

# **Cancer Patient Experience Survey**

2023 Results

## **North East London Cancer Alliance**

First published July 2024; updated November 2024 to include Integrated Care Board data

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## Executive summary

### Questions above expected range

North East London Cancer Alliance has no scores above expected range.

### Questions below expected range

	Case mix adjusted scores			England score
	2023 score	Lower expected range	Upper expected range	
Q3. Referral for diagnosis was explained in a way the patient could completely understand	<b>61%</b>	62%	71%	<b>67%</b>
Q5. Patient received all the information needed about the diagnostic test in advance	<b>89%</b>	90%	94%	<b>92%</b>
Q6. Diagnostic test staff appeared to completely have all the information they needed about the patient	<b>79%</b>	80%	87%	<b>83%</b>
Q7. Patient felt the length of time waiting for diagnostic test results was about right	<b>73%</b>	74%	81%	<b>78%</b>
Q19. Patient found advice from main contact person was very or quite helpful	<b>94%</b>	94%	97%	<b>96%</b>
Q25. A member of their care team helped the patient create a care plan to address any needs or concerns	<b>90%</b>	91%	96%	<b>94%</b>
Q26. Care team reviewed the patient's care plan with them to ensure it was up to date	<b>98%</b>	98%	100%	<b>99%</b>
Q28. Patient definitely got the right level of support for their overall health and well being from hospital staff	<b>71%</b>	71%	80%	<b>76%</b>
Q39. Patient was always able to discuss worries and fears with hospital staff while being treated as an outpatient or day case	<b>74%</b>	75%	84%	<b>79%</b>
Q41_5. Beforehand patient completely had enough understandable information about immunotherapy	<b>74%</b>	77%	90%	<b>84%</b>
Q43. Patient felt the length of waiting time at clinic and day unit for cancer treatment was about right	<b>64%</b>	72%	85%	<b>78%</b>
Q50. During treatment, the patient definitely got enough care and support at home from community or voluntary services	<b>43%</b>	46%	59%	<b>52%</b>
Q55. Patient was given enough information about the possibility and signs of cancer coming back or spreading	<b>58%</b>	58%	69%	<b>64%</b>

## Introduction

The National Cancer Patient Experience Survey 2023 is the thirteenth iteration of the survey first undertaken in 2010. It has been designed to monitor progress on cancer care; to provide information to drive local quality improvements; to assist commissioners and providers of cancer care; and to inform the work of the various charities and stakeholder groups supporting cancer patients.

The survey was undertaken by Picker on behalf of NHS England and it was overseen by a national Cancer Patient Experience Advisory Group. This Advisory Group set the principles and objectives of the survey programme and guided questionnaire development. The survey was commissioned and managed by NHS England. The survey provider, Picker, is responsible for designing, running and analysing the survey.

The 2023 survey involved 132 NHS trusts. Out of 121,121 people, 63,428 people responded to the survey, yielding a response rate of 52%.

This document was reissued in November 2024 to include an Integrated Care Board expected range summary section.

## Methodology

### Eligibility, fieldwork and survey methods

The sample for the survey included all adult (aged 16 and over) NHS patients, with a confirmed primary diagnosis of cancer, discharged from an NHS trust after an inpatient episode or day case attendance for cancer related treatment in the months of April, May and June 2023. The fieldwork for the survey was undertaken between November 2023 and February 2024.

As in the previous eight years, the survey used a mixed mode methodology. Questionnaires were sent by post, with two reminders where necessary, but also included an option to complete the questionnaire online. A Freephone helpline and email was available for respondents to opt out, ask questions about the survey, enable them to complete their questionnaire over the phone and provide access to a translation and interpreting facility for those whose first language was not English.

### Note on question comparability

The questionnaire was redeveloped for the 2021 National Cancer Patient Experience Survey. Year on year comparisons between 2021, 2022 and 2023 are included in this report for most questions. A review of the questionnaire in 2023 saw four changes being made:

- The question text for Q23 and Q42 were amended. These questions are no longer deemed comparable to 2021 and 2022.
- The long-term condition question (Q67) was amended to include “Autism or autism spectrum condition” as a response option. And the “Neurological condition” answer option was updated to include an example condition changing it to “Neurological condition, such as epilepsy”. These changes see the answer option “Neurological condition, such as epilepsy” as no longer being deemed comparable to 2021 and 2022.
- The ethnic group question (Q71) was amended to include “Roma” as an answer option. The ethnic group question is still deemed comparable to 2021 and 2022.

### How alliance and ICB results are generated

Alliance and ICB results are derived using the post code of each patient, rather than by mapping trust results to ICBs or alliances. This mapping is achieved using lookup files released by the Office for National Statistics.

Alliance and ICB results therefore reflect the experience of people referred from within the geographical footprint.

### Case-mix adjustment

Both unadjusted and adjusted scores are presented in this report. Case-mix adjusted scores allow us to account for the impact that differing patient populations might have on results. By using the case-mix adjusted estimates we can obtain a greater understanding of how an alliance is performing given their

patient population. The factors taken into account in this case-mix adjustment are Male/Female/Non-binary/Other, age, ethnicity, deprivation, and cancer type.

Unadjusted data should be used to see the actual responses from patients relating to the alliance. Case-mix adjusted data, together with expected ranges, should be used to understand whether the results are significantly higher or lower than national results taking account of the patient mix.

### Scoring methodology

Sixty-one questions from the questionnaire are scored as these questions relate directly to patient experience. For all but one question (Q59), the score shows the percentage of respondents who gave the most favourable response to a question. For Q59, respondents rate their overall care on a scale of 0 to 10, of which the average was calculated for this question's score. The percentages in this report have been rounded to the nearest percentage point. Therefore, in some cases the figures do not appear to add up to 100%.

In 2022, following a review of the scoring methodology, a change was made to the scoring of Q12 such that the response option "No, I was told by letter or email" is no longer considered neutral and is now scored as negative.

### Statistical significance

In the reporting of 2023 results, appropriate statistical tests have been undertaken to identify unadjusted scores for which the change over time is 'statistically significant'. A statistically significant difference means that the change in the result is very unlikely to have occurred by chance.

### Suppression

Data is suppressed for two reasons: to ensure unreliable results based on very small numbers of respondents are not released, and to prevent individuals being identifiable in the data.

In cases where a result is based on fewer than 10 responses, the result has been suppressed. For example, where fewer than 10 people answered a question from a particular alliance, the results are not shown for that question for that alliance.

For alliances with an eligible population of 1,000 or fewer, data relating to the respondent and their condition has been suppressed where 5 people or fewer were in a particular category. In instances where only one has been suppressed, the next lowest category has been suppressed to prevent back calculation from the total number of responses.

### Additional suppression

Additional suppression happens if only **one** alliance has a score suppressed. If this happens, we will suppress another alliance's results (both the alliance level and sub-group results for the question) based on the next lowest number of respondents for the score. We do this so that the national score cannot be used to work out the score for the individual alliance.

The same rule applies to groups in each sub-group breakdown. For example, if only one alliance has the 85+ age group suppressed for Q25 we will need to suppress another alliance's results for the 85+ age group on Q25. This suppression is based on the 85+ age group with the next lowest number of respondents for Q25.

## Understanding the results

This report shows how this alliance scored for each question in the survey compared with England results. It is aimed at helping individual alliances to understand their performance and identify areas for local improvement. Below is a description of the type of results presented within this report and how to understand them.

### Expected range charts

The expected range charts in this report show a bar with the lowest and highest score received for each question nationally. Within this bar, an expected range is given (within the grey bar) and a black diamond represents the actual score for this alliance.

Alliances whose score is above the upper limit of the expected range (in the dark blue) are positive outliers, with a score statistically significantly higher than the national mean. This indicates that the alliance performs better than what alliances of the same size and demographics are expected to perform. The opposite is true if the score is below the lower limit of the expected range (in the light

blue); these are negative outliers. For scores within the expected range (in the grey), the score is what we would expect given the alliance's size and demographics.

### **Comparability tables**

The comparability tables show the 2022 and 2023 unadjusted scores for this alliance for each scored question. The Change 2022-2023 and Change overall columns show whether the scores show a statistically significant variation between years. This is shown between 2022-2023 and as an overall between 2021-2023. An upwards arrow indicates a statistically significant increase, a downwards arrow indicates a statistically significant decrease and no arrow indicates no statistically significant change.

The adjusted 2023 score will also be presented for each scored question along with the lower and upper expected range and national score. Scores above the upper limit of the expected range will be highlighted dark blue, scores below the lower limit of the expected range will be highlighted light blue, and scores within the lower and upper limit of the expected ranges will be highlighted grey.

### **Sub-group breakdowns**

Unadjusted scores are shown for tumour group, Male/Female/Non-binary/Other, age, IMD quintile, long-term condition status and ethnicity breakdowns. Unadjusted scores for the same sub-group across different alliances may not be comparable, as they do not account for the impact that differing patient populations might have on results.

### **Tumour group tables**

The tumour group tables show the unadjusted scores for each scored question for each of the 13 tumour groups. Central nervous system is abbreviated as 'CNS' and lower gastrointestinal tract is abbreviated as 'LGT' throughout this report.

### **Age group tables**

The age group tables show the unadjusted scores for each scored question for each of the eight age groups.

### **Male/Female/Non-binary/Other tables**

These tables show the unadjusted scores for the following groups male; female; non-binary; prefer to self-describe; and prefer not to say.

### **Ethnicity tables**

The ethnicity tables show the unadjusted scores for six ethnicity groups.

### **Long-term condition status tables**

The long-term condition status tables show the unadjusted scores for two groups: those who indicate they have one or more long-term conditions and those who indicate that they have no long-term conditions.

### **IMD quintile tables**

The IMD quintile tables show the unadjusted scores for five quintiles based on relative disadvantage, with quintile 1 being the most deprived and quintile 5 being the least deprived.

### **Year on year charts**

The year on year charts show three columns representing the unadjusted scores of the last three years (2021, 2022 and 2023) for each scored question.

### **Trust expected range summary**

The number of scored questions that fell below, within and above the expected range for each trust within the alliance.

### **ICB expected range summary**

The number of scored questions that fell below, within and above the expected range for each ICB within the alliance.

## **National level and England level data**

In some cases (343 respondents in 2023), patients from outside England (from Wales, Scotland, Northern Ireland, the Channel Islands or the Isle of Man) are referred to English NHS trusts for treatment. These patients are described as 'Non-England' in other reports.

Overall response rate at response rate sections shows national level counts and response rate. For cancer alliances and its comparison at comparability tables section, all data is presented at the England level.

## **Further information**

This research was carried out in accordance with the international standard for organisations conducting market and social research (accreditation to ISO20252:2019; certificate number GB08/74322). Our statistical practice is regulated by the Office for Statistics Regulation (OSR). OSR sets the standards of trustworthiness, quality, and value in the Code of Practice for Statistics that all producers of official statistics should adhere to. You are welcome to contact us directly with any comments about how we meet these standards. Alternatively, you can contact OSR by emailing [regulation@statistics.gov.uk](mailto:regulation@statistics.gov.uk) or via the OSR website.

For more information on the methodology, please see the Technical Document. It can be viewed along with the 2023 questionnaire and survey guidance on the website at [www.ncpes.co.uk](http://www.ncpes.co.uk). For all other outputs at alliance level, please see the Excel tables and dashboards at [www.ncpes.co.uk](http://www.ncpes.co.uk).

## Response rate

### Overall response rate

969 patients responded out of a total of 2,557 patients, resulting in a response rate of 38%.

	Sample size	Adjusted sample	Completed	Response rate
Overall response rate	2,708	2,557	969	38%
National	129,231	121,121	63,438	52%

### Respondents by survey type

	Number of respondents
Paper	731
Online	238
Phone	0
Translation service	0
<b>Total</b>	<b>969</b>

### Respondents by tumour group

	Number of respondents
Brain / CNS	3
Breast	225
Colorectal / LGT	119
Gynaecological	59
Haematological	99
Head and neck	30
Lung	63
Prostate	117
Sarcoma	7
Skin	20
Upper gastro	38
Urological	68
Other	121
<b>Total</b>	<b>969</b>

