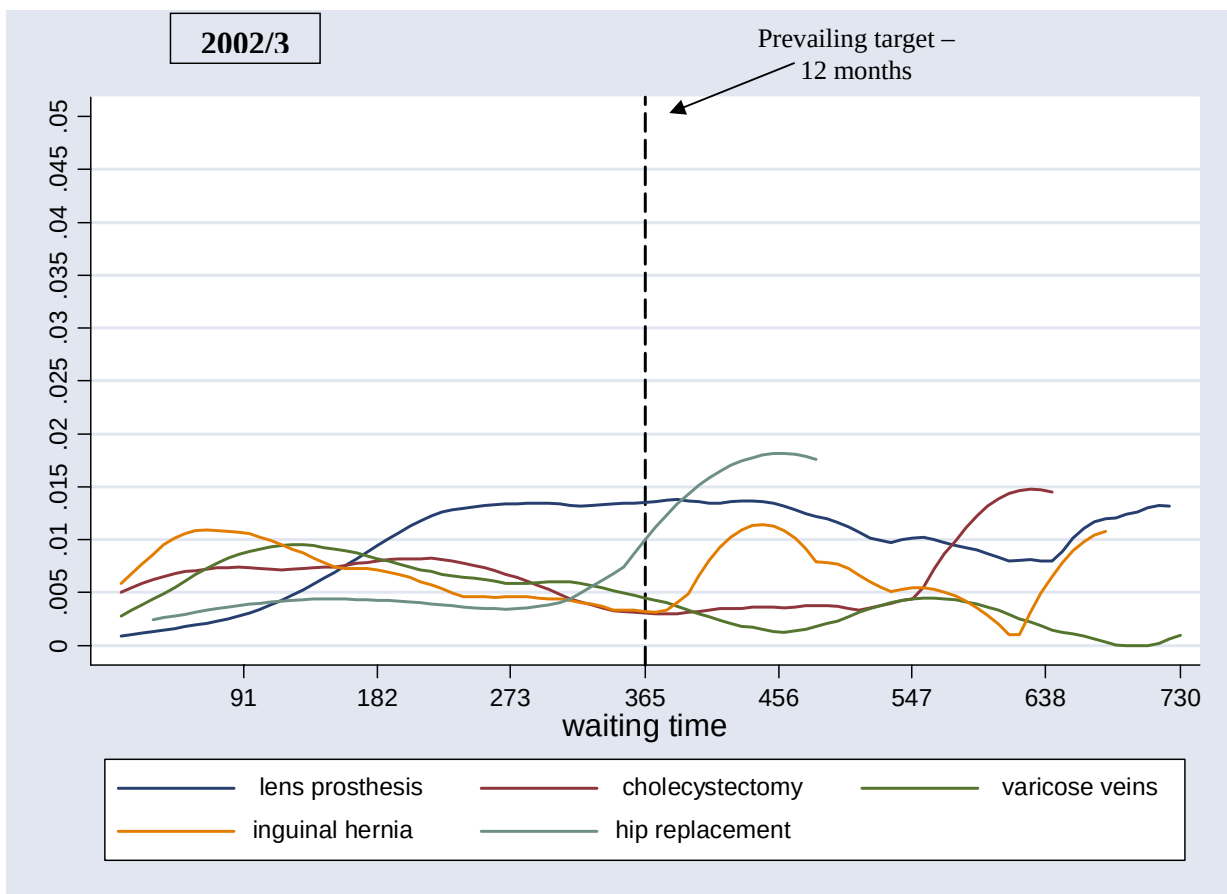
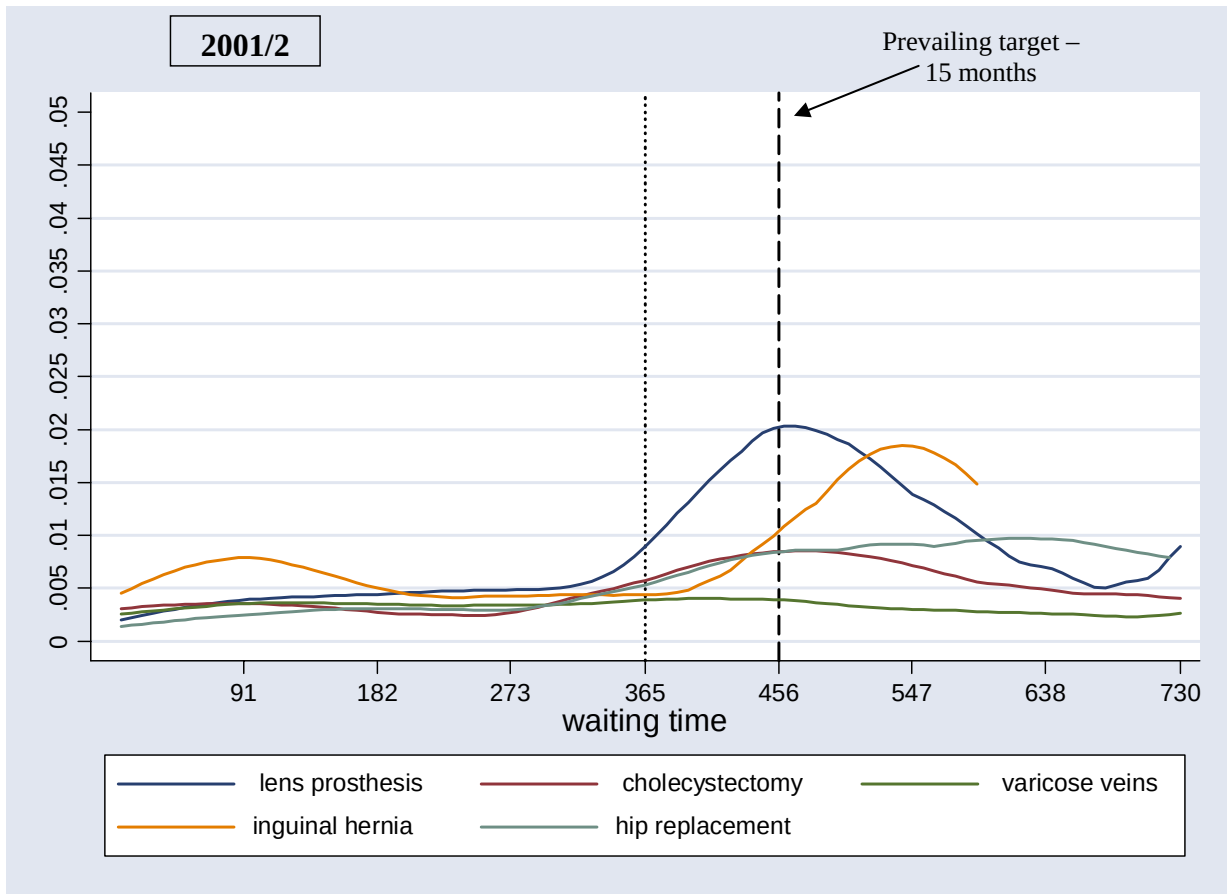


Figure 14b: Hazard rates for five operations in Royal Free



The other five NHS trusts adopt different behavioral responses to meet the 15 month target for 2001/2 and the 12 month target for 2002/3. Some of the estimated survival and hazard functions by overall waiting time, by specialty and different operative procedures during 2001/2 and 2002/3 are illustrated in appendix I. Moreover, the proportion of patients leaving the list for admission with respect to the time they had to wait is displayed in table 1.

Generally, Southampton, Guys and Manchester trusts have the longest waiting lists, followed by Royal Free and Nottingham whose patients have to wait moderate time periods to be treated. Birmingham and Newcastle providers perform the best of all seven trusts as their waiting times as a whole, by different specialty and by the majority of the operations tested are the shortest.

Although the majority of providers manage more effectively their waiting lists during the second financial year, not all of them show signs of improvement; for example, lens prosthesis waiting lists for Manchester increased during 2002/3. We also observe providers to be offering improved surgical treatment in terms of waiting for some operative procedures and not for others. Guys hospital falls in that category; for example, the rate at which patients leave the lens prosthesis waiting list for treatment is quicker for 2002/3 than the previous year, but the exact opposite effect characterises the waiting times distribution of cholecystectomy. Clearly, in addition to some universal management tactics to reduce waiting times for elective surgery, trusts also develop appropriate measures to deal with the separate surgical waiting lists. Besides the importance of other factors such as the natural history of each disease and its clinical evolution and severity, this behavioural response by trusts might explain part of the waiting times variations.

Differences among trusts reflect the level and type of activity employed by them at any given time. Some providers exhibit greater activity to tackle the excessive waits of their patients and offer better quality health care than others (e.g. the cases of Birmingham and Newcastle trusts). Some manage only the longest waiters while others appear to tackle the whole spectrum of the waiting times distribution. Although not commonly seen, there are trusts in which the probability of admission remains constant with respect to the time period patients spent on lists (e.g. the case of Royal Free).

Table 1: Proportions of patients leaving lists for admission with respect to the time they already had to wait

	2001/02																	
	<i>overall waiting (days)</i>		<i>by specialty (waiting times, days)</i>						<i>by operation (waiting times, days)</i>									
			general surgery		orthopaedics		ophthalmology		lens prosthesis		cholecystectomy		varicose vein		inguinal hernia		hip replacement	
<i>Proportion of patients leaving list</i>	50%	75%	50%	75%	50%	75%	50%	75%	50%	75%	50%	75%	50%	75%	50%	75%	50%	75%
trust																		
Royal Free	90	217	49	126	112	327	162	301	178	323	103	376	139	353	85	176	250	432
Nottingham	94	247	61	142	112	235	134	282	263	312	162	315	109	204	132	238	286	397
Southampton	144	236	71	174	218	433	167	200	180	203	156	365	352	412	137	236	479	507
Guys	149	308	158	414	163	311	137	220	150	218	123	221	322	359	222	474	244	470
Birmingham	57	143	34	63	111	259	194	253	205	256	47	73	55	95	46	77	213	287
Newcastle	75	125	24	62	-	-	89	134	91	128	109	193	-	-	69	128	-	-
Manchester	130	266	62	163	140	416	174	274	212	293	152	409	369	452	127	405	250	519

	2002/03																	
	<i>overall waiting (days)</i>		<i>by specialty (waiting times, days)</i>						<i>by operation (waiting times, days)</i>									
			general surgery		orthopaedics		ophthalmology		lens prosthesis		cholecystectomy		varicose vein		inguinal hernia		hip replacement	
<i>Proportion of patients leaving list</i>	50%	75%	50%	75%	50%	75%	50%	75%	50%	75%	50%	75%	50%	75%	50%	75%	50%	75%
trust																		
Royal Free	97	201	49	94	113	308	180	239	188	244	78	175	105	170	69	133	179	380
Nottingham	101	228	72	174	130	270	106	228	205	246	231	335	96	142	125	253	236	342
Southampton	155	267	68	202	220	382	188	243	217	250	161	336	375	421	151	244	298	404
Guys	126	283	161	330	183	324	75	150	67	120	213	347	144	204	262	354	229	366
Birmingham	65	153	40	71	119	234	144	198	153	201	49	79	72	117	47	94	181	262
Newcastle	73	120	22	64	-	-	87	131	87	123	87	123	-	-	96	187	-	-
Manchester	135	311	86	257	137	321	171	321	275	349	157	324	297	357	173	328	278	413

Survival analysis with covariates

For the AFT models, the dependent variable is waiting time until admission; for the PH models it is the hazard rate. For both models, the independent variables are Age, which was calculated as age minus 50, the average for the sample, and a series of dummy variables representing sex, admission category, main specialty, patient classification and ethnicity. This defines a “reference group” of people that are male, 50 years old, NHS patients, admitted in general surgery, admitted to inpatients and white. The dummy variables are therefore Female; Private patient; Orthopaedics; Ophthalmology; Day case; Black; Indian; and Other ethnicity,

Table 2: Accelerated Failure Time Models

	<i>Exponential</i>		<i>Weibull</i>		<i>Log-normal</i>		<i>Log-logistic</i>	
T	Coef.	P> z	Coef.	P> z	Coef.	P> z	Coef.	P> z
Age	.0005929	0.000	.0006069	0.000	.000834	0.000	.0005195	0.000
Female	.0076326	0.001	.0079359	0.001	.0026647	0.362	-.0064518	0.024
Private patient	-1.293627	0.000	-1.288302	0.000	-1.416566	0.000	-1.471763	0.000
Orthopaedics	.4333493	0.000	.4277083	0.000	.7011317	0.000	.748772	0.000
Ophthalmology	.4204603	0.000	.4141631	0.000	.6748901	0.000	.7456852	0.000
Day case	-.2501083	0.000	-.2502762	0.000	-.1594466	0.000	-.2244831	0.000
Black	-.0047612	0.633	-.0047061	0.629	-.0128596	0.291	-.0161637	0.176
Indian	-.0441387	0.000	-.0442778	0.000	-.0204272	0.026	-.0373673	0.000
Other ethnicity	-.2797464	0.000	-.2782713	0.000	-.3077501	0.000	-.3418351	0.000
Cons	4.916261	0.000	4.929258	0.000	4.16032	0.000	4.262388	0.000
ln_p			.0237477	0.000				
P			1.024032					
1/p			.976532					
ln_sig					.2005121	0.000		
Sigma					1.222028			
ln_gam							-.3788122	0.000
Gamma							.6846742	
Log likelihood	-1104677.4		-1104356.9		-1150796.2		-1147369	

Table 2 shows the results using four different AFT models. In interpreting these, it should be noted that the very large sample size means that the statistical significance of a variable’s coefficient is a poor guide to its practical significance. The only coefficients that are not statistically significant are those for the black ethnic group, in all specifications, and for females in the AFT Log-normal specification. The models as a whole are also statistically significant with high log-likelihood values.

