UK Renal Registry 17th Annual Report: Chapter 11 Centre Variation in Access to Renal Transplantation in the UK (2008–2010)

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Key Words

Centre variation · Comorbidity · Donor after brainstem death · Donor after cardiac death · Equity of access · Living kidney donor · Outcomes · Patient factors · Quality improvement · Renal transplantation · Transplant waiting list

Abstract

Background: Renal transplantation is recognised as being the optimal treatment modality for many patients with established renal failure. This analysis aimed to explore inter-centre variation in access to renal transplantation in the UK. **Methods:** Transplant activity and waiting list data were obtained from NHS Blood and Transplant, demographic and laboratory data were obtained from the UK Renal Registry. All incident RRT patients starting treatment between 1st January 2008 and 31st December 2010 from

71 renal centres were considered for inclusion. The cohort was followed until 31st December 2012 (or until transplantation or death, whichever was earliest). **Results:** Age, ethnicity and primary renal diagnosis were associated with both accessing the kidney transplant waiting list and receiving a kidney transplant. A patient starting dialysis in a non-transplanting renal centre was less likely to be registered for transplantation (OR 0.74, 95% CI 0.68–0.81) or receive a transplant from a donor after cardiac death or a living kidney donor (OR 0.75, 95% CI 0.67–0.84) compared with patients cared for in transplanting renal centres. Once registered for kidney transplantation, patients in both transplanting and non-transplanting renal centres had an equal chance of receiving a transplant from a donor after brainstem death (OR 0.93, 95% CI 0.78 to 1.10). **Conclusion:** There was wide variation in access to kidney transplantation between UK renal centres which cannot be explained by differences in case mix.

Introduction

For 'suitable' patients with established renal failure, renal transplantation confers both better quality of life and life expectancy than dialysis [1–3] and is the preferred modality of renal replacement therapy. Achieving prompt and timely activation on the waiting list is important not least because increasing length of time on dialysis adversely affects graft and patient survival, but also because the current organ allocation algorithm introduced in April 2006 takes time spent on the waiting list into account when allocating deceased donor kidneys in the UK [4]. Thus, centres that achieve earlier listing for transplantation provide an advantage for their patients compared with centres that take longer.

This analysis aims to evaluate whether equity of access to the renal transplant list exists for patients with end stage renal disease across the UK, whether centres differ in the time taken to activate suitable patients on the waiting list and whether equity exists in the receipt of a renal transplant once the patient is on the transplant list (that is, the conversion efficiency from being on the waiting list to receiving a transplant). Patient specific and independent variables that influenced access to the waiting list or transplantation were analysed.

Methods

Study population

All adult patients starting renal replacement therapy ($N = 20,076$) between 1st January 2008 and 31st December 2010 in renal centres ($N = 71$) returning data to the UK Renal Registry (UKRR) were considered for inclusion. For the analysis of the proportion of a centre's patients included on the waiting list, patients aged 65 years or above ($n = 9,908$), patients with inappropriate activation and early suspension as described below ($n = 55$) and patients listed for multi-organ transplants other than pancreas ($n = 34$) were excluded, resulting in a final cohort of 10,079 patients. These patients were followed to 31st December 2012 or until they were put on the waiting list for kidney transplant alone, kidney plus pancreas transplant, or death, whichever was earliest. For the analysis of the proportion transplanted, all patients from the incident cohort who were activated on the waiting list before 31st December 2011 ($N = 5,239$) were followed until 31st December 2013, to estimate the proportion transplanted with a kidney alone or kidney plus pancreas within two years of inclusion on the waiting list.

Exclusions

Patients listed for multi-organ transplants other than pancreas were excluded as were those who were suspended for more than 30 days within 90 days of first activation. The latter avoided any

potential bias from centres that may activate patients on the transplant list and then immediately suspend them before more permanent activation at a later date after more formal medical assessment of the patient's fitness.