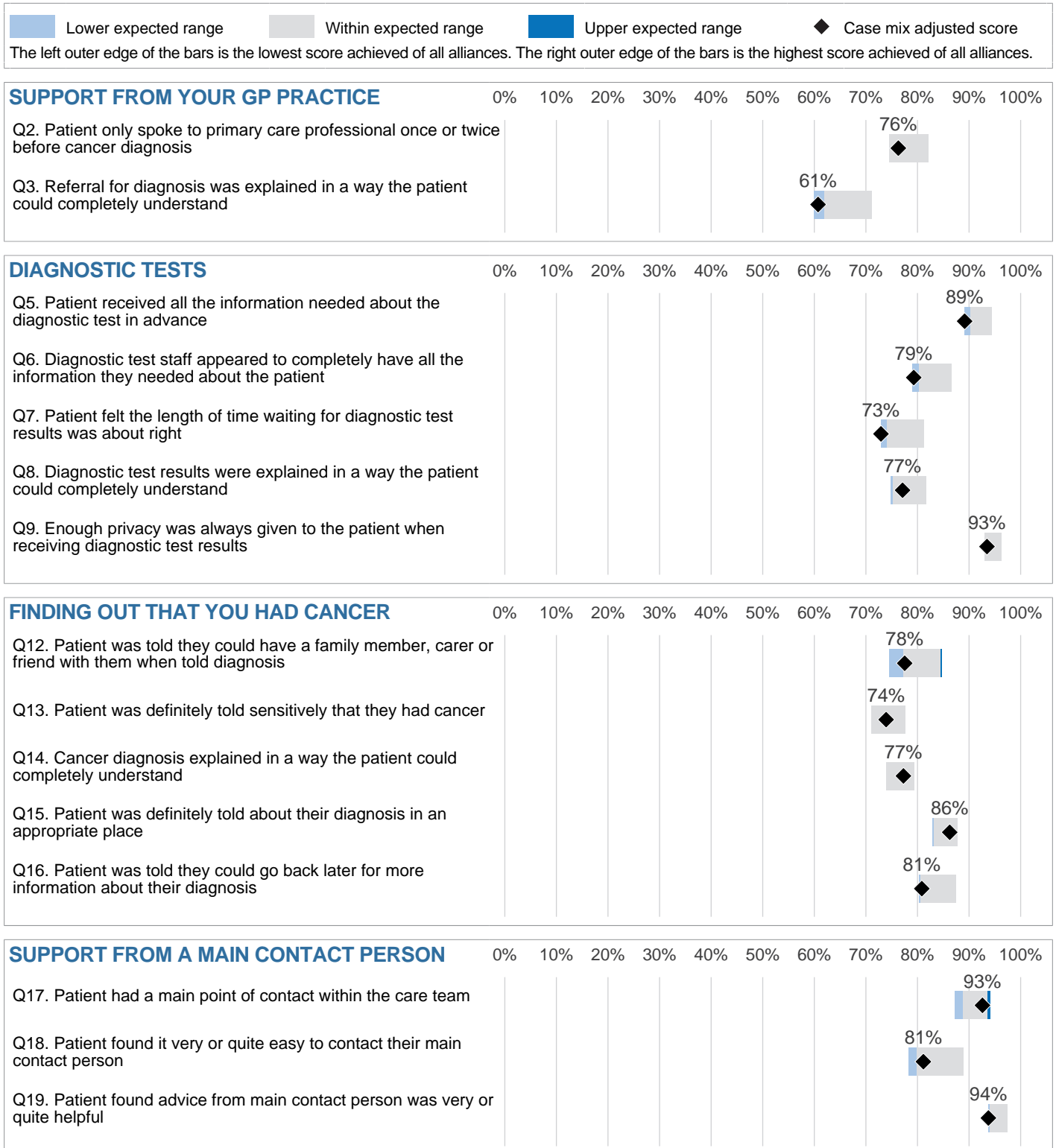


## Respondents by ethnicity

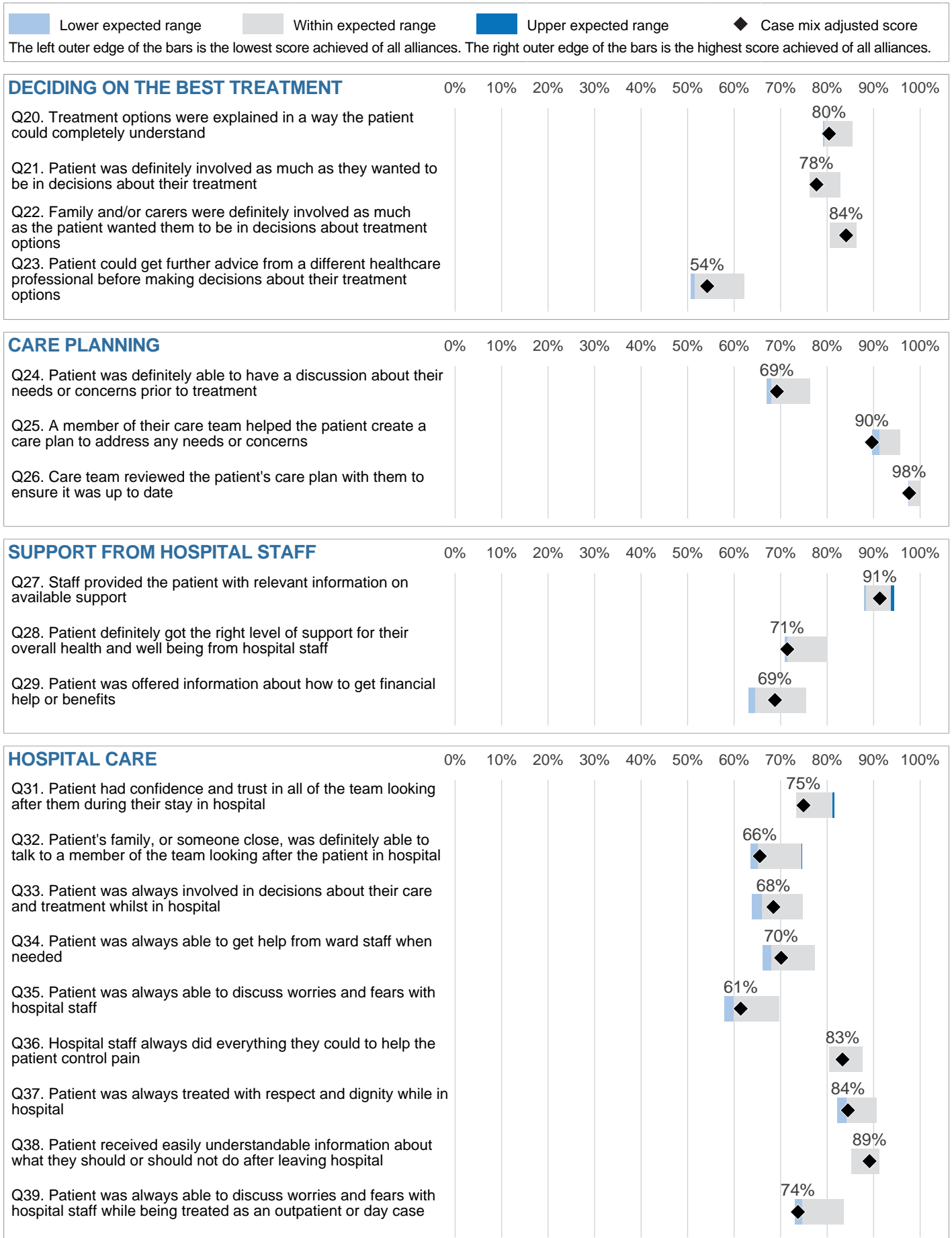
	Number of respondents
<b>White</b>	
English / Welsh / Scottish / Northern Irish / British	572
Irish	17
Gypsy or Irish Traveller	*
Roma	*
Any other White background	65
<b>Mixed / Multiple Ethnic Groups</b>	
White and Black Caribbean	8
White and Black African	8
White and Asian	*
Any other Mixed / multiple ethnic background	6
<b>Asian or Asian British</b>	
Indian	37
Pakistani	24
Bangladeshi	20
Chinese	8
Any other Asian background	15
<b>Black / African / Caribbean / Black British</b>	
African	48
Caribbean	35
Any other Black / African / Caribbean background	*
<b>Other Ethnicity</b>	
Arab	*
Any other ethnic group	10
<b>Not given</b>	
Not given	88
<b>Total</b>	<b>969</b>

\* indicates the count is not shown due to suppression

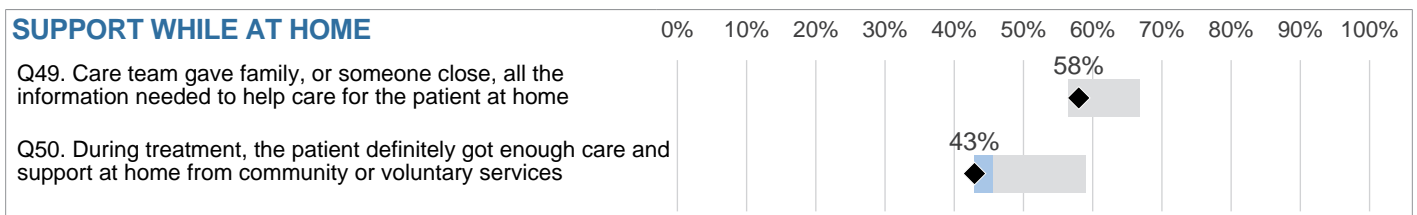
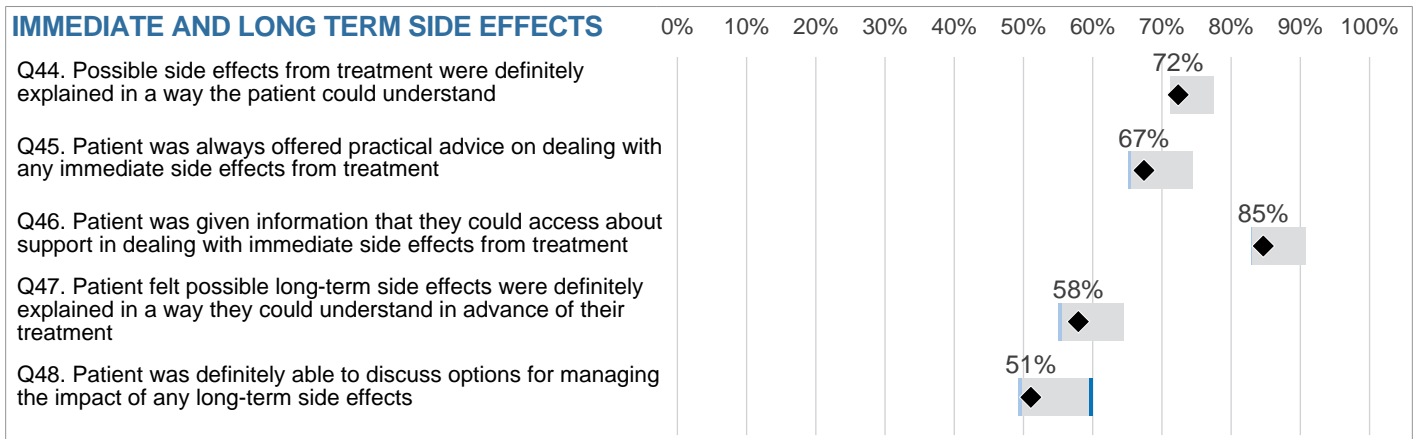
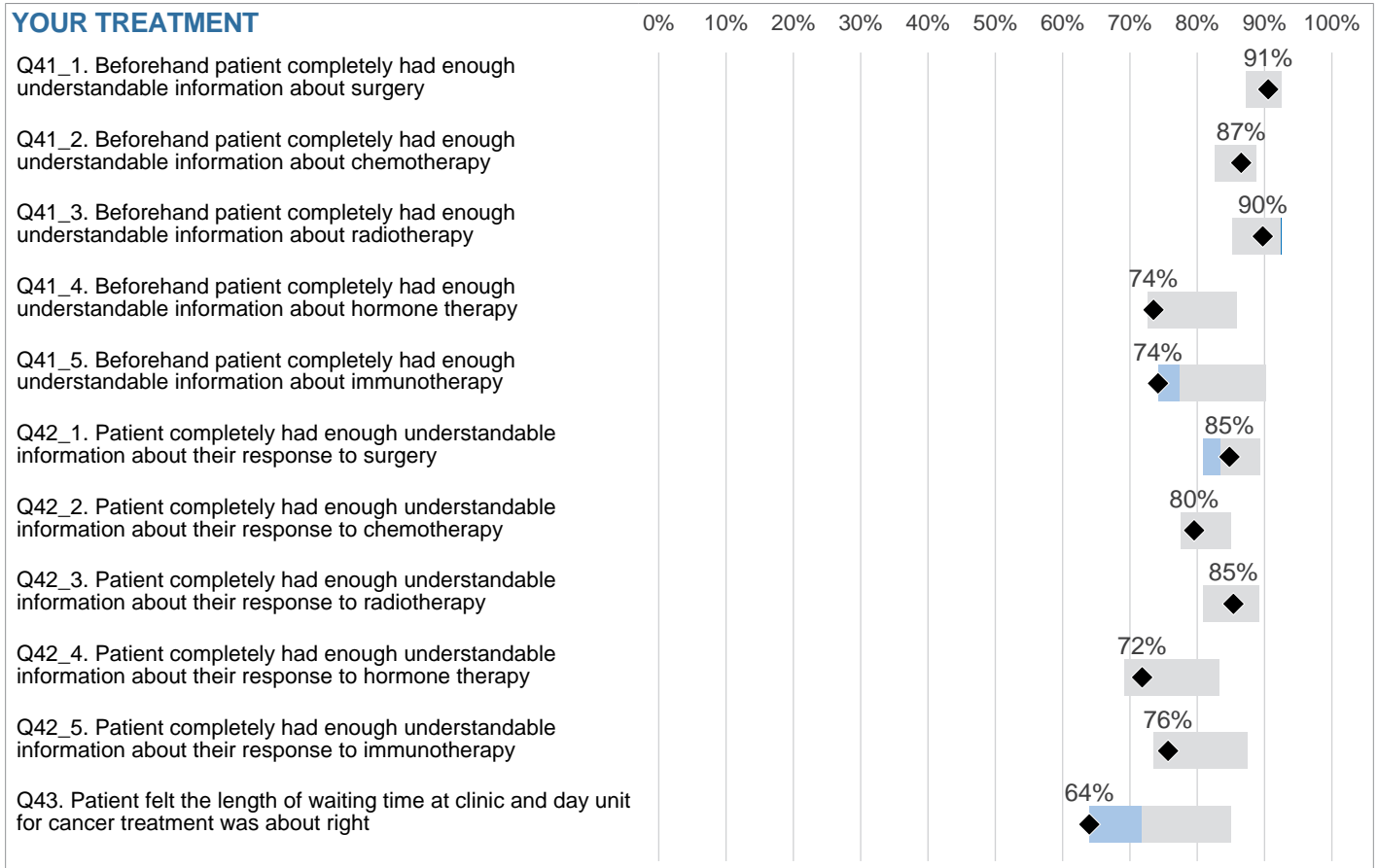
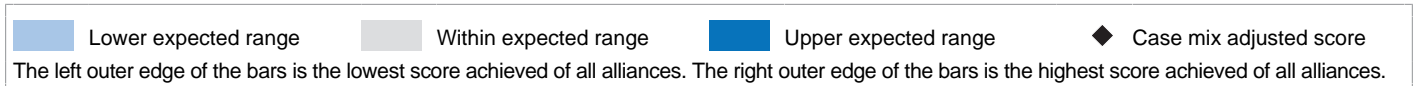
## Expected range charts



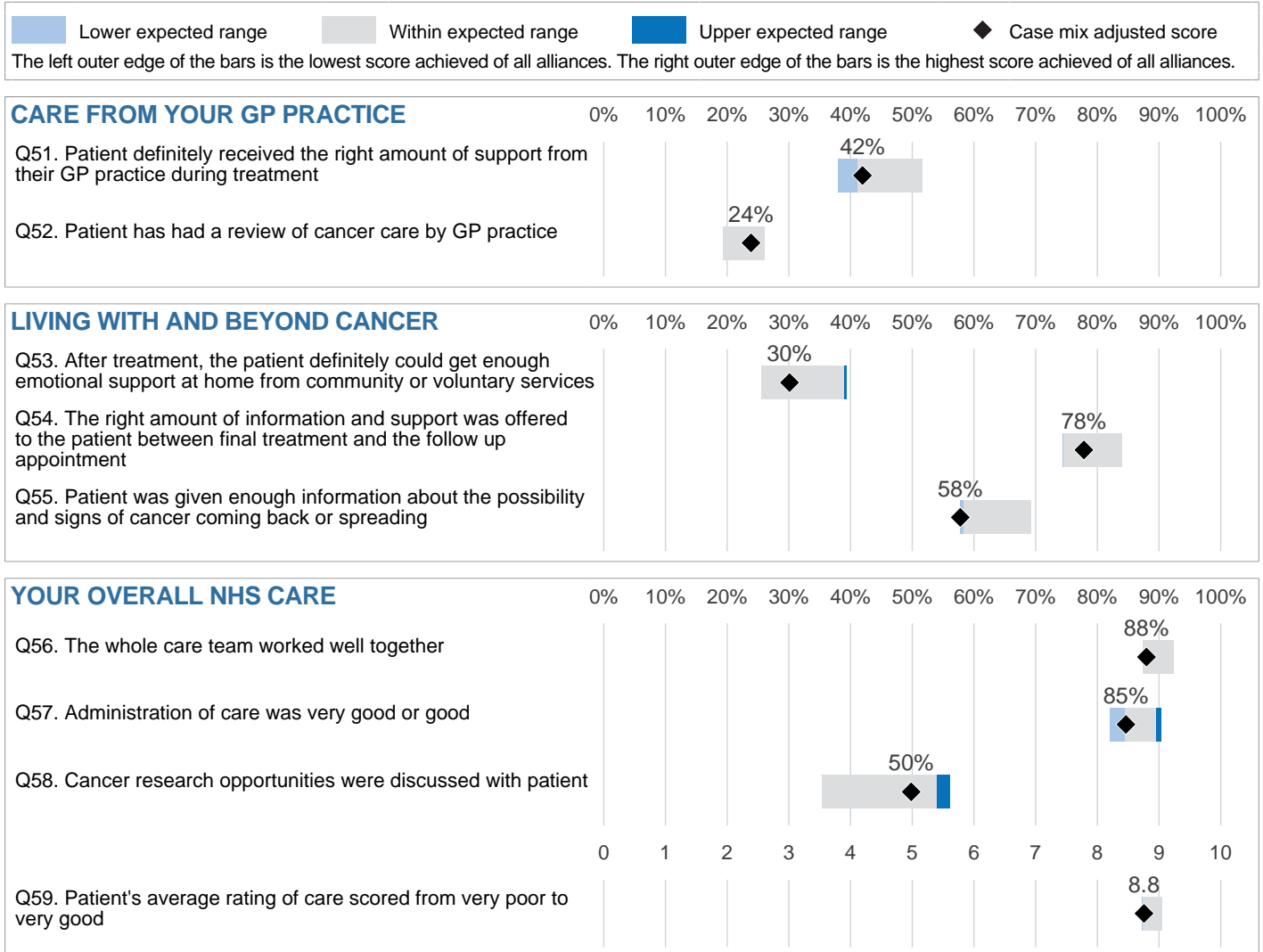
## Expected range charts



## Expected range charts



## Expected range charts



## Comparability tables

\* Indicates where a score is not available due to suppression or a low base size. ▲ or ▼

\*\* No score available for 2022.

Change 2022-2023: Indicates where 2023 score is significantly higher or lower than 2022 score.  
Change overall: Indicates significant change overall (2021, 2022, and 2023).