Because the models produce similar results, we will discuss only the AFT-Exponential model. The antilog of the constant term is the average waiting time for the reference group as defined above, ie: $e^{4.91} = 136.5$ days. Although age has a significant coefficient, this translates into an increase of less than one day’s waiting time for a one-year increase in age, other things being equal. The changes in waiting times due to being female or Indian are also less than one day. Such differences are obviously of no account, but other findings are of more interest. Other things being

equal, private patients wait on average 99 fewer days; Orthopaedic and Ophthalmology patients wait on average 74 and 71 more days; day case patients wait on average 39 fewer days; “Other” ethnic groups wait on average 44 fewer days.

<i>Table 3: Proportional Hazard Models</i>							
	<i>Exponential</i>		<i>Weibull</i>		<i>Gompertz</i>		<i>Cox</i>
t	Haz. Ratio	P> z	Haz. Ratio	P> z	Haz. Ratio	P> z	Haz. Ratio P> z
Age	.9994073	0.000	.9993787	0.000	.9993014	0.000	.9992734 0.000
Female	.9923965	0.001	.9919063	0.001	.9911281	0.000	.9875895 0.001
Private patient	3.645988	0.000	3.74066	0.000	3.751843	0.000	3.594825 0.000
Orthopaedics	.648334	0.000	.6453342	0.000	.6463513	0.000	.6437025 0.000
Ophthalmology	.6567444	0.000	.6543478	0.000	.6590856	0.000	.6672428 0.000
Day case	1.284164	0.000	1.292128	0.000	1.297126	0.000	1.319649 0.000
Black	1.004773	0.633	1.004831	0.629	1.005844	0.559	.9981193 0.850
Indian	1.045127	0.000	1.046385	0.000	1.048196	0.000	1.044269 0.000
Other ethnicity	1.322794	0.000	1.329707	0.000	1.328031	0.000	1.339732 0.000
ln_p			.0237477	0.000			
p			1.024032				
1/p			.976532				
gamma					.0003357	0.000	
Log likelihood	-1104677.4		-1104356.9		-1103564.3		

Table 3 shows the results from the PH models. All covariates are again statistically significant apart from the black ethnic group. The results from the parametric models are consistent statistically, functionally and quantitatively with those from the Cox regression.

Overall, there is very little to choose between these models, so the results presented above concerning the impact of the independent variables may be taken as representative. Some variables have no real impact on waiting times, such as age and sex; however, some, such as whether the patient is NHS or private, have an impact on waiting times that is significant in both statistical and practical terms.

Discussion

This study analysed waiting time data in a number of ways which suggest the potential usefulness of time-to-event analysis. The first part of the analyses that we report captures the whole set of patients admissions for elective surgery from all over UK using around 2 million record level information for each financial year. The second part of the analyses examines data at less aggregated levels – at different providers by different specialties and operative procedures.

From the point of view of the aims of this study, the first question was: how have behavioural responses to the targets affected the distribution of waiting times? The survival curve analysis confirmed the findings that would be possible from a simple comparison of waiting times, that more demanding targets were associated with

reductions in average waiting times. The value added is that the analysis showed more clearly how these reductions were achieved in terms of alterations in the shape of the waiting time distribution, rather than simply the central tendency. This finding was demonstrated more clearly by the more detailed and refined analyses of the paper. Waiting times distributions did vary at the level of each trust revealing much more details of trusts' behavioural tactics and waiting lists management over time.

The second question was: How is the probability of admission for any given waiting time affected by the targets? The hazard curve analysis casts light on two aspects of this. First, that hazard rates are in general not constant over time. The reasons for the existence of differences over time might be due, for example to the characteristics of the operation or underlying disease, for example increasing severity and urgency due to delays might lead to a simple increasing probability over time. However, the existence of peaks, where probability rises then falls, suggests that other factors are taken into account. The fact that these in many cases coincide with target waiting times suggests that waiting list management is a major factor. Secondly, the peaks do change over time in line with changes in targets.

The third question was: To what extent are clinical distortions evident in the pattern of admissions? There is less evidence available on this. In general hazard rate peaks which were at or higher than target waiting times moved towards new targets over time, whilst those which were already shorter than targets did not move or in some cases appeared later in the distribution. This provides some indication that waiting list management does involve adjustment of relative priorities.

The fourth question was: Can variations between individuals' waiting times be explained by clinical, patient or provider-level characteristics? These results suggest that they can, although more analyses and further data are needed to answer this properly. From an equity point of view, it is useful to know that characteristics such as age and sex do not affect waiting times in any important way, and that we can be confident about those findings because of the very large sample size. Some findings suggest that more investigation is required, for example the difference in waiting times for "Other" ethnic groups. However, some large differences are of immediate interest. Particularly interesting is the very large difference between NHS and private patients - which in this context is private patients using NHS accommodation or services - suggesting that private patients have a considerable advantage in access compared to NHS patients, even though the two groups are competing users for exactly the same facilities and services.

A limitation of this study as a test of the potential advantages of time-to-event analysis is that data only included completed patient spells, with no censored observations. This is related to the way data were collected and reported by HES: Exits from the waiting list were restricted to admissions and excluded such reasons as dying, moving away, or no longer need treatment. A further limitation to the study is that some observations suggested extremely long waits, in some cases greater than three years; whilst this is possible, it is more likely to be due to

problems with patient record coding. We took account for such outliers by restricting analysis at waiting time up to 3 years and truncating graphs at 2 years to reveal details about the main part of the distribution, away from the long right-hand tail. Also, it is possible that Trusts may have to some extent met their targets by adjustments such as reclassifying patients that have been included on waiting lists as planned cases and reclassifying day-cases as outpatients (NAO, 2001). We have not yet explored this issue, but there is clearly potential to do so.

The final question concerned the methods themselves. As stated, we believe that we have demonstrated that time-to-event analysis does in practice provide the additional insights that it promises. It should therefore be more widely applied to waiting time data. Further analysis at less aggregated levels would be particularly useful, such as by referring GP, PCT and admitting consultant. This has three potential uses. First, it may provide an additional set of variables which explain variations in the distribution of waiting times, and therefore waiting time management. Secondly, it might offer a tool by which these individuals and organisations can learn about the practical effects of their management policies. Thirdly, it might offer a tool for assessing performance, for example for hospitals in respect of consultants' admitting behaviour and PCTs in respect of hospitals' admitting behaviour and GPs' referring behaviour.

The policy implications of our findings are important. How physicians, consultants and managers respond to the implementation of waiting time targets is decisive for the success of policies to tackle long waiting. As noted, the observation that increased elective surgery activity for patients waiting the exact times as the prevailing targets supports the suggestion that clinical distortions might have taken place. As it is inappropriate to concentrate on less urgent routine cases in preference to those who require more urgent treatment so as to meet the targets, this phenomenon may have important clinical and policy implications. To assess the practical significance of this, it would be necessary to link the distribution of waiting times to quality of life information, such as urgency, severity of illness and rates of deterioration. In addition, future research that would relate specific HRGs with cost could establish whether there is any association between the cost of different type of surgeries and trends in waiting times of the relevant patients. Such analyses would help in deciding whether or not it is necessary to introduce a more standardised prioritisation system using explicit criteria, along the lines of the "admission index" suggested by Culyer and Cullis (1976), which could incorporate clinical, social and economic factors.