Data analysed

Information on start date of renal replacement therapy and relevant patient level data including age (grouped as 18–29, 30–39, 40–49, 50–59, 60–64), gender, ethnicity (White, non-White, missing) and (primary renal diagnosis (PRD) classified as: patient with diabetes, patient without diabetes, missing) came from the UKRR. The date of activation on the kidney transplant waiting list, date of transplantation, or both came from the UK Transplant Registry held by the Organ Donation and Transplantation Directorate of NHS Blood and Transplant.

Statistical methods

A logistic regression model was developed to identify the influence of patient specific variables including age, gender, ethnicity and PRD, on the probability of access to the transplant list and receipt of a transplant once on the waiting list. After adjusting for patient specific variables, the percentage of patients activated on the transplant list and the percentage of patients on the waiting list who achieved a transplant in each centre was determined. The overall effect of the centre associated with each analysis was assessed by including renal centre as a random effect in the risk-adjusted logistic regression model. The extent of variation between centres was determined by using a log likelihood ratio test that provided the change in the value of -2 LogL on inclusion of the random centre effect. SAS 9.4 was used for analyses; a p value of less than 5% was considered significant.

To analyse access to the transplant list, the proportion of incident patients with end stage renal disease in each centre who were subsequently activated on the waiting list within two years of starting renal replacement therapy (RRT) was identified. All patients who achieved live donor transplantation without prior activation on the national transplant waiting list were assumed to have been activated for the purposes of this analysis. Time to activation on the waiting list was defined as the interval between the start of RRT and the date of activation on the waiting list. Patients achieving pre-emptive deceased donor transplantation were considered to have been activated on the same day as starting RRT i.e. a time to activation of zero days. Patients achieving pre-emptive live donor transplantation without prior activation on the national transplant list were considered to have been 'active' on the list for an arbitrary time of six months. This was to take into account an average of six months required by most centres to complete live donor fitness evaluation and hence the likelihood that the intended recipient was considered fit for transplantation (and by inference suitable to be active on the waiting list) for that duration. This was done to account for different centre practices with regard to listing patients on the deceased donor list prior to receiving a living donor transplant.

The median time to activation was estimated from the Kaplan-Meier plot for patients at each renal centre, with the event as the date of activation and censoring at death or on 31st December 2012, whichever was earlier. Data from patients who did not achieve activation were included in the calculation of median times using this method, thus providing a meaningful estimate of the true time to activation. Including only those patients activated would produce

a biased estimate. The overall centre effect associated with time to activation was calculated by including renal centre as a variable in a risk-adjusted Cox regression model.

To analyse the difference between centres in renal transplantation once listed, the percentage of patients activated on the waiting list who received a renal transplant within two years of being activated was estimated (conversion efficiency). The conversion efficiency for receiving a transplant from a donor after brainstem death or a donor after cardiac death/living kidney donor was analysed separately. Receipt of a kidney from a donor after brainstem death was predominantly influenced by national allocation policy, whereas receipt from a donor after cardiac death/live donor kidney was much more dependent on local transplant centre practices. For the cohort under consideration, donor after cardiac death transplantation was predominantly a locally managed service.

Funnel plots are used to present the results for each outcome of interest, providing a visual comparison of each centre's performance compared with its peers. Where relevant, the funnel plots are adjusted for patient specific variables influencing that outcome. The solid black straight line in each funnel plot shows the overall average together with the 95% and 99.8% confidence intervals, which correspond to two and three standard deviations from the mean. Each point on the plot represents one renal centre. With 71 centres included, for each outcome of interest, three centres would be predicted to fall between the 95% and 99.8% confidence intervals and no centre should fall outside the 99.8% confidence interval. Centres ($n = 3$) with fewer than 10 patients activated on the waiting list are not included in the funnel plots.

The analysis methodology described above is identical to a previous independent peer reviewed publication [5].