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<span style="background-color: #e6e6e6; border: 1px solid black; display: inline-block; width: 15px; height: 10px;"></span> Adjusted score between upper and lower expected ranges
<span style="background-color: #1f77b4; border: 1px solid black; display: inline-block; width: 15px; height: 10px;"></span> Adjusted score above upper expected range

SUPPORT FROM YOUR GP PRACTICE	Unadjusted scores						Case mix adjusted scores			England score
	2022 n	2022 score	2023 n	2023 score	Change 2022-2023	Change overall	2023 score	Lower expected range	Upper expected range	
Q2. Patient only spoke to primary care professional once or twice before cancer diagnosis	375	72%	447	72%			76%	74%	82%	78%
Q3. Referral for diagnosis was explained in a way the patient could completely understand	533	59%	600	60%			61%	62%	71%	67%

DIAGNOSTIC TESTS	Unadjusted scores						Case mix adjusted scores			England score
	2022 n	2022 score	2023 n	2023 score	Change 2022-2023	Change overall	2023 score	Lower expected range	Upper expected range	
Q5. Patient received all the information needed about the diagnostic test in advance	700	91%	779	88%			89%	90%	94%	92%
Q6. Diagnostic test staff appeared to completely have all the information they needed about the patient	750	79%	809	77%			79%	80%	87%	83%
Q7. Patient felt the length of time waiting for diagnostic test results was about right	754	75%	819	72%			73%	74%	81%	78%
Q8. Diagnostic test results were explained in a way the patient could completely understand	756	77%	821	75%			77%	75%	82%	78%
Q9. Enough privacy was always given to the patient when receiving diagnostic test results	758	94%	827	93%			93%	93%	96%	95%

FINDING OUT THAT YOU HAD CANCER	Unadjusted scores						Case mix adjusted scores			England score
	2022 n	2022 score	2023 n	2023 score	Change 2022-2023	Change overall	2023 score	Lower expected range	Upper expected range	
Q12. Patient was told they could have a family member, carer or friend with them when told diagnosis	811	75%	881	78%		▲	78%	77%	84%	81%
Q13. Patient was definitely told sensitively that they had cancer	866	71%	941	74%			74%	71%	78%	74%
Q14. Cancer diagnosis explained in a way the patient could completely understand	871	75%	942	76%			77%	74%	79%	77%
Q15. Patient was definitely told about their diagnosis in an appropriate place	868	86%	948	86%			86%	83%	88%	86%
Q16. Patient was told they could go back later for more information about their diagnosis	761	82%	839	81%			81%	81%	87%	84%

SUPPORT FROM A MAIN CONTACT PERSON	Unadjusted scores						Case mix adjusted scores			England score
	2022 n	2022 score	2023 n	2023 score	Change 2022-2023	Change overall	2023 score	Lower expected range	Upper expected range	
Q17. Patient had a main point of contact within the care team	840	92%	916	93%			93%	89%	94%	91%
Q18. Patient found it very or quite easy to contact their main contact person	721	78%	785	80%			81%	80%	89%	84%
Q19. Patient found advice from main contact person was very or quite helpful	745	94%	799	93%			94%	94%	97%	96%




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\*\* No score available for 2022.

▲ or ▼

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Change overall: Indicates significant change overall (2021, 2022, and 2023).

	Adjusted score below lower expected range
	Adjusted score between upper and lower expected ranges
	Adjusted score above upper expected range

DECIDING ON THE BEST TREATMENT	Unadjusted scores						Case mix adjusted scores			England score
	2022 n	2022 score	2023 n	2023 score	Change 2022-2023	Change overall	2023 score	Lower expected range	Upper expected range	
Q20. Treatment options were explained in a way the patient could completely understand	817	<b>79%</b>	900	<b>79%</b>			<b>80%</b>	79%	86%	<b>82%</b>
Q21. Patient was definitely involved as much as they wanted to be in decisions about their treatment	861	<b>76%</b>	930	<b>76%</b>			<b>78%</b>	76%	83%	<b>80%</b>
Q22. Family and/or carers were definitely involved as much as the patient wanted them to be in decisions about treatment options	713	<b>80%</b>	813	<b>83%</b>		▲	<b>84%</b>	81%	86%	<b>83%</b>
Q23. Patient could get further advice from a different healthcare professional before making decisions about their treatment options	**	**	563	<b>55%</b>			<b>54%</b>	52%	62%	<b>57%</b>

CARE PLANNING	Unadjusted scores						Case mix adjusted scores			England score
	2022 n	2022 score	2023 n	2023 score	Change 2022-2023	Change overall	2023 score	Lower expected range	Upper expected range	
Q24. Patient was definitely able to have a discussion about their needs or concerns prior to treatment	776	<b>67%</b>	861	<b>69%</b>			<b>69%</b>	68%	76%	<b>72%</b>
Q25. A member of their care team helped the patient create a care plan to address any needs or concerns	482	<b>90%</b>	548	<b>90%</b>			<b>90%</b>	91%	96%	<b>94%</b>
Q26. Care team reviewed the patient's care plan with them to ensure it was up to date	370	<b>99%</b>	411	<b>98%</b>			<b>98%</b>	98%	100%	<b>99%</b>

SUPPORT FROM HOSPITAL STAFF	Unadjusted scores						Case mix adjusted scores			England score
	2022 n	2022 score	2023 n	2023 score	Change 2022-2023	Change overall	2023 score	Lower expected range	Upper expected range	
Q27. Staff provided the patient with relevant information on available support	744	<b>89%</b>	829	<b>91%</b>		▲	<b>91%</b>	88%	94%	<b>91%</b>
Q28. Patient definitely got the right level of support for their overall health and well being from hospital staff	862	<b>74%</b>	942	<b>71%</b>			<b>71%</b>	71%	80%	<b>76%</b>
Q29. Patient was offered information about how to get financial help or benefits	522	<b>63%</b>	608	<b>68%</b>			<b>69%</b>	65%	76%	<b>70%</b>

## Comparability tables

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▲ or ▼

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HOSPITAL CARE	Unadjusted scores						Case mix adjusted scores			England score
	2022 n	2022 score	2023 n	2023 score	Change 2022-2023	Change overall	2023 score	Lower expected range	Upper expected range	
Q31. Patient had confidence and trust in all of the team looking after them during their stay in hospital	420	81%	457	75%			75%	73%	81%	77%
Q32. Patient's family, or someone close, was definitely able to talk to a member of the team looking after the patient in hospital	337	64%	363	66%		▲	66%	65%	75%	70%
Q33. Patient was always involved in decisions about their care and treatment whilst in hospital	403	68%	441	68%			68%	66%	75%	70%
Q34. Patient was always able to get help from ward staff when needed	406	73%	443	70%			70%	68%	77%	73%
Q35. Patient was always able to discuss worries and fears with hospital staff	393	64%	424	61%			61%	60%	70%	65%
Q36. Hospital staff always did everything they could to help the patient control pain	355	84%	408	83%			83%	80%	88%	84%
Q37. Patient was always treated with respect and dignity while in hospital	419	86%	449	84%			84%	84%	91%	87%
Q38. Patient received easily understandable information about what they should or should not do after leaving hospital	407	88%	435	89%			89%	85%	91%	88%
Q39. Patient was always able to discuss worries and fears with hospital staff while being treated as an outpatient or day case	744	72%	819	72%			74%	75%	84%	79%

YOUR TREATMENT	Unadjusted scores						Case mix adjusted scores			England score
	2022 n	2022 score	2023 n	2023 score	Change 2022-2023	Change overall	2023 score	Lower expected range	Upper expected range	
Q41_1. Beforehand patient completely had enough understandable information about surgery	504	85%	514	90%	▲		91%	87%	92%	90%
Q41_2. Beforehand patient completely had enough understandable information about chemotherapy	449	84%	495	87%			87%	83%	89%	86%
Q41_3. Beforehand patient completely had enough understandable information about radiotherapy	259	87%	293	89%			90%	85%	92%	89%
Q41_4. Beforehand patient completely had enough understandable information about hormone therapy	122	75%	145	73%			74%	73%	86%	79%
Q41_5. Beforehand patient completely had enough understandable information about immunotherapy	105	84%	127	75%			74%	77%	90%	84%
Q42_1. Patient completely had enough understandable information about their response to surgery	**	**	516	84%			85%	84%	89%	86%
Q42_2. Patient completely had enough understandable information about their response to chemotherapy	**	**	499	80%			80%	78%	85%	81%
Q42_3. Patient completely had enough understandable information about their response to radiotherapy	**	**	287	85%			85%	81%	89%	85%
Q42_4. Patient completely had enough understandable information about their response to hormone therapy	**	**	141	72%			72%	69%	83%	76%
Q42_5. Patient completely had enough understandable information about their response to immunotherapy	**	**	125	75%			76%	74%	87%	81%
Q43. Patient felt the length of waiting time at clinic and day unit for cancer treatment was about right	845	67%	901	62%		▼	64%	72%	85%	78%



## Comparability tables

\* Indicates where a score is not available due to suppression or a low base size. ▲ or ▼

\*\* No score available for 2022.

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<span style="background-color: #d9e1f2; border: 1px solid #000; display: inline-block; width: 15px; height: 10px;"></span> Adjusted score below lower expected range
<span style="background-color: #e6e6e6; border: 1px solid #000; display: inline-block; width: 15px; height: 10px;"></span> Adjusted score between upper and lower expected ranges
<span style="background-color: #0070c0; border: 1px solid #000; display: inline-block; width: 15px; height: 10px;"></span> Adjusted score above upper expected range

IMMEDIATE AND LONG TERM SIDE EFFECTS	Unadjusted scores						Case mix adjusted scores			England score
	2022 n	2022 score	2023 n	2023 score	Change 2022-2023	Change overall	2023 score	Lower expected range	Upper expected range	
Q44. Possible side effects from treatment were definitely explained in a way the patient could understand	816	<b>72%</b>	886	<b>73%</b>			<b>72%</b>	71%	78%	<b>74%</b>
Q45. Patient was always offered practical advice on dealing with any immediate side effects from treatment	776	<b>66%</b>	839	<b>67%</b>			<b>67%</b>	66%	74%	<b>70%</b>
Q46. Patient was given information that they could access about support in dealing with immediate side effects from treatment	656	<b>84%</b>	697	<b>84%</b>			<b>85%</b>	83%	91%	<b>87%</b>
Q47. Patient felt possible long-term side effects were definitely explained in a way they could understand in advance of their treatment	769	<b>57%</b>	850	<b>59%</b>			<b>58%</b>	56%	65%	<b>60%</b>
Q48. Patient was definitely able to discuss options for managing the impact of any long-term side effects	679	<b>51%</b>	741	<b>51%</b>			<b>51%</b>	50%	59%	<b>55%</b>

SUPPORT WHILE AT HOME	Unadjusted scores						Case mix adjusted scores			England score
	2022 n	2022 score	2023 n	2023 score	Change 2022-2023	Change overall	2023 score	Lower expected range	Upper expected range	
Q49. Care team gave family, or someone close, all the information needed to help care for the patient at home	599	<b>58%</b>	640	<b>58%</b>			<b>58%</b>	56%	67%	<b>62%</b>
Q50. During treatment, the patient definitely got enough care and support at home from community or voluntary services	369	<b>47%</b>	437	<b>41%</b>			<b>43%</b>	46%	59%	<b>52%</b>

CARE FROM YOUR GP PRACTICE	Unadjusted scores						Case mix adjusted scores			England score
	2022 n	2022 score	2023 n	2023 score	Change 2022-2023	Change overall	2023 score	Lower expected range	Upper expected range	
Q51. Patient definitely received the right amount of support from their GP practice during treatment	482	<b>39%</b>	562	<b>41%</b>			<b>42%</b>	41%	52%	<b>46%</b>
Q52. Patient has had a review of cancer care by GP practice	808	<b>24%</b>	873	<b>26%</b>			<b>24%</b>	19%	26%	<b>23%</b>

LIVING WITH AND BEYOND CANCER	Unadjusted scores						Case mix adjusted scores			England score
	2022 n	2022 score	2023 n	2023 score	Change 2022-2023	Change overall	2023 score	Lower expected range	Upper expected range	
Q53. After treatment, the patient definitely could get enough emotional support at home from community or voluntary services	211	<b>25%</b>	247	<b>30%</b>			<b>30%</b>	26%	39%	<b>32%</b>
Q54. The right amount of information and support was offered to the patient between final treatment and the follow up appointment	382	<b>76%</b>	443	<b>76%</b>			<b>78%</b>	75%	84%	<b>79%</b>
Q55. Patient was given enough information about the possibility and signs of cancer coming back or spreading	694	<b>55%</b>	739	<b>56%</b>			<b>58%</b>	58%	69%	<b>64%</b>

YOUR OVERALL NHS CARE	Unadjusted scores						Case mix adjusted scores			England score
	2022 n	2022 score	2023 n	2023 score	Change 2022-2023	Change overall	2023 score	Lower expected range	Upper expected range	
Q56. The whole care team worked well together	823	<b>88%</b>	874	<b>88%</b>			<b>88%</b>	87%	92%	<b>90%</b>
Q57. Administration of care was very good or good	850	<b>85%</b>	930	<b>85%</b>			<b>85%</b>	84%	90%	<b>87%</b>
Q58. Cancer research opportunities were discussed with patient	534	<b>49%</b>	604	<b>51%</b>			<b>50%</b>	35%	54%	<b>45%</b>
Q59. Patient's average rating of care scored from very poor to very good	826	<b>8.7</b>	917	<b>8.7</b>			<b>8.8</b>	8.7	9.0	<b>8.9</b>

## Tumour group tables

\* Indicates where a score is not available due to suppression or a low base size.

	Tumour group													
	Brain / CNS	Breast	Colorectal / LGT	Gynaecological	Haematological	Head and neck	Lung	Prostate	Sarcoma	Skin	Upper gastro	Urological	Other	All cancers
Q2. Patient only spoke to primary care professional once or twice before cancer diagnosis	*	90%	57%	72%	65%	78%	65%	78%	*	67%	58%	65%	68%	<b>72%</b>
Q3. Referral for diagnosis was explained in a way the patient could completely understand	*	75%	55%	54%	48%	54%	38%	64%	*	71%	65%	63%	49%	<b>60%</b>

	Tumour group													
	Brain / CNS	Breast	Colorectal / LGT	Gynaecological	Haematological	Head and neck	Lung	Prostate	Sarcoma	Skin	Upper gastro	Urological	Other	All cancers
Q5. Patient received all the information needed about the diagnostic test in advance	*	85%	82%	89%	90%	83%	94%	89%	*	87%	97%	96%	93%	<b>88%</b>
Q6. Diagnostic test staff appeared to completely have all the information they needed about the patient	*	74%	79%	74%	83%	58%	77%	86%	*	82%	84%	76%	75%	<b>77%</b>
Q7. Patient felt the length of time waiting for diagnostic test results was about right	*	73%	70%	60%	82%	63%	65%	75%	*	76%	63%	75%	71%	<b>72%</b>
Q8. Diagnostic test results were explained in a way the patient could completely understand	*	75%	81%	72%	78%	60%	81%	77%	*	94%	69%	77%	71%	<b>75%</b>
Q9. Enough privacy was always given to the patient when receiving diagnostic test results	*	94%	94%	96%	95%	80%	91%	94%	*	88%	97%	90%	91%	<b>93%</b>

	Tumour group													
	Brain / CNS	Breast	Colorectal / LGT	Gynaecological	Haematological	Head and neck	Lung	Prostate	Sarcoma	Skin	Upper gastro	Urological	Other	All cancers
Q12. Patient was told they could have a family member, carer or friend with them when told diagnosis	*	82%	84%	86%	82%	67%	84%	68%	*	67%	77%	70%	77%	<b>78%</b>
Q13. Patient was definitely told sensitively that they had cancer	*	78%	80%	78%	77%	70%	71%	68%	*	80%	63%	64%	72%	<b>74%</b>
Q14. Cancer diagnosis explained in a way the patient could completely understand	*	80%	81%	69%	82%	69%	77%	77%	*	80%	69%	71%	71%	<b>76%</b>
Q15. Patient was definitely told about their diagnosis in an appropriate place	*	92%	85%	90%	90%	79%	84%	85%	*	85%	81%	82%	83%	<b>86%</b>
Q16. Patient was told they could go back later for more information about their diagnosis	*	84%	88%	78%	83%	91%	81%	77%	*	84%	71%	76%	78%	<b>81%</b>

## Tumour group tables

\* Indicates where a score is not available due to suppression or a low base size.