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Appendix I

Figure 1: Survival curves of overall waiting times for seven trusts

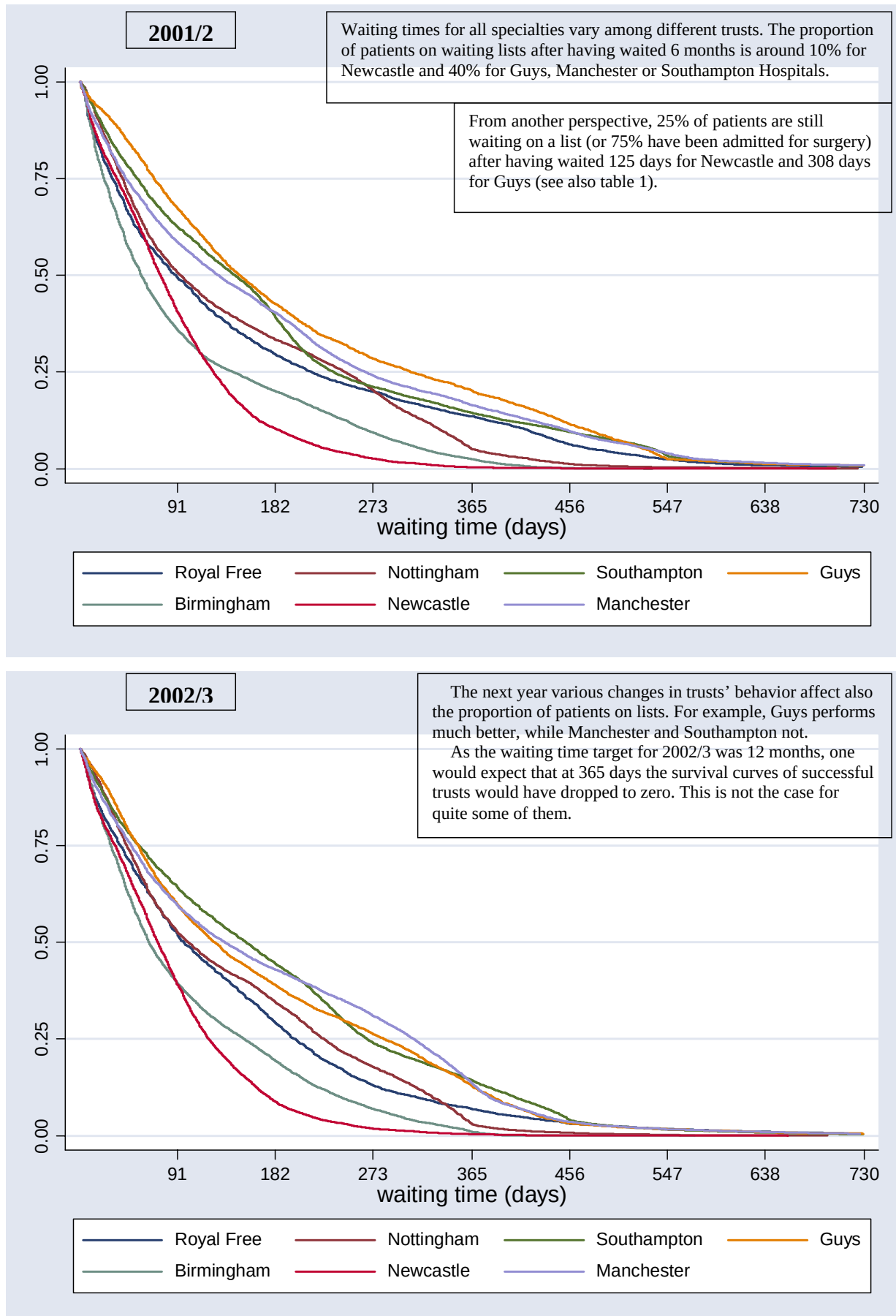


Figure 2: Two quite different behaviour responses by Guys and Manchester Trusts

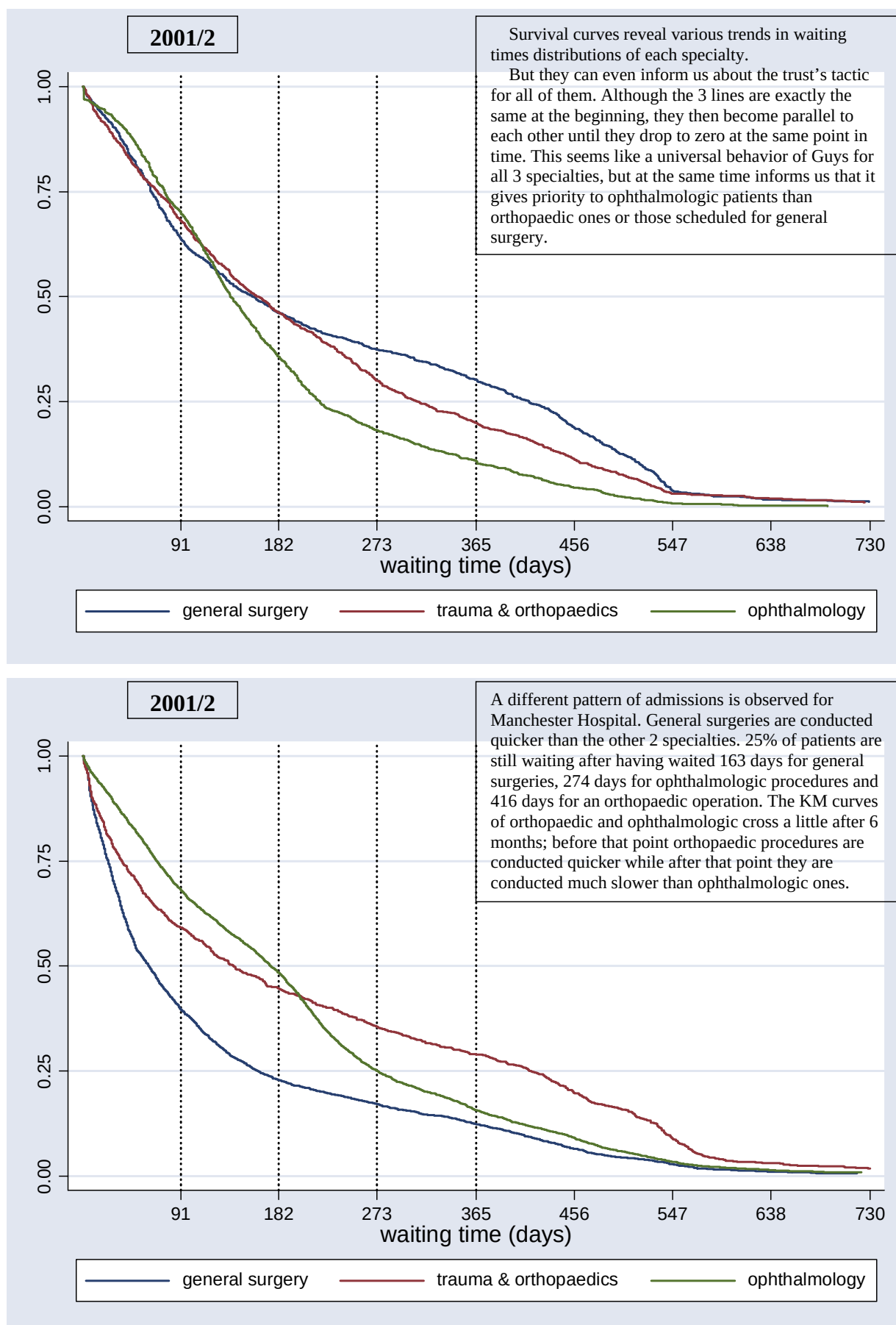


Figure 3: KM curves for three specialties in Guys and Manchester for the next financial year

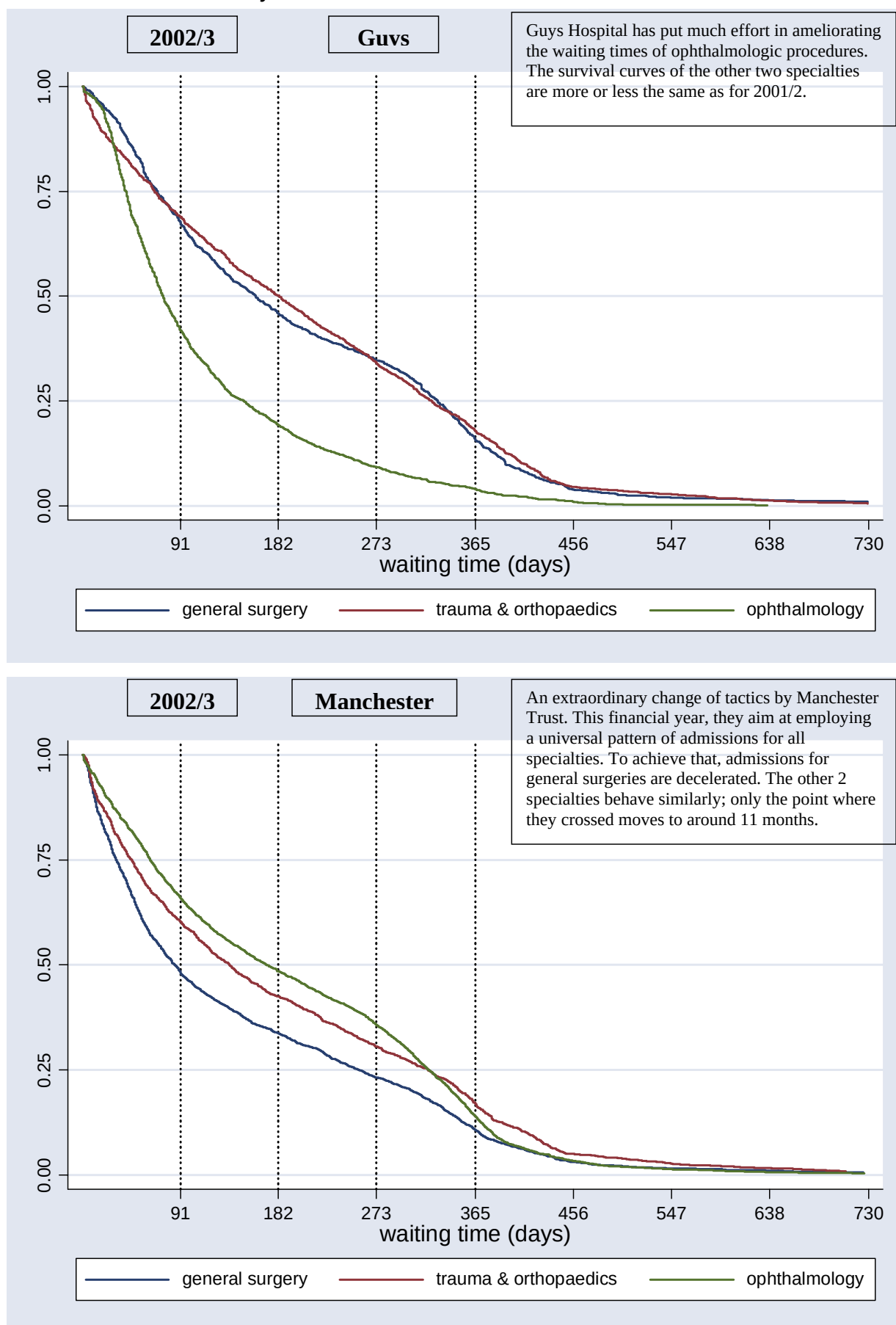


Figure 4: Shift of hazard curve's peaks towards shorter targets in Guys

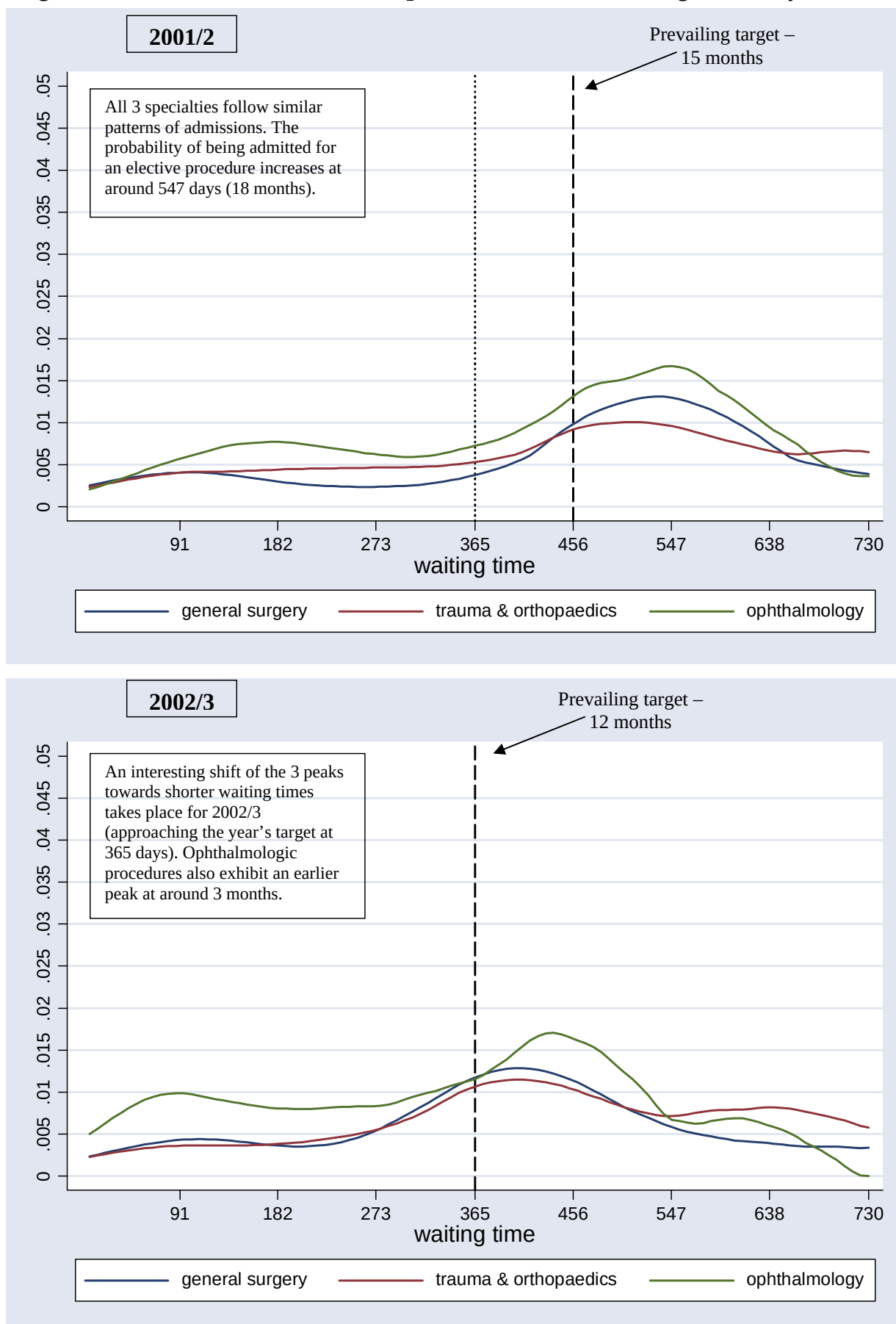


Figure 5: Hazard rates for three specialties in Manchester

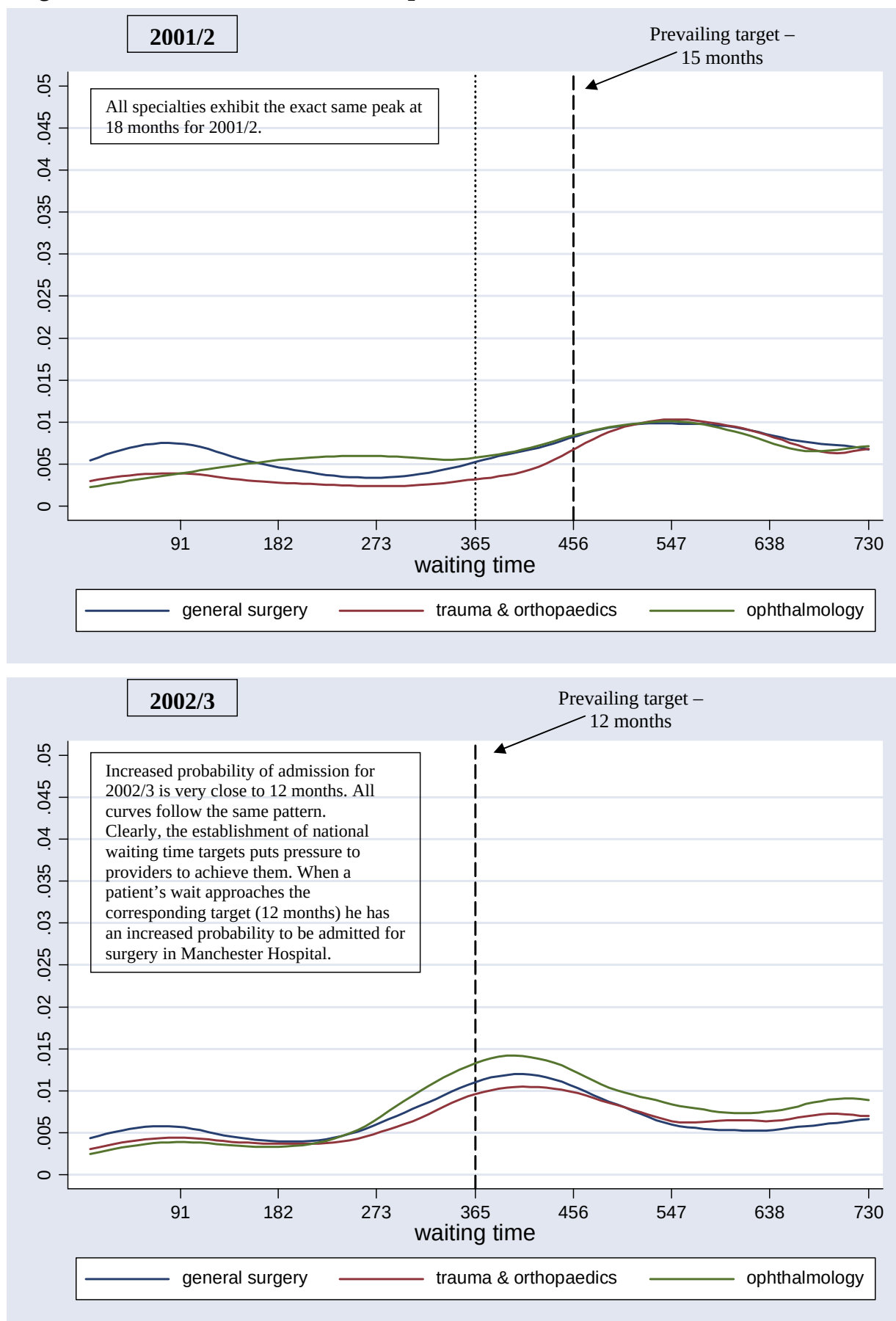


Figure 6: The KM curves for three specialties in Newcastle are almost identical for both years