Results

The results of the logistic regression model analysis of patient characteristics influencing access to the waiting

list are presented in table 11.1. Ethnicity data were missing for 11% of patients and PRD for 6% of patients.

Tables 11.2 and 11.3 show the results of the logistic regression analysis of factors influencing the likelihood of receiving a transplant from a donor after brainstem death and the analysis of factors influencing receipt of a transplant from a donor after cardiac death or a living kidney donor. Ethnicity data were missing for 8.1% of patients and PRD for 6.1% of patients (table 11.3).

A patient starting dialysis in a non-transplanting renal centre was less likely to be registered for transplantation (OR 0.74, 95% CI 0.68–0.81) or receive a transplant from a donor after cardiac death or a living kidney donor (OR 0.75, 95% CI 0.67–0.84) compared with patients managed in transplanting renal centres. Once registered for kidney transplantation, patients in both transplanting and non-transplanting renal centres had an equal chance of receiving a transplant from a donor after brainstem death (OR 0.93, 95% CI 0.78–1.10).

After adjusting for patient specific variables that were shown to influence outcome (age, ethnicity, gender, PRD), significant centre effects were identified for the probability of being activated on the waiting list (figure 11.1, table 11.4) (change in $-2 \log L = 188.6$, df (degrees of freedom) = 1, $p < 0.0001$).

After adjustment for patient variables, significant centre differences were seen in the probability of receiving a renal transplant from a donor after brain stem death (figure 11.2, table 11.5) (change in $-2 \log L = 13.3$, df = 1, $p = 0.0003$) or a donor after cardiac death/living kidney donor (figure 11.3, table 11.5) (change in $-2 \log L = 136.1$, df = 1,

Table 11.1. Patient factors influencing activation on the national kidney transplant waiting list within two years of RRT start

Factor	Category (at baseline)	Patients N (%)	Odds ratio	95% CI	P value
Age	(18–29)	899 (8.9)	1	ref	n/a
	30–39	1,425 (14.1)	1.06	0.88–1.27	0.57
	40–49	2,400 (23.8)	0.70	0.59–0.83	<0.0001
	50–59	3,239 (32.1)	0.36	0.31–0.43	<0.0001
	60–64	2,116 (21.0)	0.20	0.17–0.24	<0.0001
Ethnicity	(White)	6,697 (66.5)	1	ref	n/a
	Non-White	2,274 (22.6)	0.80	0.72–0.89	<0.0001
	Missing	1,108 (11.0)	0.50	0.43–0.57	<0.0001
Gender	(Male)	6,086 (60.4)	1	ref	n/a
	Female	3,993 (39.6)	0.95	0.88–1.04	0.28
PRD	(Non-diabetic)	7,017 (69.6)	1	ref	n/a
	Diabetic	2,455 (24.4)	0.44	0.40–0.49	<0.0001
	Missing	607 (6.0)	0.84	0.71–1.00	0.06

ref – reference category, n/a – not applicable

Table 11.2. Patient factors affecting the probability of receiving a transplant from a donor after brainstem death within two years of registration on the national kidney transplant waiting list

Factor	Category (at baseline)	Patients N (%)	Odds ratio	95% CI	P value
Age	(18–29)	646 (12.3)	1	ref	n/a
	30–39	1,014 (19.4)	1.05	0.80–1.38	0.7
	40–49	1,457 (27.8)	0.78	0.60–1.01	0.06
	50–59	1,444 (27.6)	0.40	0.30–0.54	<0.0001
	60–64	678 (12.9)	0.23	0.15–0.35	<0.0001
Ethnicity	(White)	3,639 (69.5)	1	ref	n/a
	Non-White	1,176 (22.5)	0.65	0.52–0.81	0.0001
	Missing	424 (8.1)	1.05	0.78–1.40	0.7
Gender	(Male)	3,180 (60.7)	1	ref	n/a
	Female	2,059 (39.3)	1.32	1.11–1.56	0.002
PRD	(Non-diabetic)	4,045 (77.2)	1	ref	n/a
	Diabetic	874 (16.7)	3.79	3.13–4.57	<0.0001
	Missing	320 (6.1)	1.04	0.71–1.52	0.8

ref – reference category, n/a – not applicable

$p < 0.0001$). As shown, several centres fall outside the 95% and 99.8% confidence intervals.