	Tumour group													
	Brain / CNS	Breast	Colorectal / LGT	Gynaecological	Haematological	Head and neck	Lung	Prostate	Sarcoma	Skin	Upper gastro	Urological	Other	All cancers
Q17. Patient had a main point of contact within the care team	*	95%	92%	96%	98%	100%	92%	83%	*	84%	94%	88%	95%	<b>93%</b>
Q18. Patient found it very or quite easy to contact their main contact person	*	76%	77%	82%	88%	84%	86%	72%	*	83%	77%	85%	83%	<b>80%</b>
Q19. Patient found advice from main contact person was very or quite helpful	*	93%	90%	94%	97%	88%	100%	96%	*	100%	87%	92%	92%	<b>93%</b>

	Tumour group													
	Brain / CNS	Breast	Colorectal / LGT	Gynaecological	Haematological	Head and neck	Lung	Prostate	Sarcoma	Skin	Upper gastro	Urological	Other	All cancers
Q20. Treatment options were explained in a way the patient could completely understand	*	82%	82%	80%	84%	62%	88%	73%	*	84%	79%	78%	75%	<b>79%</b>
Q21. Patient was definitely involved as much as they wanted to be in decisions about their treatment	*	73%	75%	80%	82%	67%	82%	77%	*	95%	74%	71%	76%	<b>76%</b>
Q22. Family and/or carers were definitely involved as much as the patient wanted them to be in decisions about treatment options	*	81%	86%	88%	84%	76%	87%	77%	*	78%	86%	78%	84%	<b>83%</b>
Q23. Patient could get further advice from a different healthcare professional before making decisions about their treatment options	*	55%	53%	43%	61%	63%	62%	48%	*	*	61%	49%	54%	<b>55%</b>

	Tumour group													
	Brain / CNS	Breast	Colorectal / LGT	Gynaecological	Haematological	Head and neck	Lung	Prostate	Sarcoma	Skin	Upper gastro	Urological	Other	All cancers
Q24. Patient was definitely able to have a discussion about their needs or concerns prior to treatment	*	65%	70%	81%	71%	69%	70%	67%	*	57%	57%	67%	72%	<b>69%</b>
Q25. A member of their care team helped the patient create a care plan to address any needs or concerns	*	87%	92%	87%	92%	89%	86%	87%	*	*	91%	91%	93%	<b>90%</b>
Q26. Care team reviewed the patient's care plan with them to ensure it was up to date	*	95%	96%	100%	98%	100%	100%	97%	*	*	100%	97%	98%	<b>98%</b>

	Tumour group													
	Brain / CNS	Breast	Colorectal / LGT	Gynaecological	Haematological	Head and neck	Lung	Prostate	Sarcoma	Skin	Upper gastro	Urological	Other	All cancers
Q27. Staff provided the patient with relevant information on available support	*	93%	86%	96%	97%	77%	93%	89%	*	79%	97%	91%	90%	<b>91%</b>
Q28. Patient definitely got the right level of support for their overall health and well being from hospital staff	*	67%	66%	71%	79%	74%	77%	70%	*	65%	73%	69%	70%	<b>71%</b>
Q29. Patient was offered information about how to get financial help or benefits	*	69%	61%	88%	73%	77%	74%	49%	*	*	52%	64%	75%	<b>68%</b>

## Tumour group tables

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	Tumour group													
	Brain/ CNS	Breast	Colorectal/ LGT	Gynaecological	Haematological	Head and neck	Lung	Prostate	Sarcoma	Skin	Upper gastro	Urological	Other	All cancers
Q31. Patient had confidence and trust in all of the team looking after them during their stay in hospital	*	76%	77%	62%	65%	67%	90%	87%	*	*	72%	73%	73%	<b>75%</b>
Q32. Patient's family, or someone close, was definitely able to talk to a member of the team looking after the patient in hospital	*	65%	58%	50%	78%	69%	67%	94%	*	*	71%	55%	68%	<b>66%</b>
Q33. Patient was always involved in decisions about their care and treatment whilst in hospital	*	69%	60%	67%	67%	50%	85%	86%	*	*	61%	67%	69%	<b>68%</b>
Q34. Patient was always able to get help from ward staff when needed	*	81%	69%	52%	68%	67%	80%	86%	*	*	63%	62%	67%	<b>70%</b>
Q35. Patient was always able to discuss worries and fears with hospital staff	*	66%	61%	43%	75%	57%	68%	74%	*	*	57%	46%	64%	<b>61%</b>
Q36. Hospital staff always did everything they could to help the patient control pain	*	84%	78%	71%	83%	88%	94%	93%	*	*	90%	75%	81%	<b>83%</b>
Q37. Patient was always treated with respect and dignity while in hospital	*	89%	76%	69%	85%	78%	85%	95%	*	*	74%	87%	81%	<b>84%</b>
Q38. Patient received easily understandable information about what they should or should not do after leaving hospital	*	83%	93%	92%	91%	71%	94%	95%	*	*	83%	93%	93%	<b>89%</b>
Q39. Patient was always able to discuss worries and fears with hospital staff while being treated as an outpatient or day case	*	67%	72%	82%	83%	77%	75%	76%	*	53%	68%	60%	73%	<b>72%</b>

	Tumour group													
	Brain/ CNS	Breast	Colorectal/ LGT	Gynaecological	Haematological	Head and neck	Lung	Prostate	Sarcoma	Skin	Upper gastro	Urological	Other	All cancers
Q41_1. Beforehand patient completely had enough understandable information about surgery	*	90%	90%	93%	*	76%	93%	98%	*	83%	94%	88%	93%	<b>90%</b>
Q41_2. Beforehand patient completely had enough understandable information about chemotherapy	*	84%	88%	89%	88%	*	88%	82%	*	*	95%	75%	89%	<b>87%</b>
Q41_3. Beforehand patient completely had enough understandable information about radiotherapy	*	89%	97%	90%	86%	77%	93%	88%	*	*	*	*	90%	<b>89%</b>
Q41_4. Beforehand patient completely had enough understandable information about hormone therapy	*	66%	*	*	*	*	*	79%	*	*	*	*	86%	<b>73%</b>
Q41_5. Beforehand patient completely had enough understandable information about immunotherapy	*	56%	*	*	90%	*	86%	*	*	*	*	59%	83%	<b>75%</b>
Q42_1. Patient completely had enough understandable information about their response to surgery	*	84%	85%	91%	*	59%	80%	90%	*	92%	95%	83%	86%	<b>84%</b>
Q42_2. Patient completely had enough understandable information about their response to chemotherapy	*	72%	81%	89%	85%	*	84%	68%	*	*	78%	69%	87%	<b>80%</b>
Q42_3. Patient completely had enough understandable information about their response to radiotherapy	*	84%	86%	89%	86%	75%	93%	88%	*	*	*	*	88%	<b>85%</b>
Q42_4. Patient completely had enough understandable information about their response to hormone therapy	*	66%	*	*	*	*	*	72%	*	*	*	*	85%	<b>72%</b>
Q42_5. Patient completely had enough understandable information about their response to immunotherapy	*	64%	*	*	100%	*	82%	*	*	*	*	52%	81%	<b>75%</b>
Q43. Patient felt the length of waiting time at clinic and day unit for cancer treatment was about right	*	47%	77%	70%	62%	85%	59%	80%	*	63%	46%	62%	60%	<b>62%</b>

## Tumour group tables

\* Indicates where a score is not available due to suppression or a low base size.

	Tumour group													
	Brain / CNS	Breast	Colorectal / LGT	Gynaecological	Haematological	Head and neck	Lung	Prostate	Sarcoma	Skin	Upper gastro	Urological	Other	All cancers
Q44. Possible side effects from treatment were definitely explained in a way the patient could understand	*	71%	77%	80%	81%	69%	68%	71%	*	100%	66%	68%	68%	<b>73%</b>
Q45. Patient was always offered practical advice on dealing with any immediate side effects from treatment	*	64%	70%	68%	74%	60%	70%	66%	*	88%	63%	65%	67%	<b>67%</b>
Q46. Patient was given information that they could access about support in dealing with immediate side effects from treatment	*	83%	80%	89%	90%	86%	86%	75%	*	83%	96%	78%	87%	<b>84%</b>
Q47. Patient felt possible long-term side effects were definitely explained in a way they could understand in advance of their treatment	*	61%	61%	62%	66%	72%	59%	63%	*	71%	41%	52%	50%	<b>59%</b>
Q48. Patient was definitely able to discuss options for managing the impact of any long-term side effects	*	49%	58%	50%	54%	52%	55%	51%	*	54%	52%	53%	46%	<b>51%</b>

	Tumour group													
	Brain / CNS	Breast	Colorectal / LGT	Gynaecological	Haematological	Head and neck	Lung	Prostate	Sarcoma	Skin	Upper gastro	Urological	Other	All cancers
Q49. Care team gave family, or someone close, all the information needed to help care for the patient at home	*	51%	59%	63%	64%	43%	59%	67%	*	*	63%	40%	61%	<b>58%</b>
Q50. During treatment, the patient definitely got enough care and support at home from community or voluntary services	*	35%	36%	59%	51%	50%	59%	40%	*	*	57%	26%	34%	<b>41%</b>

	Tumour group													
	Brain / CNS	Breast	Colorectal / LGT	Gynaecological	Haematological	Head and neck	Lung	Prostate	Sarcoma	Skin	Upper gastro	Urological	Other	All cancers
Q51. Patient definitely received the right amount of support from their GP practice during treatment	*	37%	45%	60%	41%	29%	36%	47%	*	18%	38%	53%	40%	<b>41%</b>
Q52. Patient has had a review of cancer care by GP practice	*	23%	32%	21%	22%	32%	28%	30%	*	21%	43%	20%	27%	<b>26%</b>

## Tumour group tables

\* Indicates where a score is not available due to suppression or a low base size.

	Tumour group													
	Brain / CNS	Breast	Colorectal / LGT	Gynaecological	Haematological	Head and neck	Lung	Prostate	Sarcoma	Skin	Upper gastro	Urological	Other	All cancers
Q53. After treatment, the patient definitely could get enough emotional support at home from community or voluntary services	*	23%	28%	39%	35%	45%	39%	22%	*	*	50%	33%	29%	<b>30%</b>
Q54. The right amount of information and support was offered to the patient between final treatment and the follow up appointment	*	70%	82%	78%	76%	79%	65%	80%	*	80%	67%	76%	83%	<b>76%</b>
Q55. Patient was given enough information about the possibility and signs of cancer coming back or spreading	*	49%	54%	61%	75%	41%	59%	53%	*	81%	59%	53%	61%	<b>56%</b>

	Tumour group													
	Brain / CNS	Breast	Colorectal / LGT	Gynaecological	Haematological	Head and neck	Lung	Prostate	Sarcoma	Skin	Upper gastro	Urological	Other	All cancers
Q56. The whole care team worked well together	*	88%	91%	91%	90%	84%	90%	85%	*	89%	85%	85%	90%	<b>88%</b>
Q57. Administration of care was very good or good	*	84%	81%	82%	92%	74%	85%	84%	*	89%	89%	82%	88%	<b>85%</b>
Q58. Cancer research opportunities were discussed with patient	*	48%	54%	63%	57%	43%	36%	44%	*	*	68%	68%	42%	<b>51%</b>
Q59. Patient's average rating of care scored from very poor to very good	*	8.6	8.5	8.7	9.0	8.5	8.8	8.9	*	8.5	8.5	8.5	8.7	<b>8.7</b>

## Age group tables

\* Indicates where a score is not available due to suppression or a low base size.

	Age								
	16 - 24	25 - 34	35 - 44	45 - 54	55 - 64	65 - 74	75 - 84	85+	All
<b>SUPPORT FROM YOUR GP PRACTICE</b>									
Q2. Patient only spoke to primary care professional once or twice before cancer diagnosis	*	*	59%	77%	68%	71%	79%	100%	<b>72%</b>
Q3. Referral for diagnosis was explained in a way the patient could completely understand	*	73%	52%	64%	55%	66%	55%	58%	<b>60%</b>

	Age								
	16 - 24	25 - 34	35 - 44	45 - 54	55 - 64	65 - 74	75 - 84	85+	All
<b>DIAGNOSTIC TESTS</b>									
Q5. Patient received all the information needed about the diagnostic test in advance	*	70%	85%	88%	87%	89%	90%	91%	<b>88%</b>
Q6. Diagnostic test staff appeared to completely have all the information they needed about the patient	*	80%	71%	69%	77%	81%	79%	75%	<b>77%</b>
Q7. Patient felt the length of time waiting for diagnostic test results was about right	*	50%	50%	61%	70%	77%	79%	69%	<b>72%</b>
Q8. Diagnostic test results were explained in a way the patient could completely understand	*	64%	64%	72%	73%	77%	78%	86%	<b>75%</b>
Q9. Enough privacy was always given to the patient when receiving diagnostic test results	*	92%	100%	84%	94%	91%	96%	100%	<b>93%</b>

	Age								
	16 - 24	25 - 34	35 - 44	45 - 54	55 - 64	65 - 74	75 - 84	85+	All
<b>FINDING OUT THAT YOU HAD CANCER</b>									
Q12. Patient was told they could have a family member, carer or friend with them when told diagnosis	*	69%	69%	69%	79%	79%	83%	79%	<b>78%</b>
Q13. Patient was definitely told sensitively that they had cancer	*	69%	72%	63%	76%	76%	74%	75%	<b>74%</b>
Q14. Cancer diagnosis explained in a way the patient could completely understand	*	77%	70%	73%	77%	78%	75%	86%	<b>76%</b>
Q15. Patient was definitely told about their diagnosis in an appropriate place	*	77%	87%	84%	86%	87%	85%	93%	<b>86%</b>
Q16. Patient was told they could go back later for more information about their diagnosis	*	91%	83%	82%	81%	84%	77%	74%	<b>81%</b>

	Age								
	16 - 24	25 - 34	35 - 44	45 - 54	55 - 64	65 - 74	75 - 84	85+	All
<b>SUPPORT FROM A MAIN CONTACT PERSON</b>									
Q17. Patient had a main point of contact within the care team	*	100%	94%	96%	91%	93%	92%	90%	<b>93%</b>
Q18. Patient found it very or quite easy to contact their main contact person	*	62%	76%	76%	78%	81%	82%	91%	<b>80%</b>
Q19. Patient found advice from main contact person was very or quite helpful	*	92%	86%	91%	93%	94%	95%	100%	<b>93%</b>

	Age								
	16 - 24	25 - 34	35 - 44	45 - 54	55 - 64	65 - 74	75 - 84	85+	All
<b>DECIDING ON THE BEST TREATMENT</b>									
Q20. Treatment options were explained in a way the patient could completely understand	*	62%	72%	77%	79%	82%	80%	80%	<b>79%</b>
Q21. Patient was definitely involved as much as they wanted to be in decisions about their treatment	*	62%	67%	73%	75%	79%	80%	74%	<b>76%</b>
Q22. Family and/or carers were definitely involved as much as the patient wanted them to be in decisions about treatment options	*	75%	67%	83%	81%	82%	87%	88%	<b>83%</b>
Q23. Patient could get further advice from a different healthcare professional before making decisions about their treatment options	*	*	49%	51%	54%	58%	55%	43%	<b>55%</b>

## Age group tables

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CARE PLANNING	Age								
	16 - 24	25 - 34	35 - 44	45 - 54	55 - 64	65 - 74	75 - 84	85+	All
Q24. Patient was definitely able to have a discussion about their needs or concerns prior to treatment	*	75%	58%	73%	67%	72%	68%	59%	<b>69%</b>
Q25. A member of their care team helped the patient create a care plan to address any needs or concerns	*	*	91%	91%	90%	90%	88%	89%	<b>90%</b>
Q26. Care team reviewed the patient's care plan with them to ensure it was up to date	*	*	100%	94%	97%	98%	98%	100%	<b>98%</b>