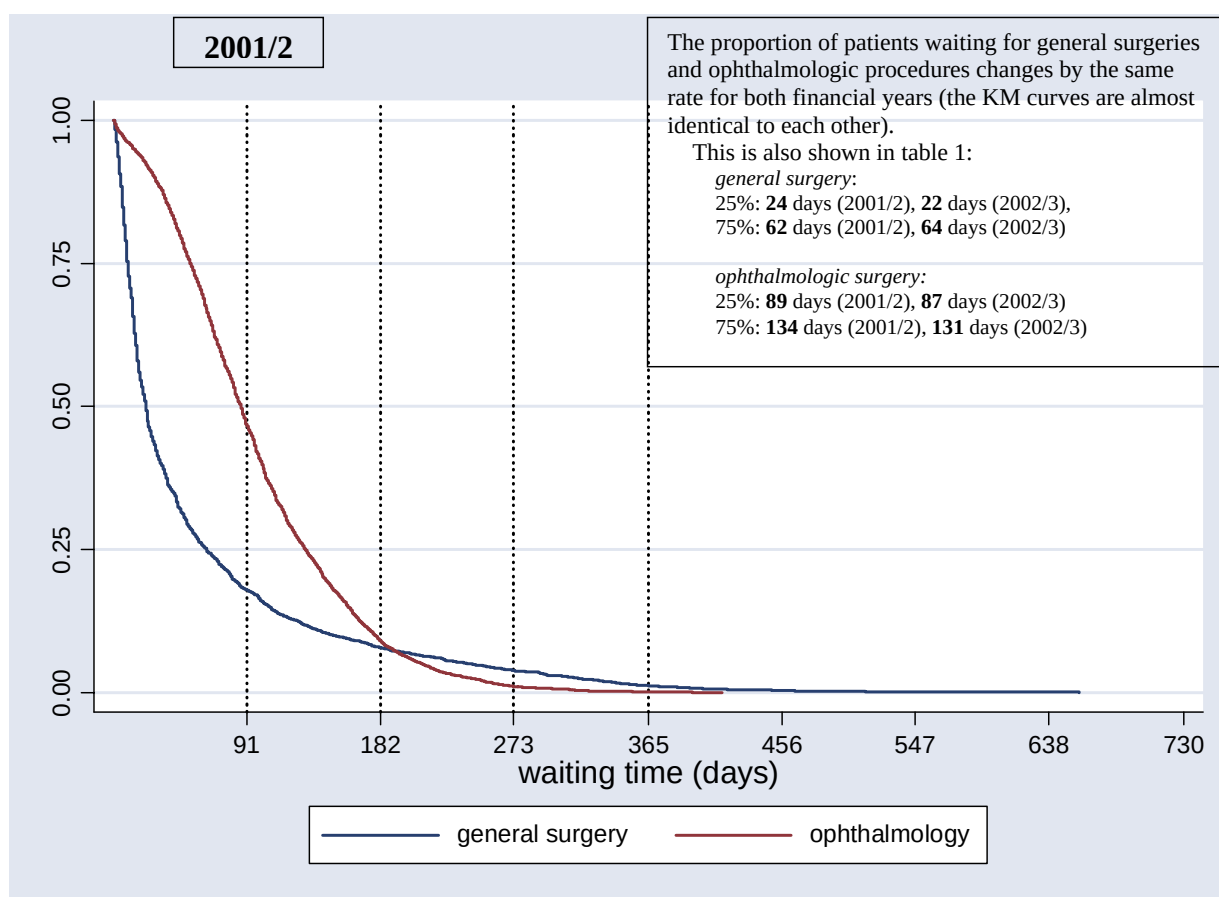
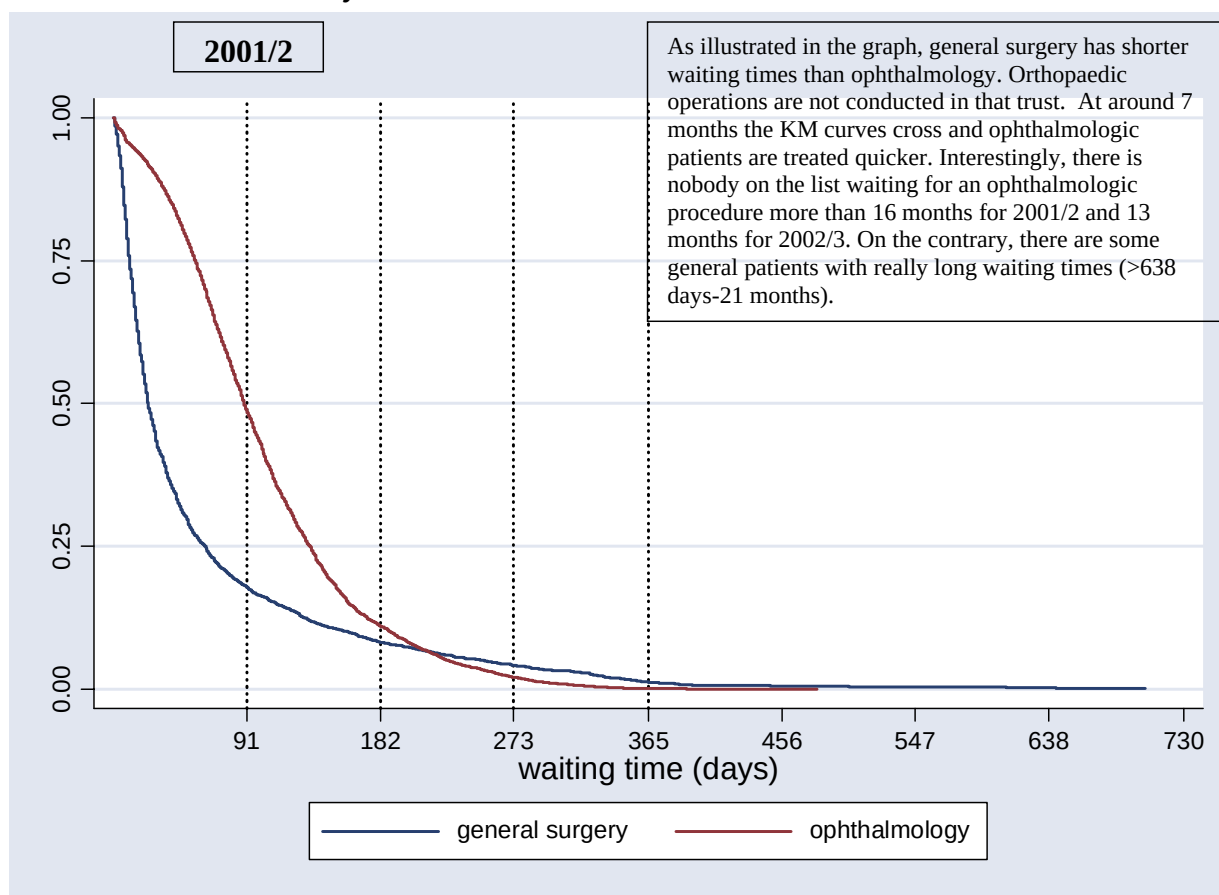


Figure 7: Hazard rates for three specialties in Newcastle

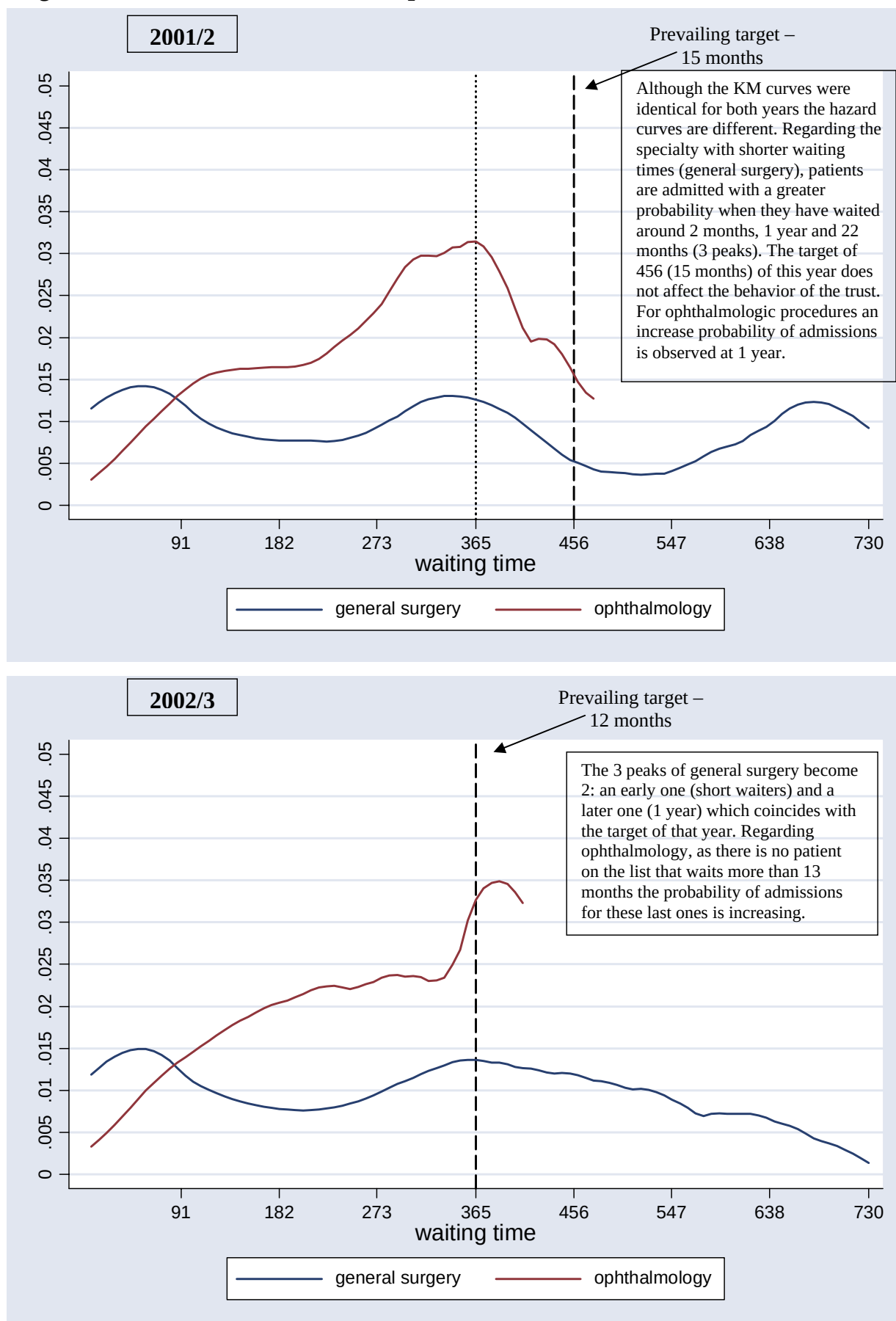


Figure 8: Moving towards a less aggregated analysis in Nottingham-Is it informative?