Figure 11.4 and table 11.6 show the unadjusted median time taken to activate patients on the transplant list for each renal centre.

The funnel plot is based on the assumption of an exponential distribution for time to activation. Although this assumption is broadly consistent with the data, the model based estimate of the national median was greater than that observed. This leads to an unusually large number of centres falling outside the lower 99.8% confidence

limit for this national rate and perhaps too few occurring outside the upper limit. However, the plot highlights those centres that have significantly longer time to activation but small numbers on the waiting list. The Cox model giving a risk-adjusted analysis of time to activation identified a significant effect of centre variation (change in $-2 \log L = 372.7$, $df = 70$, $p < 0.0001$). In general, centres with the longest unadjusted waiting times also had the longest risk-adjusted waiting times. The three centres lying outside the upper 99.8% confidence limit all had hazard ratios that indicated a significant delay in the

Table 11.3. Patient factors affecting the probability of receiving a transplant from a donor after cardiac death or living kidney donor within two years of registration on the national kidney transplant waiting list

Factor	Category (at baseline)	Patients N (%)	Odds ratio	95% CI	P value
Age	(18–29)	646 (12.3)	1	ref	n/a
	30–39	1,014 (19.4)	0.64	0.53–0.79	<0.0001
	40–49	1,457 (27.8)	0.57	0.47–0.70	<0.0001
	50–59	1,444 (27.6)	0.51	0.42–0.62	<0.0001
	60–64	678 (12.9)	0.5	0.40–0.63	<0.0001
Ethnicity	(White)	3,639 (69.5)	1	ref	n/a
	Non-White	1,176 (22.5)	0.51	0.44–0.59	<0.0001
	Missing	424 (8.1)	0.54	0.43–0.67	<0.0001
Gender	(Male)	3,180 (60.7)	1	ref	n/a
	Female	2,059 (39.3)	0.77	0.68–0.86	<0.0001
PRD	(Non-diabetic)	4,045 (77.2)	1	ref	n/a
	Diabetic	874 (16.7)	0.37	0.31–0.44	<0.0001
	Missing	320 (6.1)	1.42	1.13–1.80	0.003

ref – reference category, n/a – not applicable

Fig. 11.1. Percentage of patients wait-listed for a kidney transplant by renal centre, prior to or within two years of starting dialysis (centres with <10 patients excluded)

Table 11.4. Percentage of patients wait-listed for a kidney transplant by renal centre, prior to or within two years of starting dialysis