SUPPORT FROM HOSPITAL STAFF	Age								
	16 - 24	25 - 34	35 - 44	45 - 54	55 - 64	65 - 74	75 - 84	85+	All
Q27. Staff provided the patient with relevant information on available support	*	92%	81%	92%	92%	93%	92%	73%	<b>91%</b>
Q28. Patient definitely got the right level of support for their overall health and well being from hospital staff	*	62%	56%	68%	68%	72%	76%	69%	<b>71%</b>
Q29. Patient was offered information about how to get financial help or benefits	*	67%	71%	79%	67%	66%	66%	41%	<b>68%</b>

HOSPITAL CARE	Age								
	16 - 24	25 - 34	35 - 44	45 - 54	55 - 64	65 - 74	75 - 84	85+	All
Q31. Patient had confidence and trust in all of the team looking after them during their stay in hospital	*	*	58%	61%	73%	79%	83%	84%	<b>75%</b>
Q32. Patient's family, or someone close, was definitely able to talk to a member of the team looking after the patient in hospital	*	*	61%	51%	66%	73%	63%	69%	<b>66%</b>
Q33. Patient was always involved in decisions about their care and treatment whilst in hospital	*	*	65%	64%	68%	71%	72%	72%	<b>68%</b>
Q34. Patient was always able to get help from ward staff when needed	*	*	63%	71%	71%	70%	72%	78%	<b>70%</b>
Q35. Patient was always able to discuss worries and fears with hospital staff	*	*	57%	59%	61%	63%	58%	63%	<b>61%</b>
Q36. Hospital staff always did everything they could to help the patient control pain	*	*	79%	80%	78%	88%	86%	75%	<b>83%</b>
Q37. Patient was always treated with respect and dignity while in hospital	*	*	84%	74%	83%	83%	90%	83%	<b>84%</b>
Q38. Patient received easily understandable information about what they should or should not do after leaving hospital	*	*	88%	82%	91%	91%	90%	82%	<b>89%</b>
Q39. Patient was always able to discuss worries and fears with hospital staff while being treated as an outpatient or day case	*	58%	54%	65%	73%	72%	78%	79%	<b>72%</b>



## Age group tables

\* Indicates where a score is not available due to suppression or a low base size.

YOUR TREATMENT	Age								
	16 - 24	25 - 34	35 - 44	45 - 54	55 - 64	65 - 74	75 - 84	85+	All
Q41_1. Beforehand patient completely had enough understandable information about surgery	*	80%	75%	89%	88%	93%	96%	95%	<b>90%</b>
Q41_2. Beforehand patient completely had enough understandable information about chemotherapy	*	80%	69%	86%	91%	88%	89%	90%	<b>87%</b>
Q41_3. Beforehand patient completely had enough understandable information about radiotherapy	*	*	87%	84%	88%	94%	93%	*	<b>89%</b>
Q41_4. Beforehand patient completely had enough understandable information about hormone therapy	*	*	*	57%	78%	74%	82%	*	<b>73%</b>
Q41_5. Beforehand patient completely had enough understandable information about immunotherapy	*	*	*	65%	81%	72%	81%	*	<b>75%</b>
Q42_1. Patient completely had enough understandable information about their response to surgery	*	80%	76%	79%	84%	86%	92%	81%	<b>84%</b>
Q42_2. Patient completely had enough understandable information about their response to chemotherapy	*	70%	61%	77%	81%	80%	86%	90%	<b>80%</b>
Q42_3. Patient completely had enough understandable information about their response to radiotherapy	*	*	86%	82%	89%	87%	82%	*	<b>85%</b>
Q42_4. Patient completely had enough understandable information about their response to hormone therapy	*	*	*	56%	85%	70%	82%	*	<b>72%</b>
Q42_5. Patient completely had enough understandable information about their response to immunotherapy	*	*	*	78%	80%	68%	76%	*	<b>75%</b>
Q43. Patient felt the length of waiting time at clinic and day unit for cancer treatment was about right	*	15%	40%	56%	64%	64%	68%	78%	<b>62%</b>

IMMEDIATE AND LONG TERM SIDE EFFECTS	Age								
	16 - 24	25 - 34	35 - 44	45 - 54	55 - 64	65 - 74	75 - 84	85+	All
Q44. Possible side effects from treatment were definitely explained in a way the patient could understand	*	85%	69%	75%	73%	73%	71%	68%	<b>73%</b>
Q45. Patient was always offered practical advice on dealing with any immediate side effects from treatment	*	69%	58%	63%	69%	68%	67%	72%	<b>67%</b>
Q46. Patient was given information that they could access about support in dealing with immediate side effects from treatment	*	91%	80%	88%	83%	86%	80%	88%	<b>84%</b>
Q47. Patient felt possible long-term side effects were definitely explained in a way they could understand in advance of their treatment	*	77%	57%	58%	62%	58%	58%	50%	<b>59%</b>
Q48. Patient was definitely able to discuss options for managing the impact of any long-term side effects	*	58%	43%	43%	50%	58%	50%	43%	<b>51%</b>

SUPPORT WHILE AT HOME	Age								
	16 - 24	25 - 34	35 - 44	45 - 54	55 - 64	65 - 74	75 - 84	85+	All
Q49. Care team gave family, or someone close, all the information needed to help care for the patient at home	*	45%	42%	50%	56%	63%	59%	69%	<b>58%</b>
Q50. During treatment, the patient definitely got enough care and support at home from community or voluntary services	*	*	17%	33%	43%	45%	48%	48%	<b>41%</b>

CARE FROM YOUR GP PRACTICE	Age								
	16 - 24	25 - 34	35 - 44	45 - 54	55 - 64	65 - 74	75 - 84	85+	All
Q51. Patient definitely received the right amount of support from their GP practice during treatment	*	*	25%	37%	50%	44%	41%	24%	<b>41%</b>
Q52. Patient has had a review of cancer care by GP practice	*	38%	39%	23%	30%	26%	23%	13%	<b>26%</b>

## Age group tables

\* Indicates where a score is not available due to suppression or a low base size.

	Age								
	16 - 24	25 - 34	35 - 44	45 - 54	55 - 64	65 - 74	75 - 84	85+	All
Q53. After treatment, the patient definitely could get enough emotional support at home from community or voluntary services	*	*	22%	23%	27%	31%	40%	*	<b>30%</b>
Q54. The right amount of information and support was offered to the patient between final treatment and the follow up appointment	*	*	62%	73%	74%	82%	78%	67%	<b>76%</b>
Q55. Patient was given enough information about the possibility and signs of cancer coming back or spreading	*	30%	49%	57%	51%	56%	61%	68%	<b>56%</b>

	Age								
	16 - 24	25 - 34	35 - 44	45 - 54	55 - 64	65 - 74	75 - 84	85+	All
Q56. The whole care team worked well together	*	92%	70%	87%	87%	89%	91%	94%	<b>88%</b>
Q57. Administration of care was very good or good	*	62%	73%	85%	82%	89%	85%	88%	<b>85%</b>
Q58. Cancer research opportunities were discussed with patient	*	27%	67%	51%	58%	47%	48%	40%	<b>51%</b>
Q59. Patient's average rating of care scored from very poor to very good	*	7.6	7.6	8.6	8.6	8.8	8.9	8.9	<b>8.7</b>

## Male/Female/Non-binary/Other tables

\* Indicates where a score is not available due to suppression or a low base size.

<b>SUPPORT FROM YOUR GP PRACTICE</b>							
	Male/Female/Non-binary/Other						
	Female	Male	Non-binary	Prefer to self-describe	Prefer not to say	Not given	All
Q2. Patient only spoke to primary care professional once or twice before cancer diagnosis	73%	72%	*	*	*	70%	<b>72%</b>
Q3. Referral for diagnosis was explained in a way the patient could completely understand	62%	58%	*	*	*	48%	<b>60%</b>

<b>DIAGNOSTIC TESTS</b>							
	Male/Female/Non-binary/Other						
	Female	Male	Non-binary	Prefer to self-describe	Prefer not to say	Not given	All
Q5. Patient received all the information needed about the diagnostic test in advance	86%	93%	*	*	*	86%	<b>88%</b>
Q6. Diagnostic test staff appeared to completely have all the information they needed about the patient	74%	82%	*	*	*	75%	<b>77%</b>
Q7. Patient felt the length of time waiting for diagnostic test results was about right	71%	73%	*	*	*	68%	<b>72%</b>
Q8. Diagnostic test results were explained in a way the patient could completely understand	72%	79%	*	*	*	79%	<b>75%</b>
Q9. Enough privacy was always given to the patient when receiving diagnostic test results	92%	94%	*	*	*	96%	<b>93%</b>

<b>FINDING OUT THAT YOU HAD CANCER</b>							
	Male/Female/Non-binary/Other						
	Female	Male	Non-binary	Prefer to self-describe	Prefer not to say	Not given	All
Q12. Patient was told they could have a family member, carer or friend with them when told diagnosis	78%	77%	*	*	*	84%	<b>78%</b>
Q13. Patient was definitely told sensitively that they had cancer	73%	75%	*	*	*	77%	<b>74%</b>
Q14. Cancer diagnosis explained in a way the patient could completely understand	75%	77%	*	*	*	79%	<b>76%</b>
Q15. Patient was definitely told about their diagnosis in an appropriate place	87%	86%	*	*	*	81%	<b>86%</b>
Q16. Patient was told they could go back later for more information about their diagnosis	80%	84%	*	*	*	70%	<b>81%</b>

<b>SUPPORT FROM A MAIN CONTACT PERSON</b>							
	Male/Female/Non-binary/Other						
	Female	Male	Non-binary	Prefer to self-describe	Prefer not to say	Not given	All
Q17. Patient had a main point of contact within the care team	94%	90%	*	*	*	98%	<b>93%</b>
Q18. Patient found it very or quite easy to contact their main contact person	79%	80%	*	*	*	82%	<b>80%</b>
Q19. Patient found advice from main contact person was very or quite helpful	93%	94%	*	*	*	93%	<b>93%</b>

## Male/Female/Non-binary/Other tables

\* Indicates where a score is not available due to suppression or a low base size.

<b>DECIDING ON THE BEST TREATMENT</b>							
	Male/Female/Non-binary/Other						
	Female	Male	Non-binary	Prefer to self-describe	Prefer not to say	Not given	All
Q20. Treatment options were explained in a way the patient could completely understand	79%	80%	*	*	*	79%	<b>79%</b>
Q21. Patient was definitely involved as much as they wanted to be in decisions about their treatment	74%	78%	*	*	*	85%	<b>76%</b>
Q22. Family and/or carers were definitely involved as much as the patient wanted them to be in decisions about treatment options	83%	81%	*	*	*	89%	<b>83%</b>
Q23. Patient could get further advice from a different healthcare professional before making decisions about their treatment options	56%	54%	*	*	*	52%	<b>55%</b>

<b>CARE PLANNING</b>							
	Male/Female/Non-binary/Other						
	Female	Male	Non-binary	Prefer to self-describe	Prefer not to say	Not given	All
Q24. Patient was definitely able to have a discussion about their needs or concerns prior to treatment	68%	69%	*	*	*	68%	<b>69%</b>
Q25. A member of their care team helped the patient create a care plan to address any needs or concerns	91%	89%	*	*	*	83%	<b>90%</b>
Q26. Care team reviewed the patient's care plan with them to ensure it was up to date	97%	98%	*	*	*	100%	<b>98%</b>

<b>SUPPORT FROM HOSPITAL STAFF</b>							
	Male/Female/Non-binary/Other						
	Female	Male	Non-binary	Prefer to self-describe	Prefer not to say	Not given	All
Q27. Staff provided the patient with relevant information on available support	91%	92%	*	*	*	91%	<b>91%</b>
Q28. Patient definitely got the right level of support for their overall health and well being from hospital staff	69%	72%	*	*	*	83%	<b>71%</b>
Q29. Patient was offered information about how to get financial help or benefits	73%	61%	*	*	*	71%	<b>68%</b>

## Male/Female/Non-binary/Other tables

\* Indicates where a score is not available due to suppression or a low base size.

	Male/Female/Non-binary/Other						All
	Female	Male	Non-binary	Prefer to self-describe	Prefer not to say	Not given	
Q31. Patient had confidence and trust in all of the team looking after them during their stay in hospital	69%	80%	*	*	*	78%	<b>75%</b>
Q32. Patient's family, or someone close, was definitely able to talk to a member of the team looking after the patient in hospital	61%	68%	*	*	*	85%	<b>66%</b>
Q33. Patient was always involved in decisions about their care and treatment whilst in hospital	66%	71%	*	*	*	76%	<b>68%</b>
Q34. Patient was always able to get help from ward staff when needed	69%	72%	*	*	*	70%	<b>70%</b>
Q35. Patient was always able to discuss worries and fears with hospital staff	57%	65%	*	*	*	57%	<b>61%</b>
Q36. Hospital staff always did everything they could to help the patient control pain	81%	83%	*	*	*	95%	<b>83%</b>
Q37. Patient was always treated with respect and dignity while in hospital	81%	85%	*	*	*	91%	<b>84%</b>
Q38. Patient received easily understandable information about what they should or should not do after leaving hospital	86%	92%	*	*	*	87%	<b>89%</b>
Q39. Patient was always able to discuss worries and fears with hospital staff while being treated as an outpatient or day case	69%	74%	*	*	*	81%	<b>72%</b>

	Male/Female/Non-binary/Other						All
	Female	Male	Non-binary	Prefer to self-describe	Prefer not to say	Not given	
Q41_1. Beforehand patient completely had enough understandable information about surgery	90%	91%	*	*	*	88%	<b>90%</b>
Q41_2. Beforehand patient completely had enough understandable information about chemotherapy	86%	88%	*	*	*	90%	<b>87%</b>
Q41_3. Beforehand patient completely had enough understandable information about radiotherapy	91%	88%	*	*	*	80%	<b>89%</b>
Q41_4. Beforehand patient completely had enough understandable information about hormone therapy	72%	76%	*	*	*	*	<b>73%</b>
Q41_5. Beforehand patient completely had enough understandable information about immunotherapy	72%	79%	*	*	*	*	<b>75%</b>
Q42_1. Patient completely had enough understandable information about their response to surgery	86%	83%	*	*	*	77%	<b>84%</b>
Q42_2. Patient completely had enough understandable information about their response to chemotherapy	77%	83%	*	*	*	85%	<b>80%</b>
Q42_3. Patient completely had enough understandable information about their response to radiotherapy	85%	87%	*	*	*	70%	<b>85%</b>
Q42_4. Patient completely had enough understandable information about their response to hormone therapy	73%	69%	*	*	*	*	<b>72%</b>
Q42_5. Patient completely had enough understandable information about their response to immunotherapy	72%	78%	*	*	*	*	<b>75%</b>
Q43. Patient felt the length of waiting time at clinic and day unit for cancer treatment was about right	54%	72%	*	*	*	67%	<b>62%</b>

## Male/Female/Non-binary/Other tables

\* Indicates where a score is not available due to suppression or a low base size.