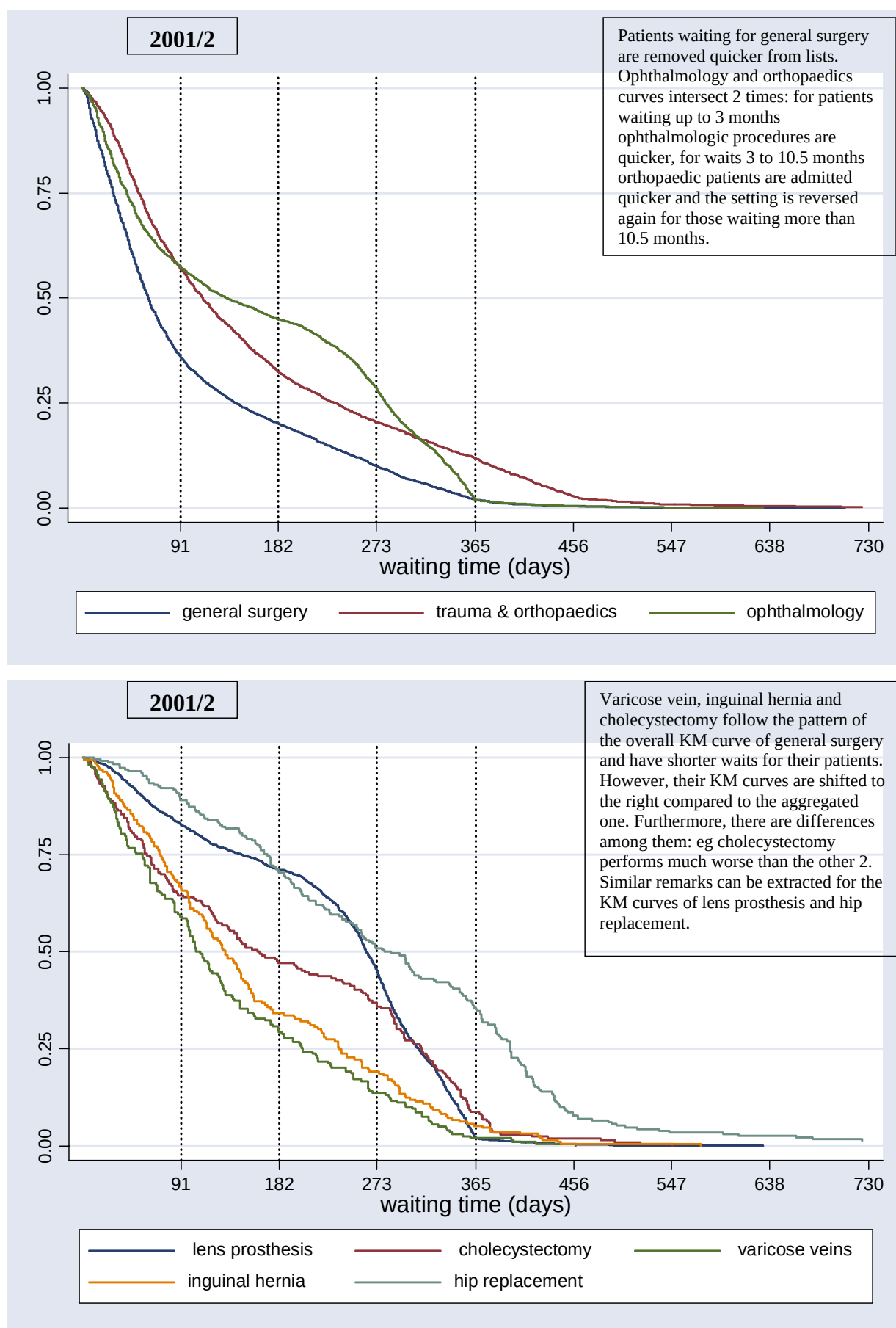


Figure 9: Survival curves by specialty and by operation in Nottingham (2002/3)

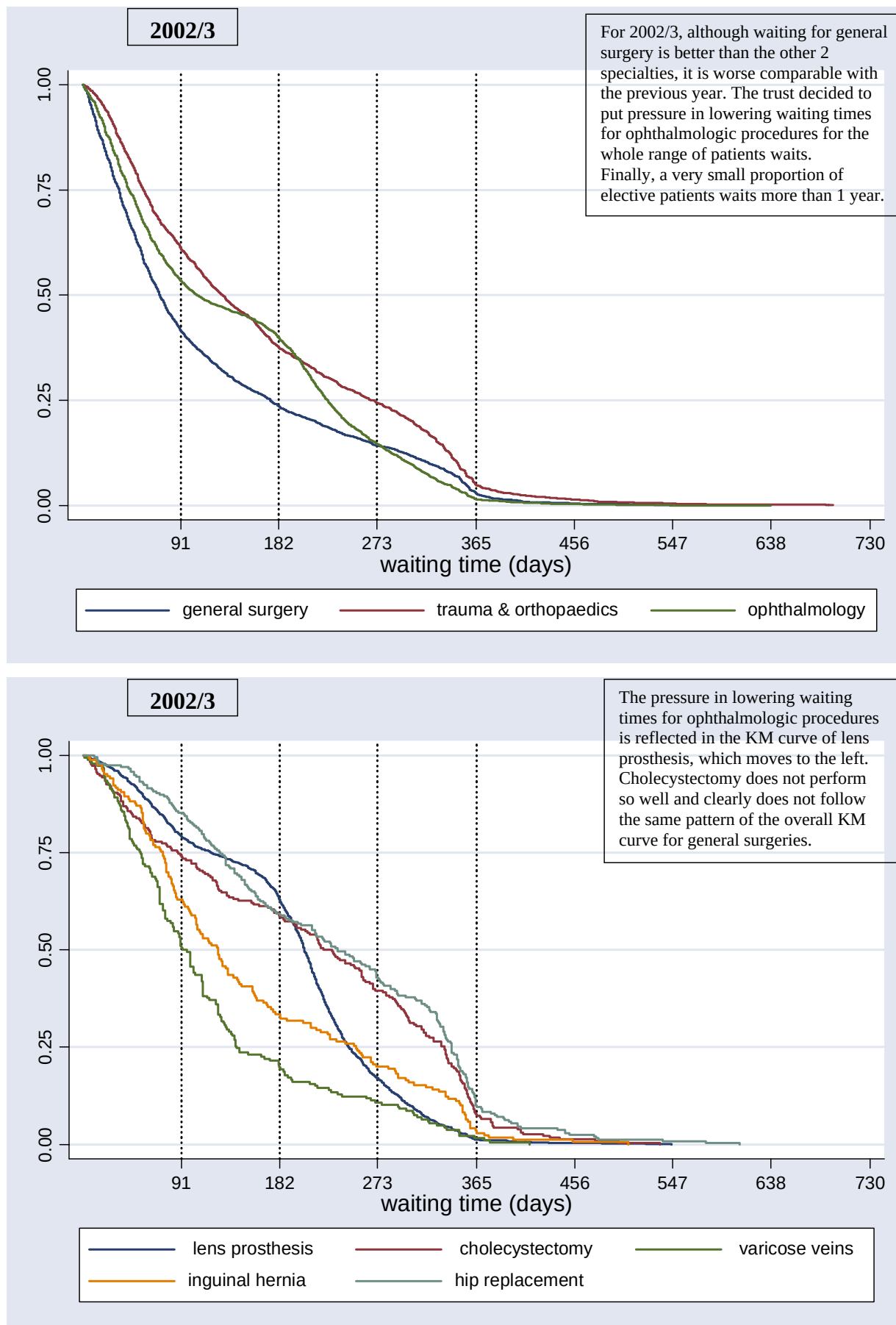


Figure 10: Hazard rates for three specialties in Nottingham

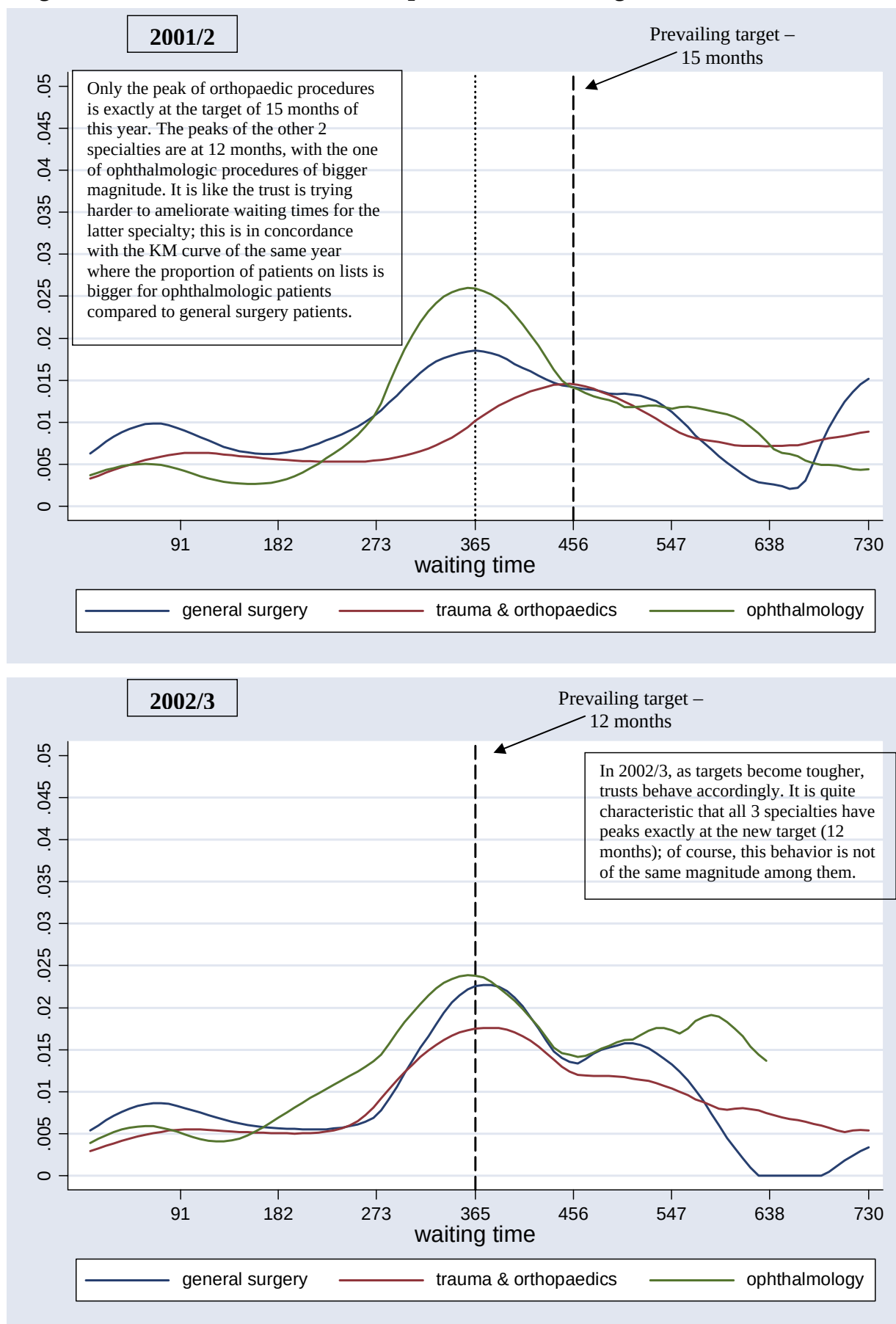


Figure 11: Hazard rates for five operations in Nottingham

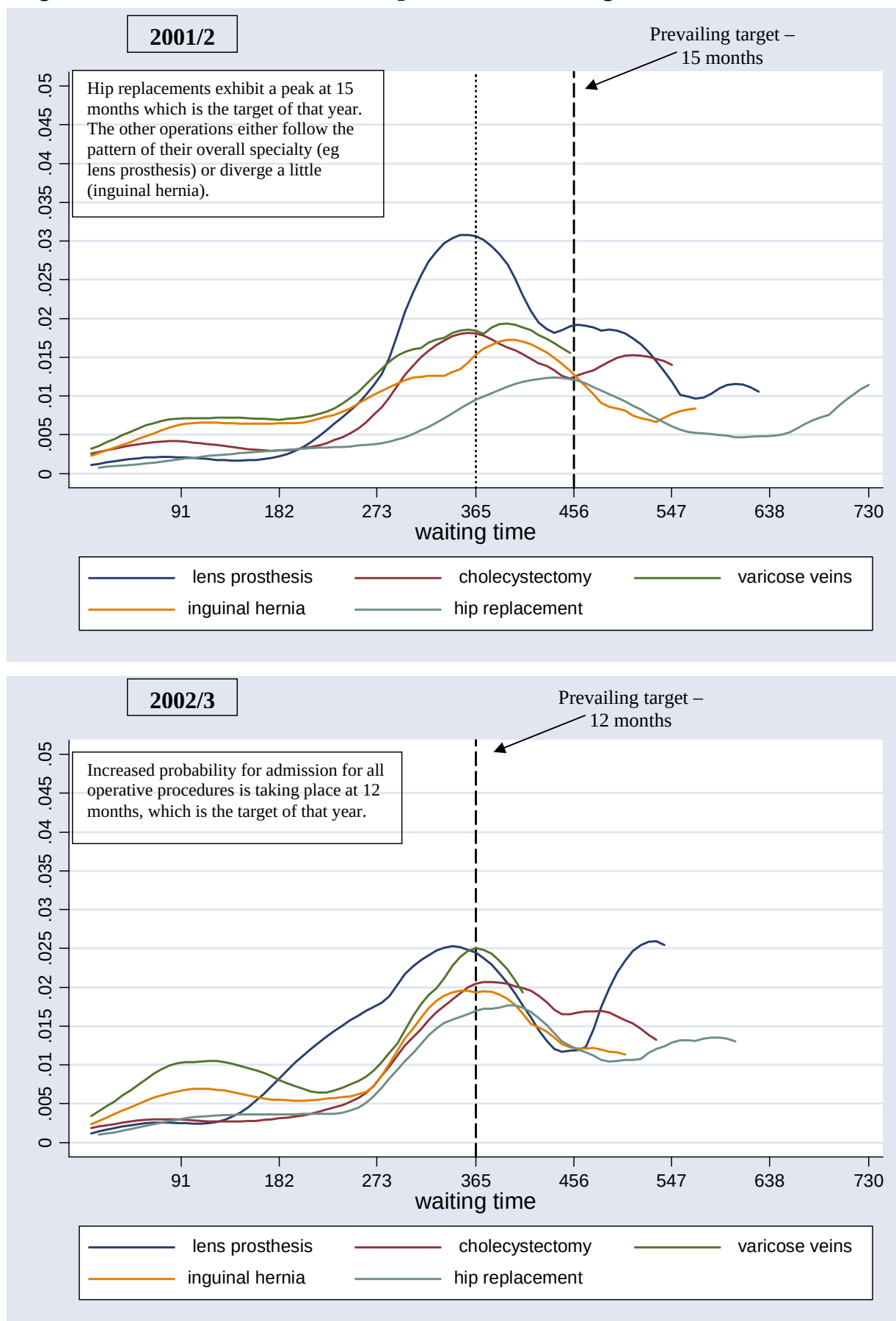


Figure 12: Moving towards a less aggregated analysis in Southampton-Is it informative?