England					Scotland				
	RRT	Registrations	% wait listed			RRT	Registrations	% wait listed	
Centre	N	N	Unadjusted	Risk-adjusted	Centre	N	N	Unadjusted	Risk-adjusted
B Heart	143	65	45.5	44.4	Plymth	96	64	66.7	65.9
B QEH	370	181	48.9	48.3	Ports	231	154	66.7	65.1
Basldn	37	10	27.0	25.2	Prestn	221	98	44.3	44.4
Bradfd	111	51	45.9	44.7	Redng	157	109	69.4	68.8
Brightn	135	62	45.9	44.3	Salford	252	145	57.5	56.7
Bristol	257	130	50.6	47.7	Sheff	237	114	48.1	46.7
Camb	153	91	59.5	56.8	Shrew	61	16	26.2	28.4
Carlisle	41	27	65.9	60.4	Stevng	177	97	54.8	52.5
Carsh	270	114	42.2	41.3	Sthend	44	25	56.8	58.2
Chelms	62	35	56.5	58.4	Stoke	120	60	50.0	48.3
Colchr	41	23	56.1	55.9	Sund	101	50	49.5	47.8
Covnt	169	79	46.7	45.3	Truro	58	35	60.3	60.9
Derby	108	41	38.0	39.4	Wirral	89	26	29.2	28.7
Donc	50	30	60.0	64.2	Wolve	131	47	35.9	34.7
Dorset	81	47	58.0	57.4	York	50	27	54.0	50.9
Dudley	69	19	27.5	27.5	N Ireland				
Exeter	149	62	41.6	42.9	Antrim	40	14	35.0	35.0
Glouc	82	45	54.9	54.7	Belfast	93	43	46.2	45.5
Hull	139	60	43.2	49.5	Newry	28	13	46.4	49.5
Ipswi	54	30	55.6	52.2	Ulster	16	7	43.8	50.4
Kent	175	103	58.9	56.2	West NI	38	21	55.3	49.2
L Barts	441	206	46.7	46.4	Scotland				
L Guys	298	168	56.4	54.2	Abrdn	88	46	52.3	64.9
L Kings	234	101	43.2	43.3	Airdrie	78	28	35.9	47.7
L Rfree	308	165	53.6	52.2	D&Gall	18	10	55.6	91.6
L St.G	158	91	57.6	55.9	Dundee	62	25	40.3	55.6
L West	568	346	60.9	62.9	Edinb	137	63	46.0	58.1
Leeds	225	120	53.3	51.4	Glasgw	257	127	49.4	64.3
Leic	346	216	62.4	61.4	Inverns	37	15	40.5	41.3
Liv Ain	60	16	26.7	28.1	Klmarnk	49	13	26.5	37.8
Liv Roy	177	96	54.2	49.3	Krkldy	41	14	34.1	46.4
M RI	246	135	54.9	51.6	Wales				
Middlbr	132	79	59.8	56.1	Bangor	37	9	24.3	27.3
Newc	162	85	52.5	50.9	Cardff	243	128	52.7	51.4
Norwch	89	46	51.7	51.2	Clwyd	22	5	22.7	23.4
Nottm	196	115	58.7	55.1	Swansea	150	56	37.3	37.1
Oxford	259	170	65.6	63.0	Wrexham	25	11	44.0	41.2

Fig. 11.2. Percentage of patients receiving a transplant from a donor after brainstem death by renal centre, within two years of transplant waiting list registration (centres with <10 patients excluded)

chance of wait-listing compared with a baseline centre that had a median time comparable to the national median.

Discussion

Centre variation

The analyses performed within this report highlight a significant centre effect in relation to the proportion of patients wait-listed with nearly 20% of centres lying outside the lower 95% confidence interval, and seven centres outside the lower 99.8% confidence interval, despite adjusting for a range of patient characteristics. Inter-centre differences are also noted in access to transplants from donors after cardiac death/living kidney donors with nine centres lying outside the lower 99.8% confidence interval.

Whilst both these outcomes are subject to individual centre practices and policies (which thus could be deemed a cause of the observed variation), one needs to interpret these results with caution as this study is limited by the lack comprehensive comorbidity data on all patients. Centres with higher prevalence rates of comorbidities would be expected to list proportionally fewer patients to reflect the fact that fewer patients are fit for transplantation. Additionally, it may take longer to activate patients in these centres due to the need for more intensive investigation and medical optimisation prior to transplantation. Indeed lack of comorbidity data limits definitive adjustment for case mix. Other patient level factors which this study also fails to adjust for include social deprivation which has been associated with

reduced access to transplantation of a range of organs, as well as the impact of primary renal diagnoses (other than diabetes), health literacy and human leucocyte antigen (HLA) sensitisation. Additionally, this study has not analysed the interplay between factors such as social deprivation and ethnicity and whether the observed differences based on ethnicity are likely to persist after adjustment for social deprivation and varying comorbidity burden in different ethnic groups. In essence, the available dataset does not permit definitive adjustment for case mix.