<b>IMMEDIATE AND LONG TERM SIDE EFFECTS</b>							
	Male/Female/Non-binary/Other						
	Female	Male	Non-binary	Prefer to self-describe	Prefer not to say	Not given	All
Q44. Possible side effects from treatment were definitely explained in a way the patient could understand	72%	74%	*	*	*	65%	<b>73%</b>
Q45. Patient was always offered practical advice on dealing with any immediate side effects from treatment	66%	69%	*	*	*	66%	<b>67%</b>
Q46. Patient was given information that they could access about support in dealing with immediate side effects from treatment	85%	84%	*	*	*	80%	<b>84%</b>
Q47. Patient felt possible long-term side effects were definitely explained in a way they could understand in advance of their treatment	58%	61%	*	*	*	56%	<b>59%</b>
Q48. Patient was definitely able to discuss options for managing the impact of any long-term side effects	50%	53%	*	*	*	56%	<b>51%</b>

<b>SUPPORT WHILE AT HOME</b>							
	Male/Female/Non-binary/Other						
	Female	Male	Non-binary	Prefer to self-describe	Prefer not to say	Not given	All
Q49. Care team gave family, or someone close, all the information needed to help care for the patient at home	54%	60%	*	*	*	74%	<b>58%</b>
Q50. During treatment, the patient definitely got enough care and support at home from community or voluntary services	40%	41%	*	*	*	50%	<b>41%</b>

<b>CARE FROM YOUR GP PRACTICE</b>							
	Male/Female/Non-binary/Other						
	Female	Male	Non-binary	Prefer to self-describe	Prefer not to say	Not given	All
Q51. Patient definitely received the right amount of support from their GP practice during treatment	40%	45%	*	*	*	34%	<b>41%</b>
Q52. Patient has had a review of cancer care by GP practice	26%	28%	*	*	*	19%	<b>26%</b>

<b>LIVING WITH AND BEYOND CANCER</b>							
	Male/Female/Non-binary/Other						
	Female	Male	Non-binary	Prefer to self-describe	Prefer not to say	Not given	All
Q53. After treatment, the patient definitely could get enough emotional support at home from community or voluntary services	31%	30%	*	*	*	25%	<b>30%</b>
Q54. The right amount of information and support was offered to the patient between final treatment and the follow up appointment	73%	79%	*	*	*	77%	<b>76%</b>
Q55. Patient was given enough information about the possibility and signs of cancer coming back or spreading	54%	59%	*	*	*	53%	<b>56%</b>

## Male/Female/Non-binary/Other tables

\* Indicates where a score is not available due to suppression or a low base size.

YOUR OVERALL NHS CARE	Male/Female/Non-binary/Other						All
	Female	Male	Non-binary	Prefer to self-describe	Prefer not to say	Not given	
Q56. The whole care team worked well together	89%	88%	*	*	*	87%	<b>88%</b>
Q57. Administration of care was very good or good	84%	85%	*	*	*	88%	<b>85%</b>
Q58. Cancer research opportunities were discussed with patient	51%	51%	*	*	*	52%	<b>51%</b>
Q59. Patient's average rating of care scored from very poor to very good	8.6	8.8	*	*	*	8.7	<b>8.7</b>

## Ethnicity tables

\* Indicates where a score is not available due to suppression or a low base size.

	Ethnicity						
	White	Mixed	Asian	Black	Other	Not given	All
<b>SUPPORT FROM YOUR GP PRACTICE</b>							
Q2. Patient only spoke to primary care professional once or twice before cancer diagnosis	77%	*	65%	53%	*	70%	<b>72%</b>
Q3. Referral for diagnosis was explained in a way the patient could completely understand	64%	46%	45%	54%	*	51%	<b>60%</b>

	Ethnicity						
	White	Mixed	Asian	Black	Other	Not given	All
<b>DIAGNOSTIC TESTS</b>							
Q5. Patient received all the information needed about the diagnostic test in advance	89%	87%	84%	90%	*	88%	<b>88%</b>
Q6. Diagnostic test staff appeared to completely have all the information they needed about the patient	79%	69%	74%	74%	*	72%	<b>77%</b>
Q7. Patient felt the length of time waiting for diagnostic test results was about right	73%	63%	58%	80%	*	73%	<b>72%</b>
Q8. Diagnostic test results were explained in a way the patient could completely understand	77%	71%	69%	71%	*	80%	<b>75%</b>
Q9. Enough privacy was always given to the patient when receiving diagnostic test results	93%	81%	92%	91%	*	94%	<b>93%</b>

	Ethnicity						
	White	Mixed	Asian	Black	Other	Not given	All
<b>FINDING OUT THAT YOU HAD CANCER</b>							
Q12. Patient was told they could have a family member, carer or friend with them when told diagnosis	77%	82%	82%	75%	90%	81%	<b>78%</b>
Q13. Patient was definitely told sensitively that they had cancer	73%	73%	75%	77%	92%	74%	<b>74%</b>
Q14. Cancer diagnosis explained in a way the patient could completely understand	76%	83%	79%	77%	75%	76%	<b>76%</b>
Q15. Patient was definitely told about their diagnosis in an appropriate place	86%	81%	89%	90%	100%	81%	<b>86%</b>
Q16. Patient was told they could go back later for more information about their diagnosis	82%	90%	84%	79%	64%	74%	<b>81%</b>

	Ethnicity						
	White	Mixed	Asian	Black	Other	Not given	All
<b>SUPPORT FROM A MAIN CONTACT PERSON</b>							
Q17. Patient had a main point of contact within the care team	93%	89%	95%	90%	92%	95%	<b>93%</b>
Q18. Patient found it very or quite easy to contact their main contact person	80%	73%	76%	85%	60%	80%	<b>80%</b>
Q19. Patient found advice from main contact person was very or quite helpful	94%	71%	97%	91%	91%	91%	<b>93%</b>

	Ethnicity						
	White	Mixed	Asian	Black	Other	Not given	All
<b>DECIDING ON THE BEST TREATMENT</b>							
Q20. Treatment options were explained in a way the patient could completely understand	80%	77%	80%	74%	100%	80%	<b>79%</b>
Q21. Patient was definitely involved as much as they wanted to be in decisions about their treatment	78%	90%	73%	71%	50%	77%	<b>76%</b>
Q22. Family and/or carers were definitely involved as much as the patient wanted them to be in decisions about treatment options	84%	89%	75%	76%	*	85%	<b>83%</b>
Q23. Patient could get further advice from a different healthcare professional before making decisions about their treatment options	59%	40%	46%	33%	*	60%	<b>55%</b>



## Ethnicity tables

\* Indicates where a score is not available due to suppression or a low base size.

CARE PLANNING	Ethnicity						
	White	Mixed	Asian	Black	Other	Not given	All
Q24. Patient was definitely able to have a discussion about their needs or concerns prior to treatment	69%	55%	70%	64%	67%	69%	<b>69%</b>
Q25. A member of their care team helped the patient create a care plan to address any needs or concerns	90%	75%	89%	92%	*	87%	<b>90%</b>
Q26. Care team reviewed the patient's care plan with them to ensure it was up to date	98%	*	100%	95%	*	97%	<b>98%</b>

SUPPORT FROM HOSPITAL STAFF	Ethnicity						
	White	Mixed	Asian	Black	Other	Not given	All
Q27. Staff provided the patient with relevant information on available support	92%	86%	92%	84%	75%	94%	<b>91%</b>
Q28. Patient definitely got the right level of support for their overall health and well being from hospital staff	69%	68%	75%	66%	58%	82%	<b>71%</b>
Q29. Patient was offered information about how to get financial help or benefits	69%	69%	71%	65%	40%	63%	<b>68%</b>

HOSPITAL CARE	Ethnicity						
	White	Mixed	Asian	Black	Other	Not given	All
Q31. Patient had confidence and trust in all of the team looking after them during their stay in hospital	75%	64%	71%	73%	*	80%	<b>75%</b>
Q32. Patient's family, or someone close, was definitely able to talk to a member of the team looking after the patient in hospital	60%	85%	65%	79%	*	80%	<b>66%</b>
Q33. Patient was always involved in decisions about their care and treatment whilst in hospital	71%	58%	62%	69%	*	66%	<b>68%</b>
Q34. Patient was always able to get help from ward staff when needed	70%	77%	75%	65%	*	69%	<b>70%</b>
Q35. Patient was always able to discuss worries and fears with hospital staff	61%	62%	62%	67%	*	58%	<b>61%</b>
Q36. Hospital staff always did everything they could to help the patient control pain	84%	82%	76%	80%	*	91%	<b>83%</b>
Q37. Patient was always treated with respect and dignity while in hospital	84%	85%	79%	81%	*	94%	<b>84%</b>
Q38. Patient received easily understandable information about what they should or should not do after leaving hospital	90%	83%	89%	91%	*	85%	<b>89%</b>
Q39. Patient was always able to discuss worries and fears with hospital staff while being treated as an outpatient or day case	74%	70%	65%	65%	55%	77%	<b>72%</b>

## Ethnicity tables

\* Indicates where a score is not available due to suppression or a low base size.

YOUR TREATMENT	Ethnicity						
	White	Mixed	Asian	Black	Other	Not given	All
Q41_1. Beforehand patient completely had enough understandable information about surgery	90%	*	92%	91%	*	90%	<b>90%</b>
Q41_2. Beforehand patient completely had enough understandable information about chemotherapy	87%	*	82%	88%	*	94%	<b>87%</b>
Q41_3. Beforehand patient completely had enough understandable information about radiotherapy	89%	*	86%	89%	*	92%	<b>89%</b>
Q41_4. Beforehand patient completely had enough understandable information about hormone therapy	72%	*	87%	*	*	73%	<b>73%</b>
Q41_5. Beforehand patient completely had enough understandable information about immunotherapy	74%	*	71%	*	*	82%	<b>75%</b>
Q42_1. Patient completely had enough understandable information about their response to surgery	84%	*	87%	87%	*	85%	<b>84%</b>
Q42_2. Patient completely had enough understandable information about their response to chemotherapy	79%	*	77%	78%	*	85%	<b>80%</b>
Q42_3. Patient completely had enough understandable information about their response to radiotherapy	86%	*	83%	81%	*	84%	<b>85%</b>
Q42_4. Patient completely had enough understandable information about their response to hormone therapy	70%	*	73%	*	*	91%	<b>72%</b>
Q42_5. Patient completely had enough understandable information about their response to immunotherapy	74%	*	62%	*	*	100%	<b>75%</b>
Q43. Patient felt the length of waiting time at clinic and day unit for cancer treatment was about right	64%	65%	56%	56%	36%	64%	<b>62%</b>

IMMEDIATE AND LONG TERM SIDE EFFECTS	Ethnicity						
	White	Mixed	Asian	Black	Other	Not given	All
Q44. Possible side effects from treatment were definitely explained in a way the patient could understand	74%	70%	73%	68%	73%	69%	<b>73%</b>
Q45. Patient was always offered practical advice on dealing with any immediate side effects from treatment	71%	55%	59%	57%	50%	66%	<b>67%</b>
Q46. Patient was given information that they could access about support in dealing with immediate side effects from treatment	85%	75%	86%	79%	*	83%	<b>84%</b>
Q47. Patient felt possible long-term side effects were definitely explained in a way they could understand in advance of their treatment	58%	48%	65%	62%	58%	61%	<b>59%</b>
Q48. Patient was definitely able to discuss options for managing the impact of any long-term side effects	51%	48%	50%	46%	40%	59%	<b>51%</b>

SUPPORT WHILE AT HOME	Ethnicity						
	White	Mixed	Asian	Black	Other	Not given	All
Q49. Care team gave family, or someone close, all the information needed to help care for the patient at home	57%	37%	56%	64%	*	65%	<b>58%</b>
Q50. During treatment, the patient definitely got enough care and support at home from community or voluntary services	43%	17%	36%	41%	*	44%	<b>41%</b>

CARE FROM YOUR GP PRACTICE	Ethnicity						
	White	Mixed	Asian	Black	Other	Not given	All
Q51. Patient definitely received the right amount of support from their GP practice during treatment	40%	43%	47%	46%	*	43%	<b>41%</b>
Q52. Patient has had a review of cancer care by GP practice	24%	33%	33%	32%	36%	25%	<b>26%</b>

## Ethnicity tables

\* Indicates where a score is not available due to suppression or a low base size.

<b>LIVING WITH AND BEYOND CANCER</b>	Ethnicity						
	White	Mixed	Asian	Black	Other	Not given	All
Q53. After treatment, the patient definitely could get enough emotional support at home from community or voluntary services	35%	*	26%	24%	*	24%	<b>30%</b>
Q54. The right amount of information and support was offered to the patient between final treatment and the follow up appointment	78%	64%	71%	70%	*	79%	<b>76%</b>
Q55. Patient was given enough information about the possibility and signs of cancer coming back or spreading	57%	57%	51%	58%	*	58%	<b>56%</b>

<b>YOUR OVERALL NHS CARE</b>	Ethnicity						
	White	Mixed	Asian	Black	Other	Not given	All
Q56. The whole care team worked well together	89%	76%	88%	85%	91%	91%	<b>88%</b>
Q57. Administration of care was very good or good	84%	87%	85%	86%	92%	88%	<b>85%</b>
Q58. Cancer research opportunities were discussed with patient	51%	44%	56%	50%	*	47%	<b>51%</b>
Q59. Patient's average rating of care scored from very poor to very good	8.8	8.4	8.1	8.4	8.5	8.7	<b>8.7</b>

## IMD quintile tables

\* Indicates where a score is not available due to suppression or a low base size.

SUPPORT FROM YOUR GP PRACTICE	IMD quintile					
	1 (most deprived)	2	3	4	5 (least deprived)	All
Q2. Patient only spoke to primary care professional once or twice before cancer diagnosis	68%	74%	69%	77%	76%	72%
Q3. Referral for diagnosis was explained in a way the patient could completely understand	55%	61%	59%	66%	57%	60%

DIAGNOSTIC TESTS	IMD quintile					
	1 (most deprived)	2	3	4	5 (least deprived)	All
Q5. Patient received all the information needed about the diagnostic test in advance	88%	89%	89%	84%	94%	88%
Q6. Diagnostic test staff appeared to completely have all the information they needed about the patient	73%	79%	74%	81%	81%	77%
Q7. Patient felt the length of time waiting for diagnostic test results was about right	73%	71%	66%	76%	74%	72%
Q8. Diagnostic test results were explained in a way the patient could completely understand	74%	73%	74%	83%	74%	75%
Q9. Enough privacy was always given to the patient when receiving diagnostic test results	91%	95%	90%	96%	91%	93%

FINDING OUT THAT YOU HAD CANCER	IMD quintile					
	1 (most deprived)	2	3	4	5 (least deprived)	All
Q12. Patient was told they could have a family member, carer or friend with them when told diagnosis	78%	78%	79%	76%	79%	78%
Q13. Patient was definitely told sensitively that they had cancer	71%	75%	76%	72%	73%	74%
Q14. Cancer diagnosis explained in a way the patient could completely understand	78%	77%	76%	76%	75%	76%
Q15. Patient was definitely told about their diagnosis in an appropriate place	84%	88%	87%	90%	79%	86%
Q16. Patient was told they could go back later for more information about their diagnosis	79%	84%	81%	76%	85%	81%

SUPPORT FROM A MAIN CONTACT PERSON	IMD quintile					
	1 (most deprived)	2	3	4	5 (least deprived)	All
Q17. Patient had a main point of contact within the care team	91%	93%	93%	94%	93%	93%
Q18. Patient found it very or quite easy to contact their main contact person	83%	82%	77%	75%	81%	80%
Q19. Patient found advice from main contact person was very or quite helpful	91%	94%	94%	93%	95%	93%

## IMD quintile tables

\* Indicates where a score is not available due to suppression or a low base size.