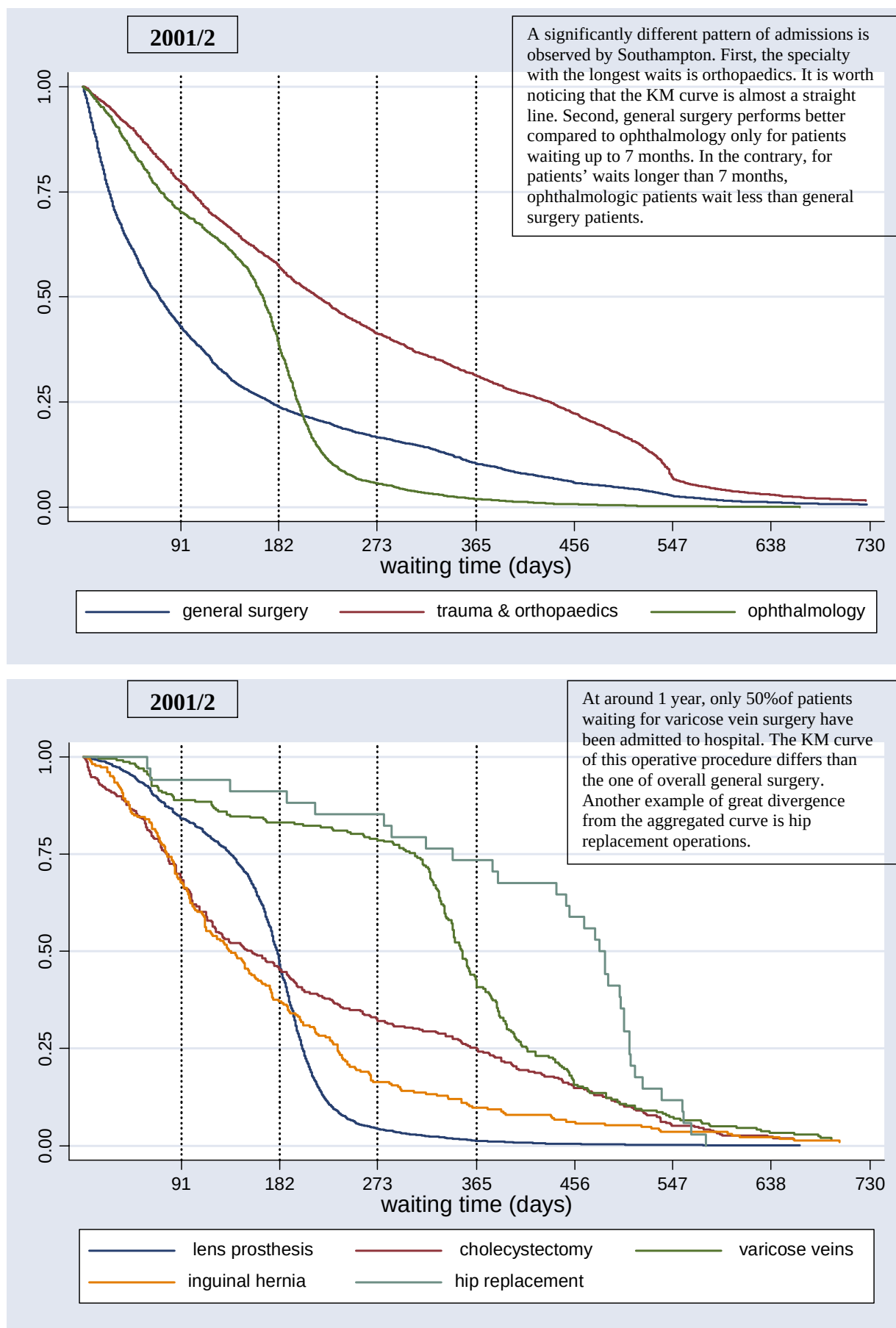


Figure 13: Southampton Trust does not seem to be influenced by national targets

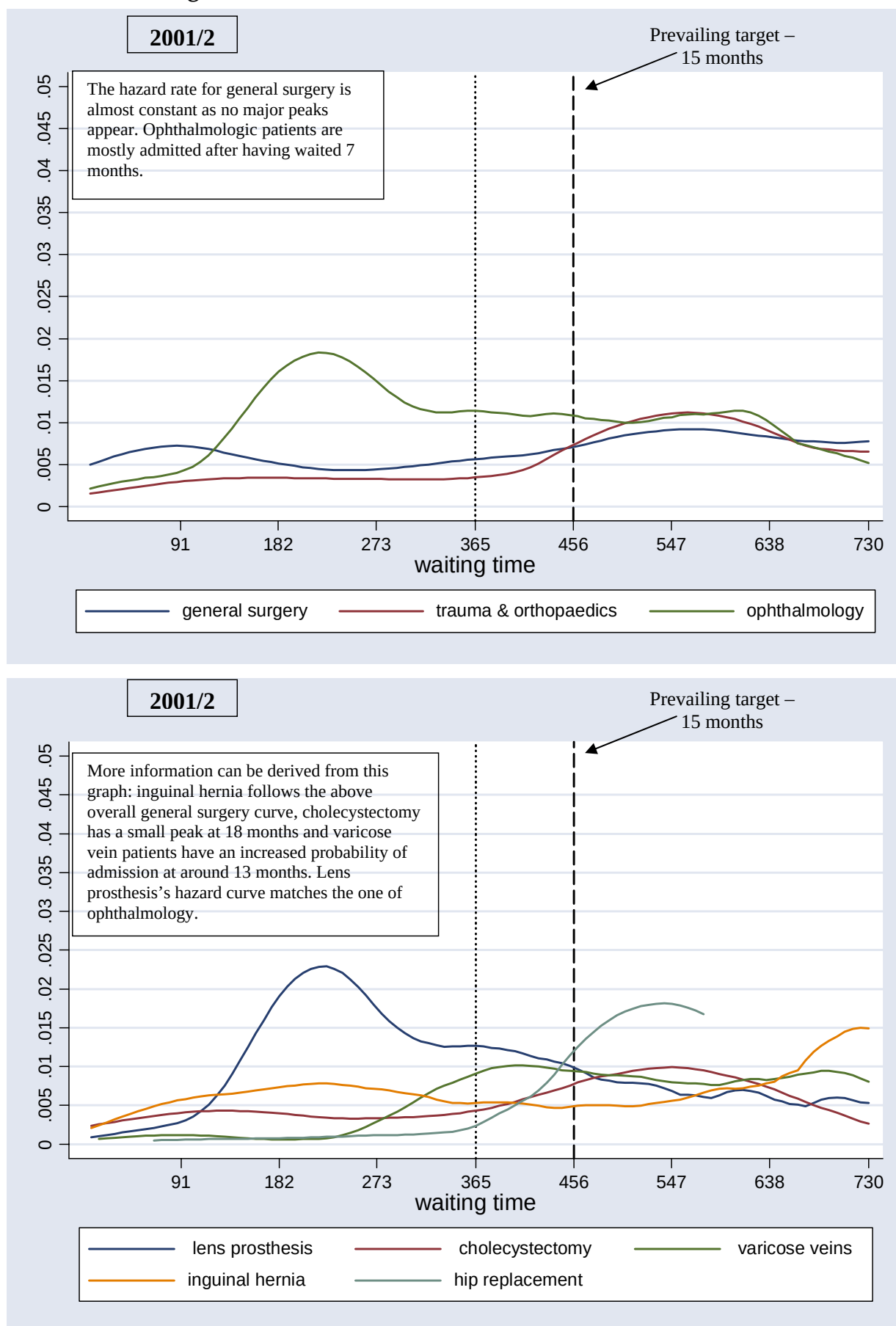


Figure 14: Survival curves by specialty and by operation in Southampton (2002/3)

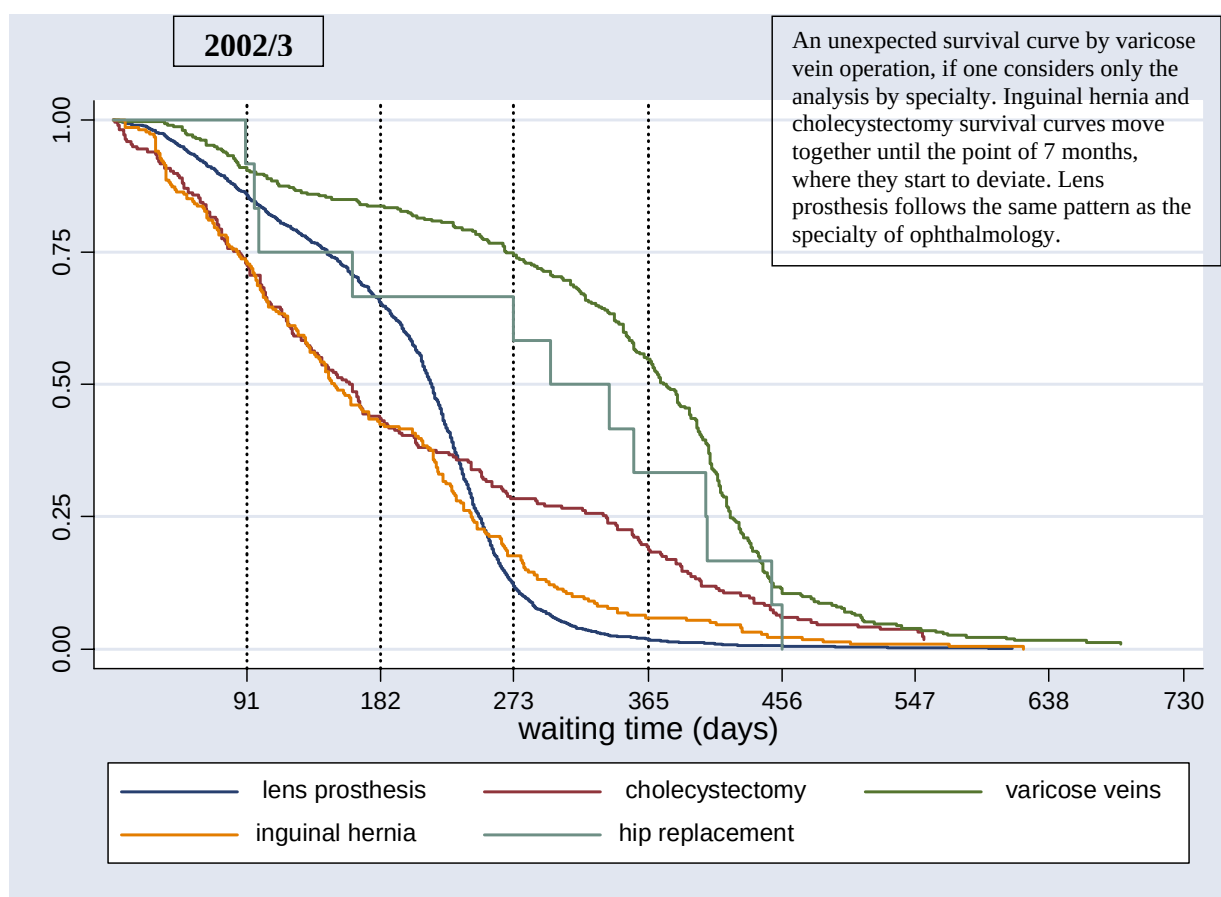
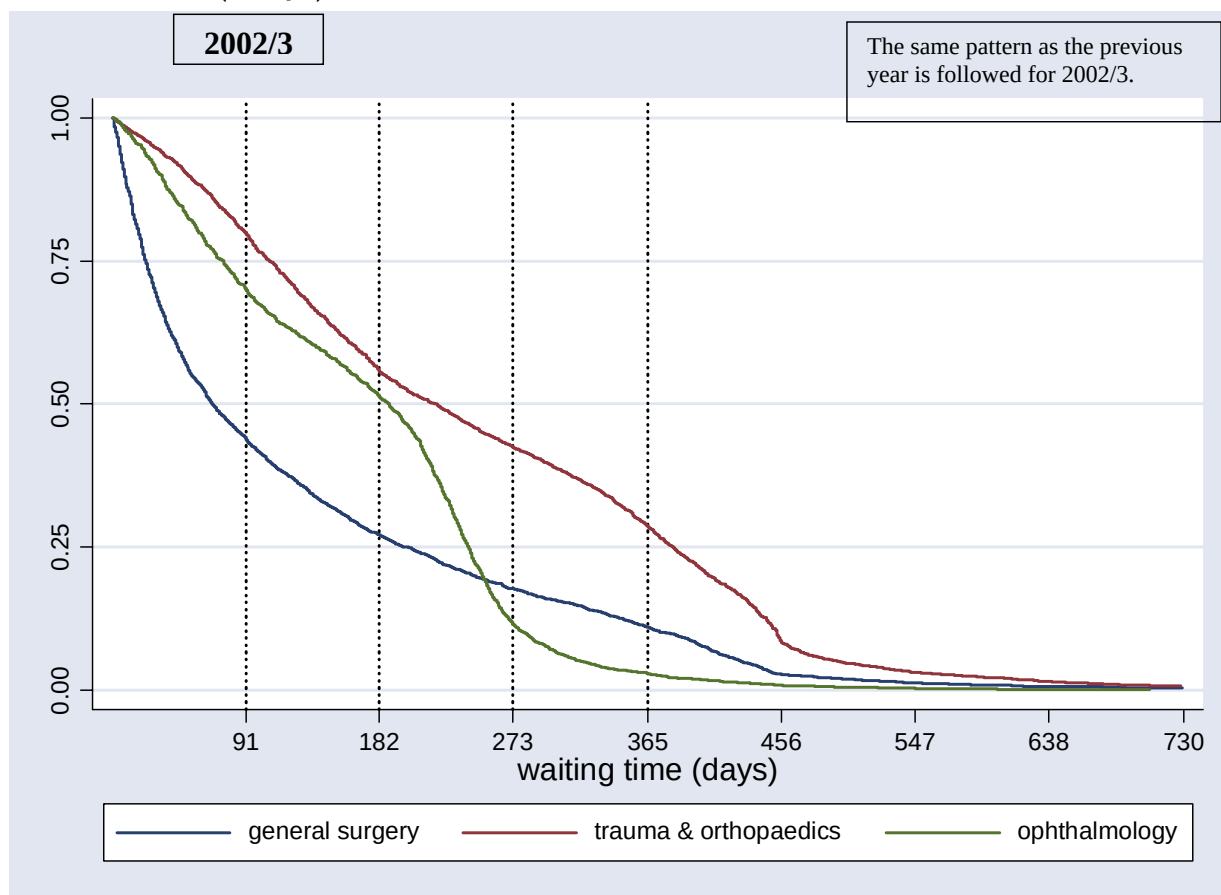