The observation that a patient starting dialysis in a non-transplanting renal centre was less likely to be registered for transplantation or receive a transplant from a donor after cardiac death (or a living kidney donor) compared with patients managed in transplanting renal centres, has been a consistent finding in similar analyses by the UKRR over the last five years. The finding implies increasing distance from the transplanting centre diminishes access to treatment. Despite the inability to conclusively adjust for case mix in our analyses, the consistent finding in the same direction, suggests centre practice patterns rather than patient dependant variables may explain this finding. Further detailed analyses to understand the reasons for such variability is currently being undertaken as part of the Access to Transplant and Transplant Outcome Measures (ATTOM) study. Once registered for kidney transplantation, patients in both transplanting and non-transplanting renal centres had an equal chance of receiving a transplant from a donor after brainstem death. This is reassuring as organ allocation is subject to the national allocation algorithm which one would expect to allocate organs equitably.

Table 11.5. Percentage of patients receiving a transplant, by donor type and renal centre, within two years of transplant waiting list registration

Centre	Organ from donor after brainstem death				Organ from living kidney donor/donor after cardiac death		
	Listed <i>N</i>	Transplanted <i>N</i>	Transplant rate (%)		Transplanted <i>N</i>	Transplant rate (%)	
			Unadjusted	Risk-adjusted		Unadjusted	Risk-adjusted
England							
B Heart	66	4	6.1	6.6	15	22.7	23.9
B QEH	186	11	5.9	6.4	56	30.1	31.2
Basldn	13	4	30.8	21.4	4	30.8	32.1
Bradfd	54	9	16.7	16.1	10	18.5	20.5
Brightn	63	10	15.9	15.1	16	25.4	23.8
Bristol	130	11	8.5	8.3	59	45.4	42.3
Camb	91	11	12.1	12.7	62	68.1	61.5
Carlis	27	5	18.5	18.3	16	59.3	50.3
Carsh	116	10	8.6	8.8	58	50.0	47.4
Chelms	35	5	14.3	15.3	19	54.3	53.7
Colchr	23	1	4.3	4.2	11	47.8	43.7
Covnt	79	7	8.9	9.7	47	59.5	56.3
Derby	42	3	7.1	7.4	3	7.1	6.6
Donc	30	3	10.0	9.3	3	10.0	9.6
Dorset	48	2	4.2	4.8	15	31.3	27.7
Dudley	19	2	10.5	10.2	2	10.5	10.5
Exeter	62	7	11.3	10.6	24	38.7	38.4
Glouc	45	7	15.6	19.2	11	24.4	23.0
Hull	61	4	6.6	6.9	22	36.1	36.1
Ipswi	30	5	16.7	17.3	18	60.0	51.9
Kent	105	12	11.4	11.7	53	50.5	46.6
L Barts	208	21	10.1	11.6	91	43.8	50.1
L Guys	171	26	15.2	15.1	99	57.9	58.8
L Kings	102	8	7.8	8.6	30	29.4	31.6
L Rfree	168	16	9.5	12.4	80	47.6	43.2
L St.G	91	6	6.6	7.1	47	51.6	53.0
L West	349	51	14.6	15.7	153	43.8	52.3
Leeds	122	21	17.2	17.7	49	40.2	38.7
Leic	216	19	8.8	9.9	85	39.4	38.5
Liv Ain	17	3	17.6	14.8	10	58.8	49.9
Liv Roy	96	13	13.5	12.2	40	41.7	39.0
M RI	136	25	18.4	16.6	53	39.0	39.5
Middlbr	81	14	17.3	15.9	49	60.5	56.5
Newc	86	16	18.6	18.2	49	57.0	53.4
Norwch	46	6	13.0	14.4	25	54.3	50.1
Nottm	115	28	24.3	24.4	38	33.0	31.3
Oxford	170	36	21.2	19.4	74	43.5	43.7
Plymth	64	7	10.9	10.9	29	45.3	42.4
Ports	156	24	15.4	13.6	51	32.7	32.2
Prestn	100	11	11.0	11.3	32	32.0	29.5
Redng	109	19	17.4	17.6	39	35.8	38.7
Salford	144	14	9.7	10.8	47	32.6	31.1
Sheff	116	5	4.3	4.4	47	40.5	38.1
Shrew	17	1	5.9	7.4	7	41.2	38.1
Stevng	100	13	13.0	12.4	51	51.0	49.8
Sthend	25	7	28.0	26.9	8	32.0	33.9
Stoke	62	6	9.7	10.1	26	41.9	37.8
Sund	50	5	10.0	9.4	32	64.0	61.1
Truro	36	2	5.6	5.4	20	55.6	53.0
Wirral	27	3	11.1	8.7	11	40.7	38.3
Wolve	47	5	10.6	9.5	14	29.8	29.1
York	27	5	18.5	13.7	9	33.3	33.4