DECIDING ON THE BEST TREATMENT	IMD quintile					
	1 (most deprived)	2	3	4	5 (least deprived)	All
Q20. Treatment options were explained in a way the patient could completely understand	78%	80%	80%	80%	79%	79%
Q21. Patient was definitely involved as much as they wanted to be in decisions about their treatment	72%	77%	74%	83%	78%	76%
Q22. Family and/or carers were definitely involved as much as the patient wanted them to be in decisions about treatment options	81%	85%	79%	84%	83%	83%
Q23. Patient could get further advice from a different healthcare professional before making decisions about their treatment options	53%	53%	59%	48%	61%	55%

CARE PLANNING	IMD quintile					
	1 (most deprived)	2	3	4	5 (least deprived)	All
Q24. Patient was definitely able to have a discussion about their needs or concerns prior to treatment	69%	65%	70%	68%	76%	69%
Q25. A member of their care team helped the patient create a care plan to address any needs or concerns	92%	88%	89%	89%	90%	90%
Q26. Care team reviewed the patient's care plan with them to ensure it was up to date	99%	96%	97%	100%	96%	98%

SUPPORT FROM HOSPITAL STAFF	IMD quintile					
	1 (most deprived)	2	3	4	5 (least deprived)	All
Q27. Staff provided the patient with relevant information on available support	87%	92%	93%	91%	91%	91%
Q28. Patient definitely got the right level of support for their overall health and well being from hospital staff	68%	72%	70%	71%	73%	71%
Q29. Patient was offered information about how to get financial help or benefits	70%	68%	75%	62%	58%	68%

HOSPITAL CARE	IMD quintile					
	1 (most deprived)	2	3	4	5 (least deprived)	All
Q31. Patient had confidence and trust in all of the team looking after them during their stay in hospital	68%	77%	80%	69%	77%	75%
Q32. Patient's family, or someone close, was definitely able to talk to a member of the team looking after the patient in hospital	57%	71%	68%	59%	67%	66%
Q33. Patient was always involved in decisions about their care and treatment whilst in hospital	60%	71%	69%	72%	70%	68%
Q34. Patient was always able to get help from ward staff when needed	61%	72%	76%	65%	81%	70%
Q35. Patient was always able to discuss worries and fears with hospital staff	51%	62%	69%	47%	76%	61%
Q36. Hospital staff always did everything they could to help the patient control pain	72%	82%	90%	89%	83%	83%
Q37. Patient was always treated with respect and dignity while in hospital	73%	84%	89%	91%	84%	84%
Q38. Patient received easily understandable information about what they should or should not do after leaving hospital	87%	89%	93%	93%	79%	89%
Q39. Patient was always able to discuss worries and fears with hospital staff while being treated as an outpatient or day case	65%	76%	74%	70%	71%	72%

## IMD quintile tables

\* Indicates where a score is not available due to suppression or a low base size.

YOUR TREATMENT	IMD quintile					
	1 (most deprived)	2	3	4	5 (least deprived)	All
Q41_1. Beforehand patient completely had enough understandable information about surgery	81%	93%	90%	94%	96%	<b>90%</b>
Q41_2. Beforehand patient completely had enough understandable information about chemotherapy	85%	86%	90%	88%	88%	<b>87%</b>
Q41_3. Beforehand patient completely had enough understandable information about radiotherapy	80%	91%	91%	95%	93%	<b>89%</b>
Q41_4. Beforehand patient completely had enough understandable information about hormone therapy	59%	81%	76%	78%	53%	<b>73%</b>
Q41_5. Beforehand patient completely had enough understandable information about immunotherapy	67%	82%	75%	82%	63%	<b>75%</b>
Q42_1. Patient completely had enough understandable information about their response to surgery	74%	85%	85%	94%	87%	<b>84%</b>
Q42_2. Patient completely had enough understandable information about their response to chemotherapy	78%	79%	75%	88%	82%	<b>80%</b>
Q42_3. Patient completely had enough understandable information about their response to radiotherapy	76%	89%	84%	86%	92%	<b>85%</b>
Q42_4. Patient completely had enough understandable information about their response to hormone therapy	61%	76%	80%	76%	53%	<b>72%</b>
Q42_5. Patient completely had enough understandable information about their response to immunotherapy	72%	85%	69%	88%	50%	<b>75%</b>
Q43. Patient felt the length of waiting time at clinic and day unit for cancer treatment was about right	62%	62%	61%	63%	65%	<b>62%</b>

IMMEDIATE AND LONG TERM SIDE EFFECTS	IMD quintile					
	1 (most deprived)	2	3	4	5 (least deprived)	All
Q44. Possible side effects from treatment were definitely explained in a way the patient could understand	69%	72%	73%	76%	74%	<b>73%</b>
Q45. Patient was always offered practical advice on dealing with any immediate side effects from treatment	66%	66%	65%	69%	72%	<b>67%</b>
Q46. Patient was given information that they could access about support in dealing with immediate side effects from treatment	83%	82%	86%	83%	88%	<b>84%</b>
Q47. Patient felt possible long-term side effects were definitely explained in a way they could understand in advance of their treatment	61%	59%	59%	53%	64%	<b>59%</b>
Q48. Patient was definitely able to discuss options for managing the impact of any long-term side effects	51%	51%	50%	47%	59%	<b>51%</b>

SUPPORT WHILE AT HOME	IMD quintile					
	1 (most deprived)	2	3	4	5 (least deprived)	All
Q49. Care team gave family, or someone close, all the information needed to help care for the patient at home	57%	56%	60%	59%	59%	<b>58%</b>
Q50. During treatment, the patient definitely got enough care and support at home from community or voluntary services	37%	37%	49%	46%	43%	<b>41%</b>

CARE FROM YOUR GP PRACTICE	IMD quintile					
	1 (most deprived)	2	3	4	5 (least deprived)	All
Q51. Patient definitely received the right amount of support from their GP practice during treatment	42%	39%	49%	38%	42%	<b>41%</b>
Q52. Patient has had a review of cancer care by GP practice	29%	26%	25%	22%	31%	<b>26%</b>

## IMD quintile tables

\* Indicates where a score is not available due to suppression or a low base size.

	IMD quintile					
	1 (most deprived)	2	3	4	5 (least deprived)	All
<b>Q53. After treatment, the patient definitely could get enough emotional support at home from community or voluntary services</b>	39%	26%	30%	22%	40%	<b>30%</b>
<b>Q54. The right amount of information and support was offered to the patient between final treatment and the follow up appointment</b>	73%	72%	81%	78%	82%	<b>76%</b>
<b>Q55. Patient was given enough information about the possibility and signs of cancer coming back or spreading</b>	57%	53%	55%	61%	61%	<b>56%</b>

	IMD quintile					
	1 (most deprived)	2	3	4	5 (least deprived)	All
<b>Q56. The whole care team worked well together</b>	84%	88%	89%	92%	88%	<b>88%</b>
<b>Q57. Administration of care was very good or good</b>	83%	85%	85%	85%	86%	<b>85%</b>
<b>Q58. Cancer research opportunities were discussed with patient</b>	52%	54%	53%	47%	42%	<b>51%</b>
<b>Q59. Patient's average rating of care scored from very poor to very good</b>	8.5	8.6	8.7	8.9	8.8	<b>8.7</b>

## Long-term condition status tables

\* Indicates where a score is not available due to suppression or a low base size.

	Long-term condition status			
	Yes	No	Not given	All
<b>SUPPORT FROM YOUR GP PRACTICE</b>				
Q2. Patient only spoke to primary care professional once or twice before cancer diagnosis	70%	76%	75%	72%
Q3. Referral for diagnosis was explained in a way the patient could completely understand	56%	67%	57%	60%

	Long-term condition status			
	Yes	No	Not given	All
<b>DIAGNOSTIC TESTS</b>				
Q5. Patient received all the information needed about the diagnostic test in advance	87%	92%	88%	88%
Q6. Diagnostic test staff appeared to completely have all the information they needed about the patient	74%	83%	79%	77%
Q7. Patient felt the length of time waiting for diagnostic test results was about right	69%	75%	74%	72%
Q8. Diagnostic test results were explained in a way the patient could completely understand	73%	77%	80%	75%
Q9. Enough privacy was always given to the patient when receiving diagnostic test results	92%	93%	95%	93%

	Long-term condition status			
	Yes	No	Not given	All
<b>FINDING OUT THAT YOU HAD CANCER</b>				
Q12. Patient was told they could have a family member, carer or friend with them when told diagnosis	77%	79%	79%	78%
Q13. Patient was definitely told sensitively that they had cancer	71%	79%	74%	74%
Q14. Cancer diagnosis explained in a way the patient could completely understand	74%	80%	79%	76%
Q15. Patient was definitely told about their diagnosis in an appropriate place	85%	91%	81%	86%
Q16. Patient was told they could go back later for more information about their diagnosis	80%	85%	77%	81%

	Long-term condition status			
	Yes	No	Not given	All
<b>SUPPORT FROM A MAIN CONTACT PERSON</b>				
Q17. Patient had a main point of contact within the care team	91%	95%	95%	93%
Q18. Patient found it very or quite easy to contact their main contact person	77%	82%	89%	80%
Q19. Patient found advice from main contact person was very or quite helpful	93%	93%	95%	93%

	Long-term condition status			
	Yes	No	Not given	All
<b>DECIDING ON THE BEST TREATMENT</b>				
Q20. Treatment options were explained in a way the patient could completely understand	79%	80%	81%	79%
Q21. Patient was definitely involved as much as they wanted to be in decisions about their treatment	74%	79%	81%	76%
Q22. Family and/or carers were definitely involved as much as the patient wanted them to be in decisions about treatment options	82%	82%	90%	83%
Q23. Patient could get further advice from a different healthcare professional before making decisions about their treatment options	50%	62%	59%	55%



## Long-term condition status tables

\* Indicates where a score is not available due to suppression or a low base size.

<b>CARE PLANNING</b>	Long-term condition status			
	Yes	No	Not given	All
Q24. Patient was definitely able to have a discussion about their needs or concerns prior to treatment	66%	72%	75%	69%
Q25. A member of their care team helped the patient create a care plan to address any needs or concerns	88%	93%	91%	90%
Q26. Care team reviewed the patient's care plan with them to ensure it was up to date	97%	98%	100%	98%

<b>SUPPORT FROM HOSPITAL STAFF</b>	Long-term condition status			
	Yes	No	Not given	All
Q27. Staff provided the patient with relevant information on available support	89%	93%	96%	91%
Q28. Patient definitely got the right level of support for their overall health and well being from hospital staff	66%	76%	83%	71%
Q29. Patient was offered information about how to get financial help or benefits	63%	76%	71%	68%

<b>HOSPITAL CARE</b>	Long-term condition status			
	Yes	No	Not given	All
Q31. Patient had confidence and trust in all of the team looking after them during their stay in hospital	70%	82%	78%	75%
Q32. Patient's family, or someone close, was definitely able to talk to a member of the team looking after the patient in hospital	61%	70%	79%	66%
Q33. Patient was always involved in decisions about their care and treatment whilst in hospital	68%	69%	71%	68%
Q34. Patient was always able to get help from ward staff when needed	69%	74%	69%	70%
Q35. Patient was always able to discuss worries and fears with hospital staff	56%	70%	59%	61%
Q36. Hospital staff always did everything they could to help the patient control pain	79%	88%	86%	83%
Q37. Patient was always treated with respect and dignity while in hospital	81%	90%	81%	84%
Q38. Patient received easily understandable information about what they should or should not do after leaving hospital	86%	96%	89%	89%
Q39. Patient was always able to discuss worries and fears with hospital staff while being treated as an outpatient or day case	70%	73%	78%	72%

## Long-term condition status tables

\* Indicates where a score is not available due to suppression or a low base size.

YOUR TREATMENT	Long-term condition status			
	Yes	No	Not given	All
Q41_1. Beforehand patient completely had enough understandable information about surgery	90%	92%	89%	90%
Q41_2. Beforehand patient completely had enough understandable information about chemotherapy	84%	90%	93%	87%
Q41_3. Beforehand patient completely had enough understandable information about radiotherapy	88%	91%	92%	89%
Q41_4. Beforehand patient completely had enough understandable information about hormone therapy	74%	72%	73%	73%
Q41_5. Beforehand patient completely had enough understandable information about immunotherapy	76%	69%	91%	75%
Q42_1. Patient completely had enough understandable information about their response to surgery	83%	87%	84%	84%
Q42_2. Patient completely had enough understandable information about their response to chemotherapy	76%	84%	85%	80%
Q42_3. Patient completely had enough understandable information about their response to radiotherapy	83%	87%	88%	85%
Q42_4. Patient completely had enough understandable information about their response to hormone therapy	71%	71%	82%	72%
Q42_5. Patient completely had enough understandable information about their response to immunotherapy	69%	83%	*	75%
Q43. Patient felt the length of waiting time at clinic and day unit for cancer treatment was about right	59%	67%	64%	62%

IMMEDIATE AND LONG TERM SIDE EFFECTS	Long-term condition status			
	Yes	No	Not given	All
Q44. Possible side effects from treatment were definitely explained in a way the patient could understand	69%	79%	74%	73%
Q45. Patient was always offered practical advice on dealing with any immediate side effects from treatment	63%	71%	76%	67%
Q46. Patient was given information that they could access about support in dealing with immediate side effects from treatment	82%	86%	88%	84%
Q47. Patient felt possible long-term side effects were definitely explained in a way they could understand in advance of their treatment	54%	69%	60%	59%
Q48. Patient was definitely able to discuss options for managing the impact of any long-term side effects	47%	56%	60%	51%

SUPPORT WHILE AT HOME	Long-term condition status			
	Yes	No	Not given	All
Q49. Care team gave family, or someone close, all the information needed to help care for the patient at home	54%	62%	68%	58%
Q50. During treatment, the patient definitely got enough care and support at home from community or voluntary services	37%	48%	51%	41%

CARE FROM YOUR GP PRACTICE	Long-term condition status			
	Yes	No	Not given	All
Q51. Patient definitely received the right amount of support from their GP practice during treatment	38%	47%	47%	41%
Q52. Patient has had a review of cancer care by GP practice	24%	29%	27%	26%

## Long-term condition status tables

\* Indicates where a score is not available due to suppression or a low base size.

<b>LIVING WITH AND BEYOND CANCER</b>	Long-term condition status			
	Yes	No	Not given	All
Q53. After treatment, the patient definitely could get enough emotional support at home from community or voluntary services	30%	35%	23%	<b>30%</b>
Q54. The right amount of information and support was offered to the patient between final treatment and the follow up appointment	73%	80%	80%	<b>76%</b>
Q55. Patient was given enough information about the possibility and signs of cancer coming back or spreading	55%	56%	65%	<b>56%</b>

<b>YOUR OVERALL NHS CARE</b>	Long-term condition status			
	Yes	No	Not given	All
Q56. The whole care team worked well together	86%	92%	91%	<b>88%</b>
Q57. Administration of care was very good or good	83%	86%	94%	<b>85%</b>
Q58. Cancer research opportunities were discussed with patient	48%	53%	60%	<b>51%</b>
Q59. Patient's average rating of care scored from very poor to very good	8.5	8.9	8.8	<b>8.7</b>

## Year on year charts

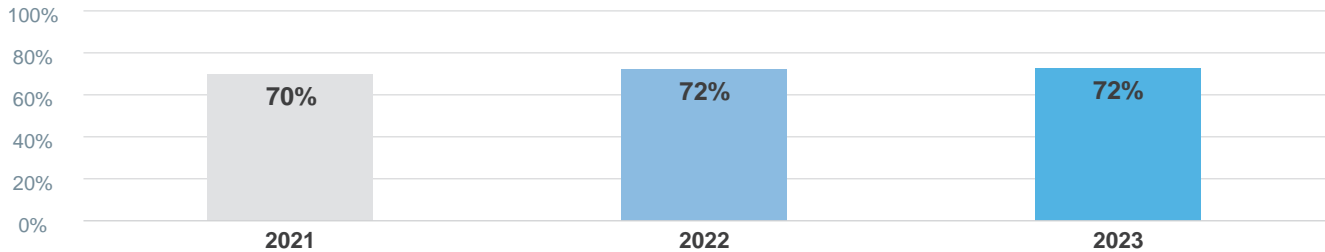
\* Indicates where a score is not available due to suppression or a low base size.

\*\* No score available for these years.

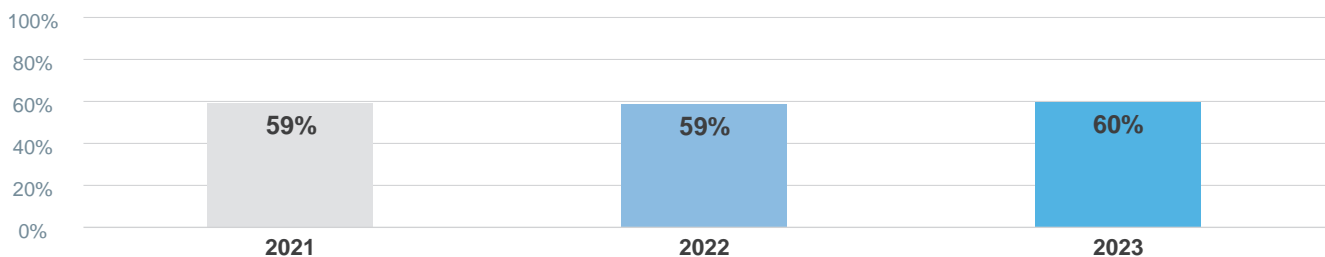
The scores are unadjusted and based on England scores only.