Figure 15: Hazard rates by specialty and by operation in Southampton (2002/3)

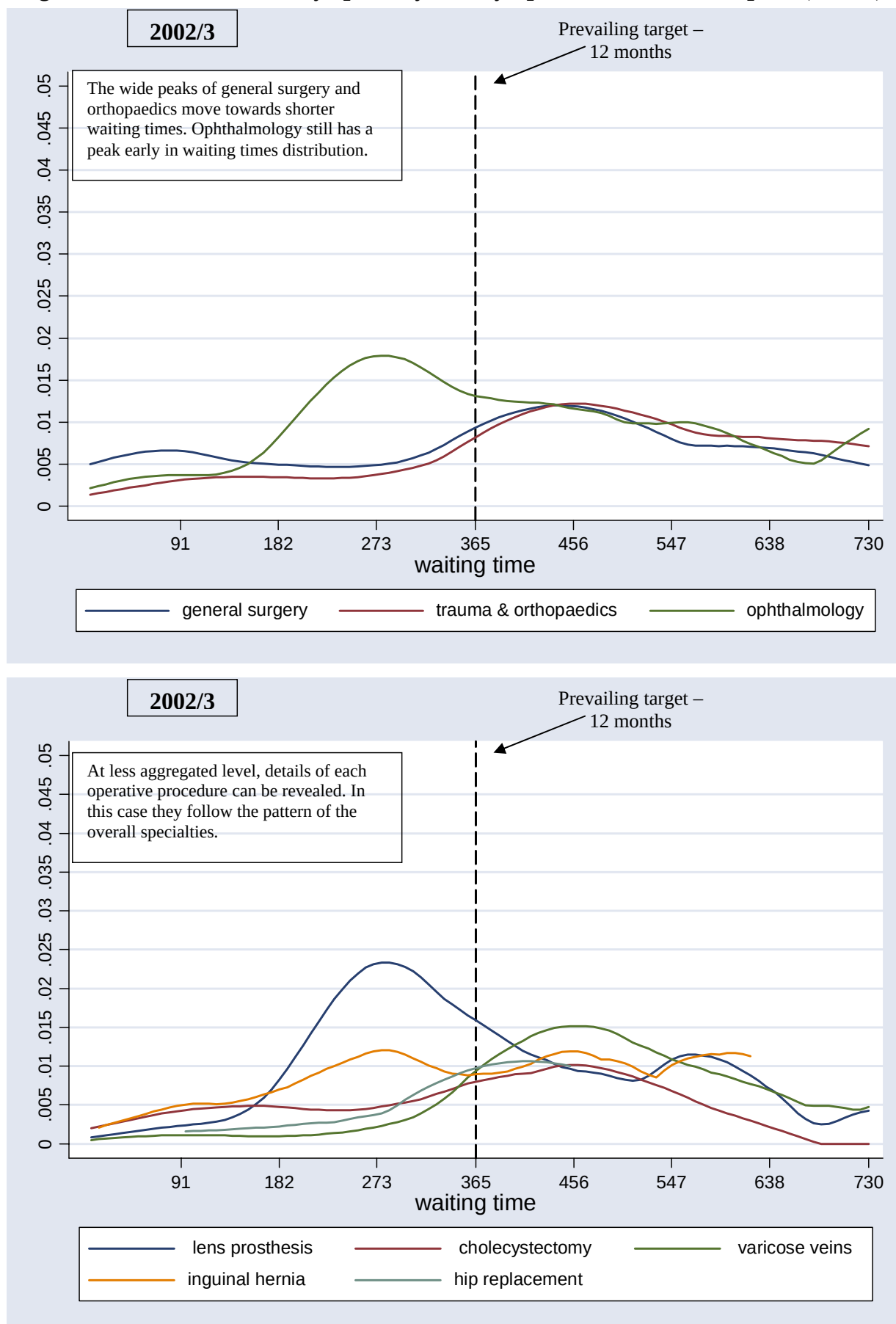


Figure 16: Waiting list management for five operations by Guys

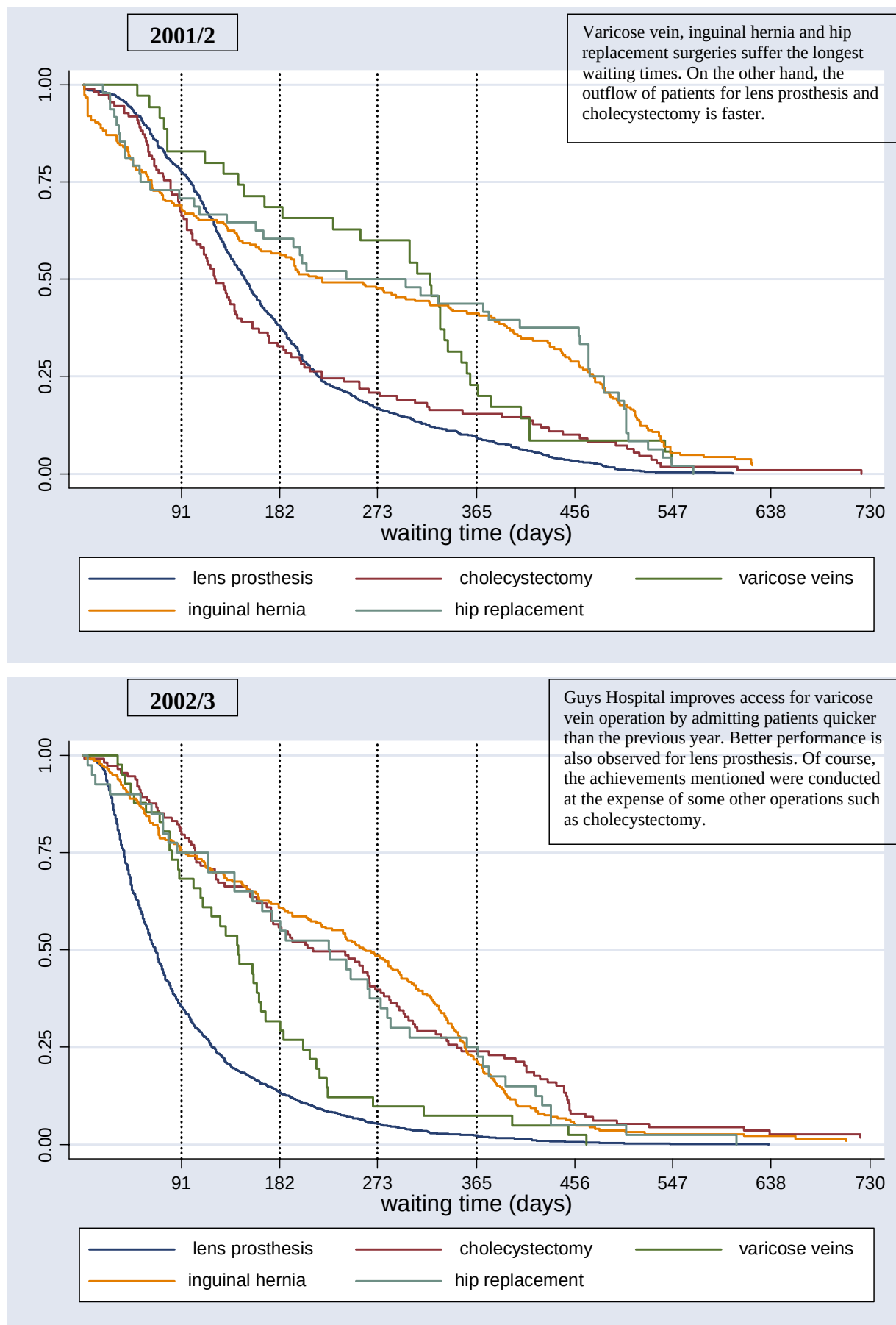


Figure 17: Hazard rates for five operations in Guys

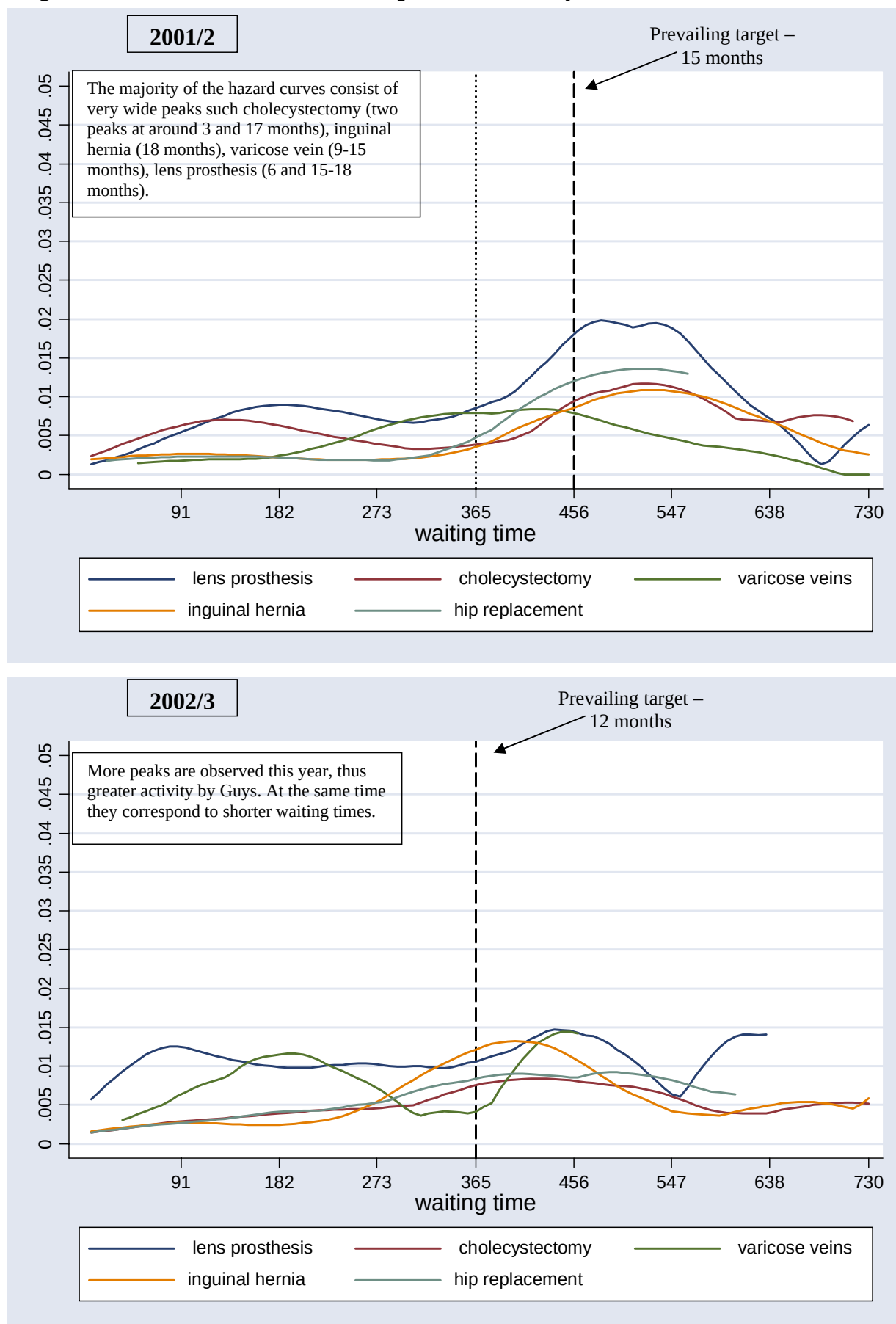


Figure 18: Different waiting list outflow in the trusts of Manchester and Newcastle