Table 11.5. Continued

Centre	Organ from donor after brainstem death				Organ from living kidney donor/donor after cardiac death		
	Listed <i>N</i>	Transplanted <i>N</i>	Transplant rate (%)		Transplanted <i>N</i>	Transplant rate (%)	
			Unadjusted	Risk-adjusted		Unadjusted	Risk-adjusted
N Ireland							
Antrim	14	1	7.1	7.0	5	35.7	30.5
Belfast	44	3	6.8	5.9	14	31.8	28.6
Newry	13	1	7.7	5.8	1	7.7	7.6
Ulster	7	0	0.0	0.0	1	14.3	14.6
West NI	21	3	14.3	16.5	5	23.8	20.5
Scotland							
Abrdn	46	7	15.2	13.8	12	26.1	31.7
Airdrie	29	7	24.1	17.5	5	17.2	23.0
D&Gall	10	2	20.0	13.9	1	10.0	17.5
Dundee	26	5	19.2	14.8	7	26.9	36.6
Edinb	66	8	12.1	10.6	29	43.9	55.9
Glasgw	130	19	14.6	12.3	40	30.8	40.5
Inverns	16	3	18.8	16.8	1	6.3	5.8
Klmarnk	14	4	28.6	26.1	1	7.1	9.0
Krkldy	14	1	7.1	5.4	2	14.3	19.9
Wales							
Bangor	9	5	55.6	70.1	2	22.2	20.4
Cardff	128	24	18.8	18.2	49	38.3	35.4
Clwyd	5	0	0.0	0.0	2	40.0	33.3
Swanse	57	11	19.3	15.9	28	49.1	48.2
Wrexm	11	3	27.3	22.9	4	36.4	31.3

Patient level factors affecting access

The finding that certain patient variables, such as increasing age, have a negative association with access to transplantation was not unexpected as the risk-benefit ratio of receiving a renal transplant alters with age. Increased comorbidity burden in older patients may require more intensive time consuming investigations prior to listing and may also deem them unsuitable in some cases. In this study 'non-White' ethnicity was

negatively associated with access to transplantation and receiving a kidney once listed (from a donor after brainstem death, and from a living kidney donor/donor after cardiac death). This may partly be explained by the importance given to HLA matching in the national allocation protocol which may have favoured a predominantly white donor pool being matched with white recipients and also the widely acknowledged lack of donors from ethnic minorities contributing to the

Fig. 11.3. Percentage of patients receiving a transplant from a living kidney donor/donor after cardiac death by renal centre, within two years of transplant waiting list registration (centres with <10 patients excluded)

Fig. 11.4. Median time to wait listing for a kidney transplant, by renal centre (centres with <10 patients excluded)

Table 11.6. Median time to wait listing for a kidney transplant, by renal centre (censoring at the earliest of death or 31st December 2012)