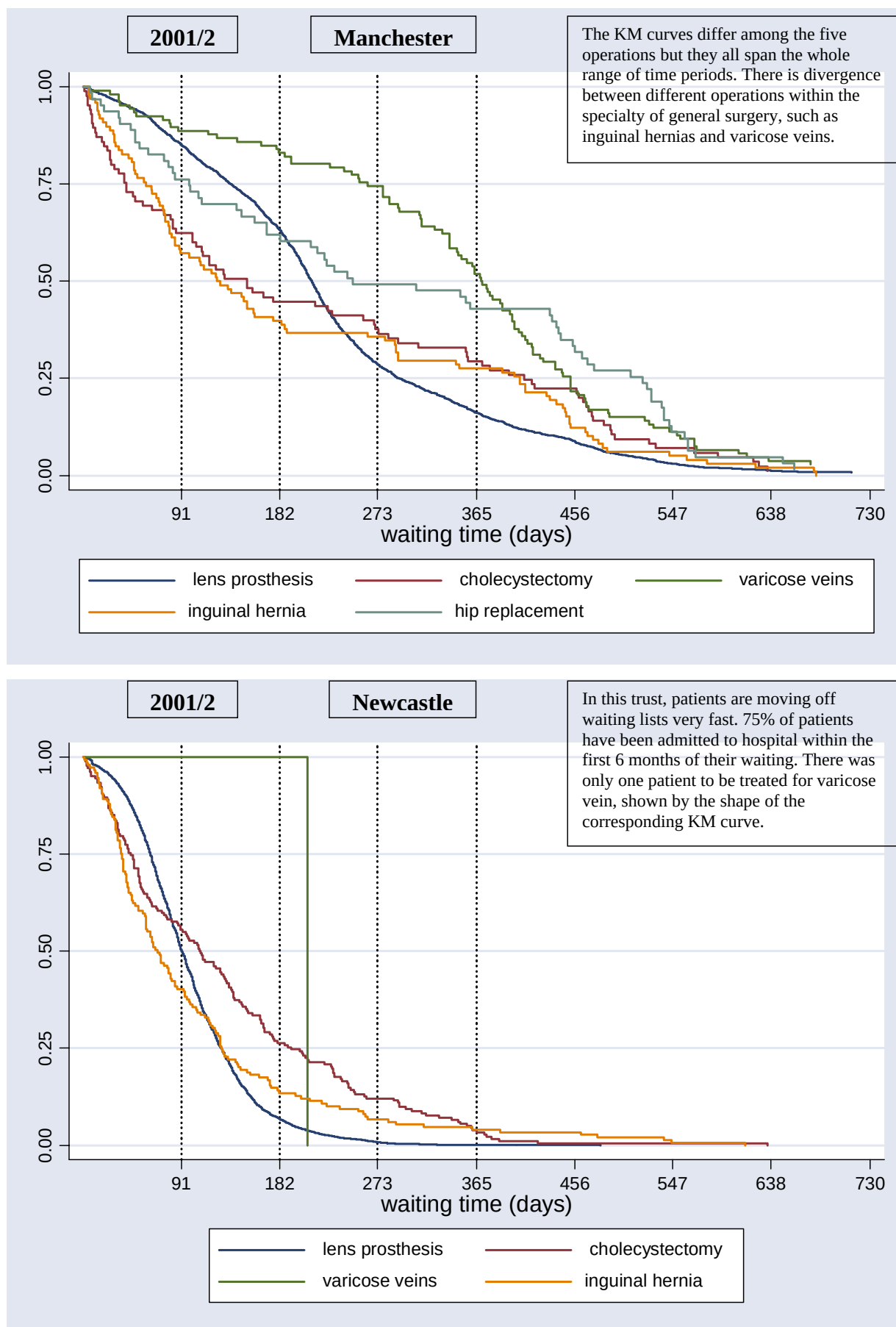


Figure 19: Survival curves by operation in Manchester and Newcastle (2002/3)

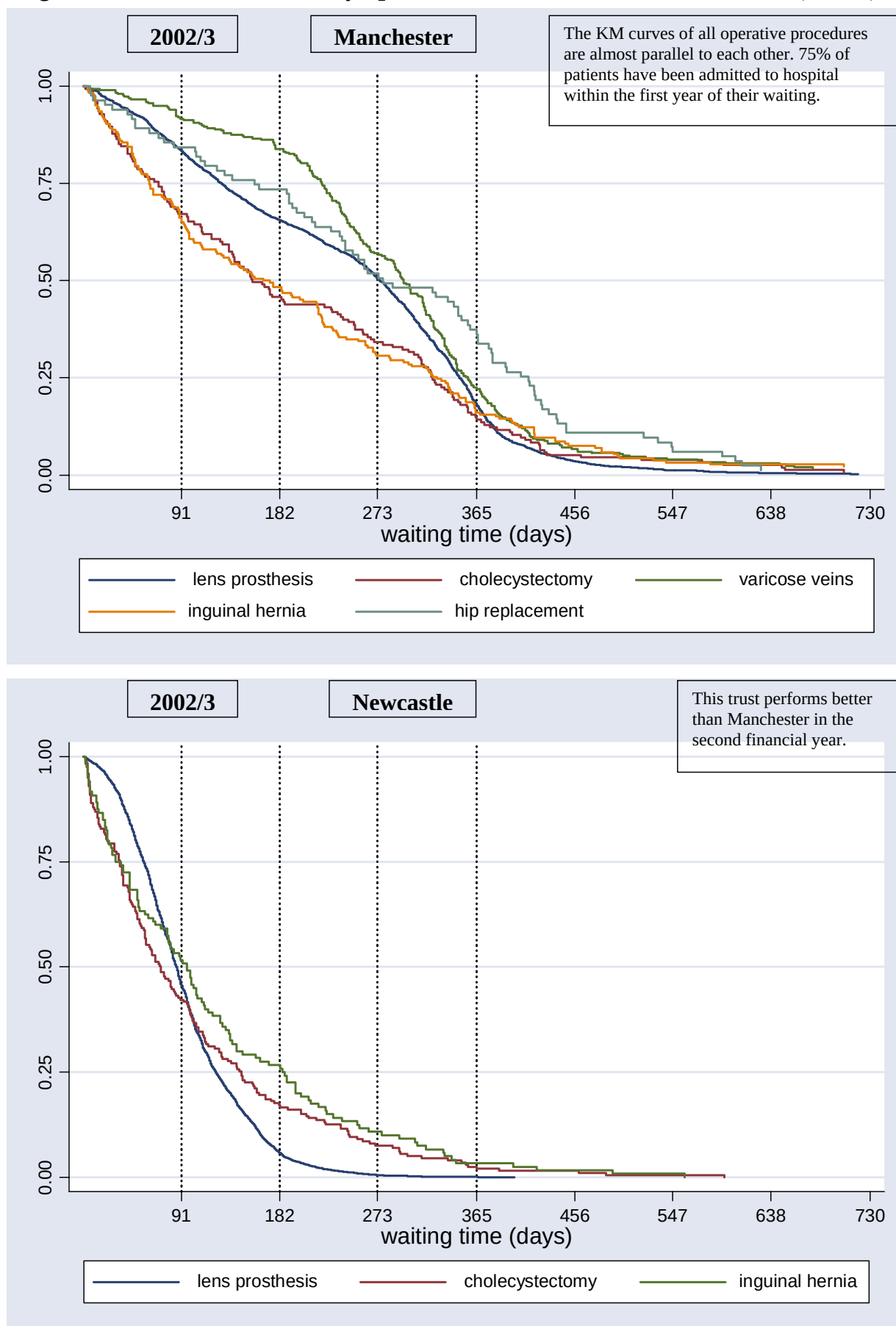


Figure 20 Hazard rates for different operations suggest greater activity by Newcastle

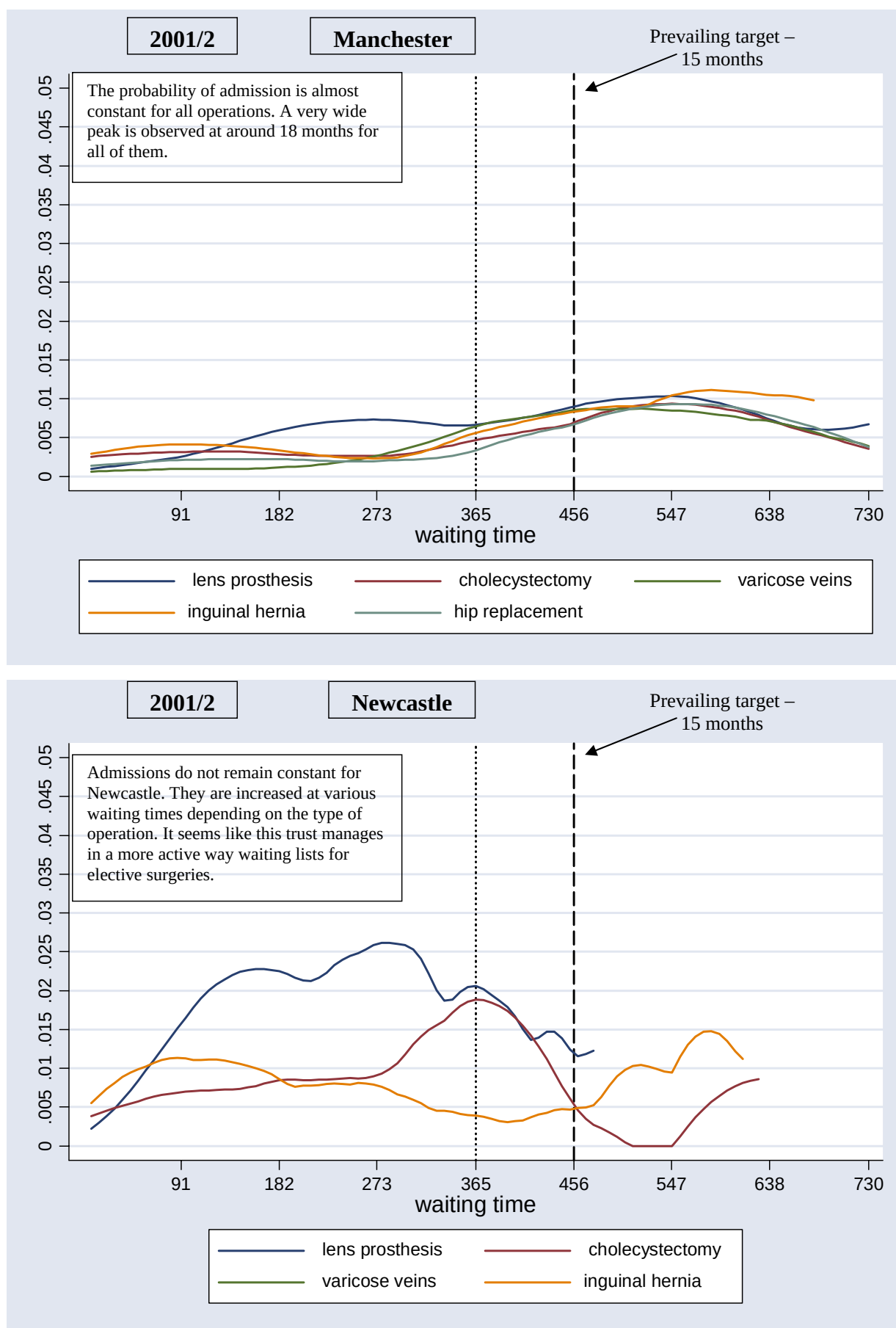


Figure 21: Hazard rates for five operations in Manchester and Newcastle (2002/3)