Centre	RRT N	Registrations N	Median time to listing (days)	Centre	RRT N	Registrations N	Median time to listing (days)
England				Plymth	96	64	157
B Heart	143	66	867*	Ports	231	156	91
B QEH	370	186	598	Prestn	221	100	797*
Basldn	37	13	923*	Redng	157	110	174
Bradfd	111	54	773	Salford	252	145	380
Brightn	135	63	811*	Sheff	237	116	769
Bristol	257	130	436	Shrew	61	17	787*
Camb	153	92	128	Stevng	177	100	302
Carlis	41	27	36	Sthend	44	25	215
Carsh	270	117	1,345*	Stoke	120	62	488
Chelms	62	35	309	Sund	101	50	413
Colchr	41	23	226	Truro	58	36	223
Covnt	169	79	712*	Wirral	89	27	835*
Derby	108	42	1,017*	Wolve	131	48	765*
Donc	50	30	147	York	50	27	472
Dorset	81	48	243	N Ireland			
Dudley	69	19	641*	Antrim	40	14	495*
Exeter	149	63	1,225*	Belfast	93	44	821
Glouc	82	45	340	Newry	28	13	683*
Hull	139	62	1,049*	Ulster	16	7	180*
Ipswi	54	30	438	West NI	38	21	397
Kent	175	105	240	Scotland			
L Barts	441	208	816*	Abrdn	88	46	365
L Guys	298	171	291	Airdrie	78	30	816*
L Kings	234	102	742*	D&Gall	18	10	335
L Rfree	308	168	364	Dundee	62	26	838*
L St.G	158	91	216	Edinb	137	66	741
L West	568	350	220	Glasgw	257	130	548
Leeds	225	122	281	Inverns	37	16	735*
Leic	346	216	116	Klmarnk	49	14	779*
Liv Ain	60	17	967*	Krkldy	41	14	258*
Liv Roy	177	98	508	Wales			
M RI	246	138	416	Bangor	37	9	716*
Middlbr	132	81	304	Cardff	243	128	311
Newc	162	86	348	Clwyd	22	5	682*
Norwch	89	46	440	Swanse	150	57	870*
Nottm	196	115	280	Wrexm	25	11	564*
Oxford	259	171	115				

*Results in **bold italics** are final event times as median times could not be estimated

donor pool. Lack of adjustment for social deprivation may also explain some of this association, as Udayaraj has previously published data showing that adjusting for social deprivation may partly explain the reduced access to transplantation seen in ethnic groups [6].

Diabetes was also seen to affect wait-listing adversely although this is not surprising as many would be subject to additional diabetic complications and increased cardiovascular risk that would need to be managed. The higher proportion of patients with diabetes receiving a transplant corresponds to an increase in the number of simultaneous kidney-pancreas transplants during the study period, as the allocation algorithm prioritised dual organ recipients.

When interpreting the analyses in this chapter it is important to consider the potential impact of missing data on the results. Missing data occurs as a result of either a renal centre failing to complete relevant fields on their renal IT system or a failure to extract this data. Missing data may not be at random; sicker patients may die more quickly, allowing inadequate time for their physician to enter relevant comorbidity data. The very process of working up and listing a patient makes it less likely that data will be missing. It is therefore perhaps not surprising that patients activated on the national kidney transplant waiting list are more likely to have ethnicity and PRD data reported ($p < 0.0001$) (table 11.1).

The UKRR is collaborating with other researchers in the National Institute for Health Research (NIHR) funded ATTOM research project to study access to kidney transplantation in greater detail. This will allow those practices identified in the better performing centres to be disseminated to other centres, thereby facilitating equity of access to transplantation across the UK.

Conclusions

This study highlights the persistence of significant centre variation in access to transplantation with respect to the proportion of patients listed and the time taken to activate suitable patients, even after correction for available relevant patient related variables. Significant differences exist between transplanting and non-transplanting centres, with increasing age, non-White ethnicity and diabetes showing a negative association in terms of accessing the transplant wait-list.

Conflicts of interest: none

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