### SUPPORT FROM YOUR GP PRACTICE

Q2. Patient only spoke to primary care professional once or twice before cancer diagnosis

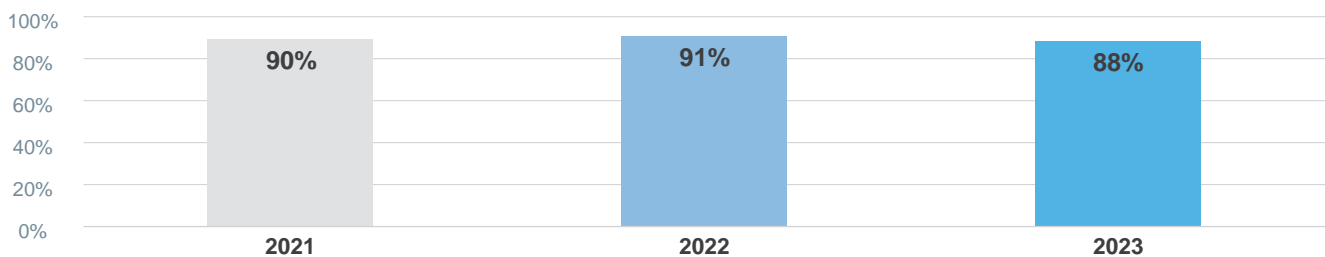


Q3. Referral for diagnosis was explained in a way the patient could completely understand

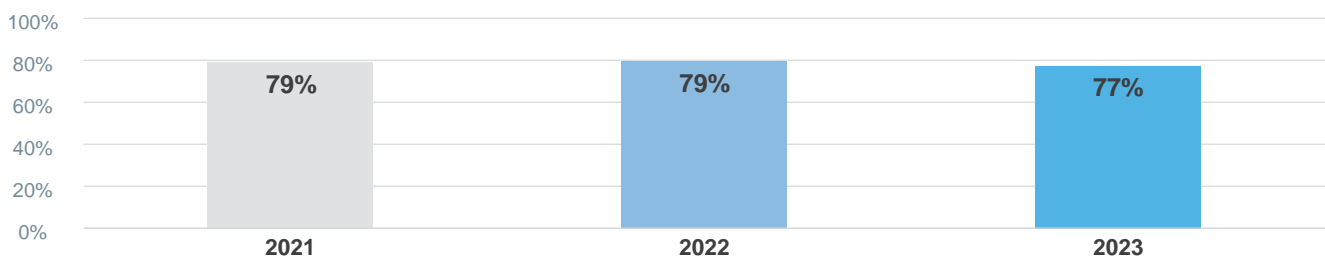


### DIAGNOSTIC TESTS

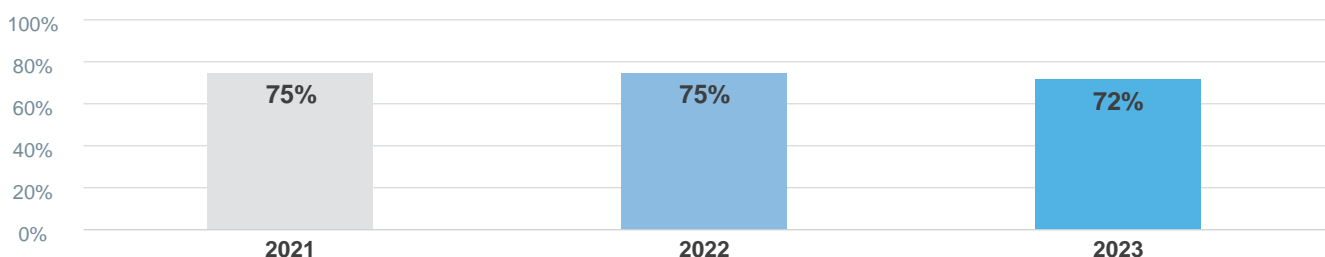
Q5. Patient received all the information needed about the diagnostic test in advance



Q6. Diagnostic test staff appeared to completely have all the information they needed about the patient



Q7. Patient felt the length of time waiting for diagnostic test results was about right



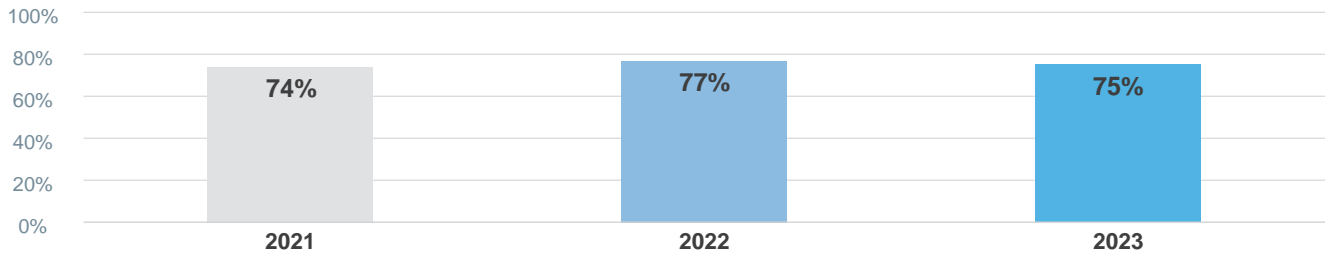
## Year on year charts

\* Indicates where a score is not available due to suppression or a low base size.

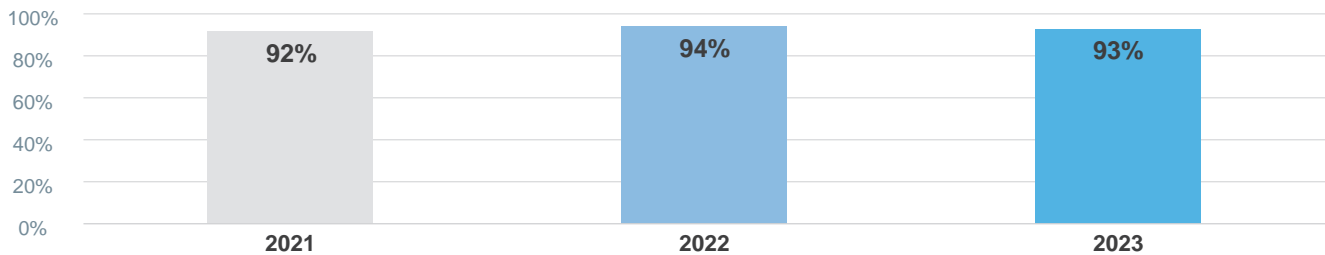
\*\* No score available for these years.

The scores are unadjusted and based on England scores only.

Q8. Diagnostic test results were explained in a way the patient could completely understand

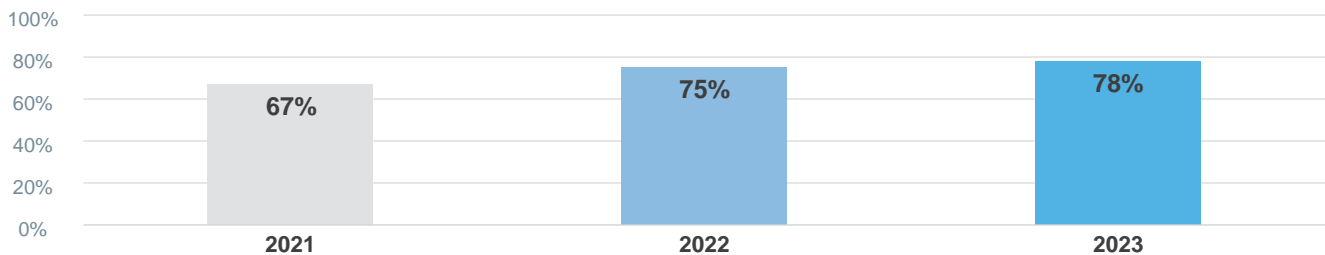


Q9. Enough privacy was always given to the patient when receiving diagnostic test results

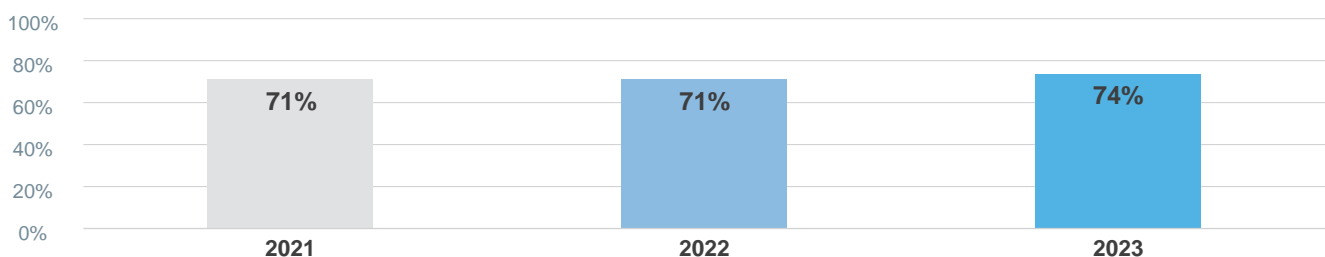


## FINDING OUT THAT YOU HAD CANCER

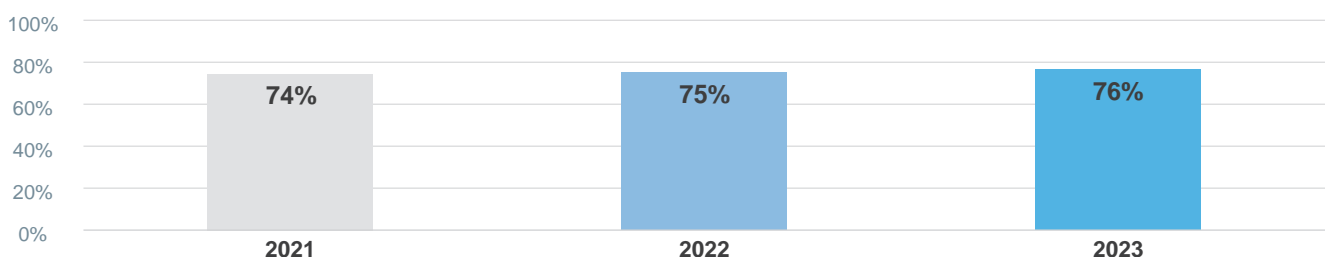
Q12. Patient was told they could have a family member, carer or friend with them when told diagnosis



Q13. Patient was definitely told sensitively that they had cancer



Q14. Cancer diagnosis explained in a way the patient could completely understand



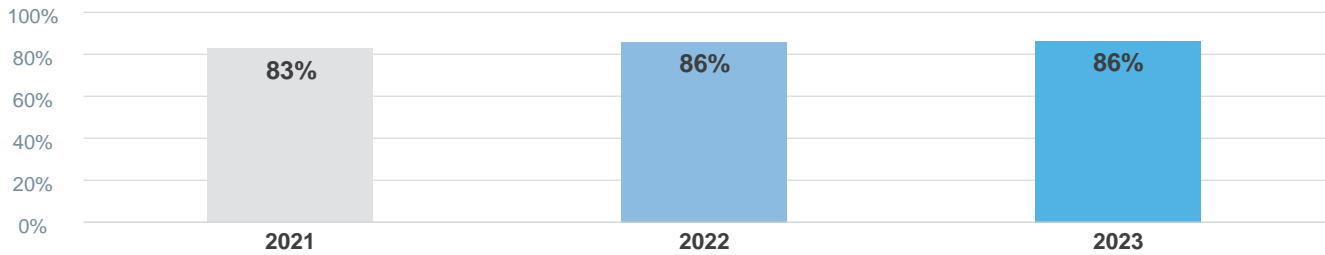
## Year on year charts

\* Indicates where a score is not available due to suppression or a low base size.

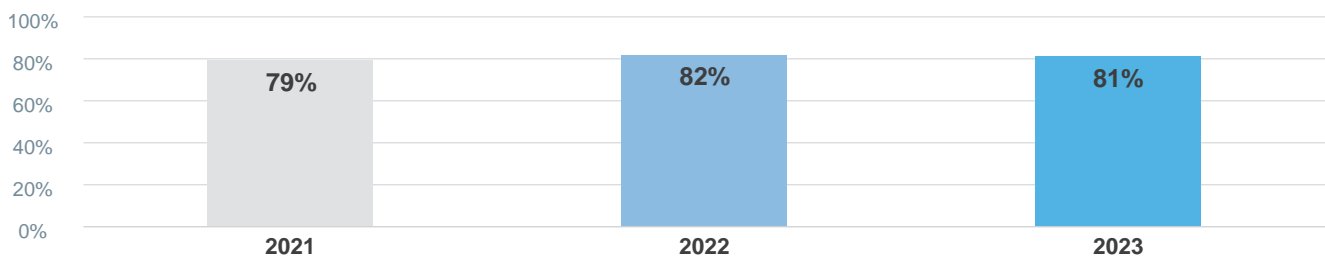
\*\* No score available for these years.

The scores are unadjusted and based on England scores only.

Q15. Patient was definitely told about their diagnosis in an appropriate place

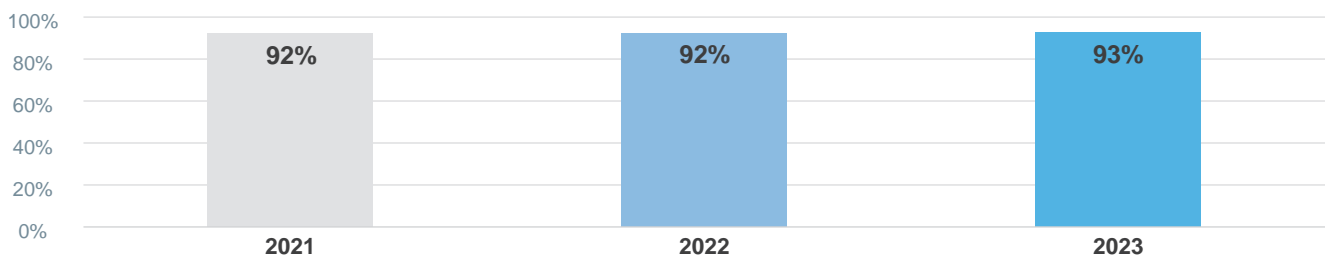


Q16. Patient was told they could go back later for more information about their diagnosis

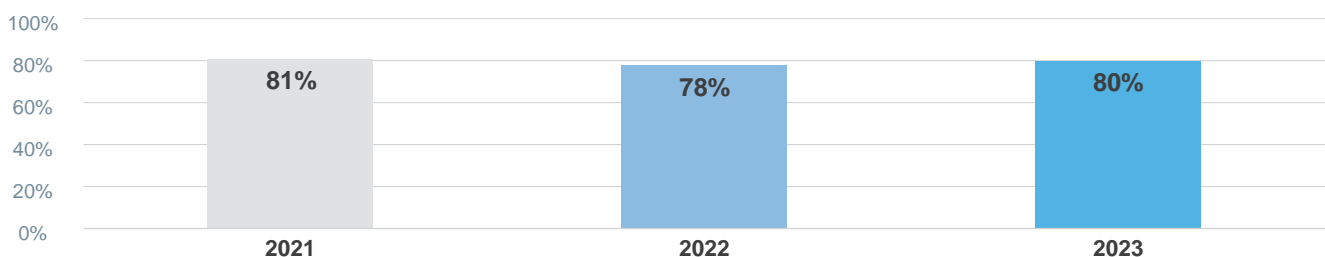


### SUPPORT FROM A MAIN CONTACT PERSON

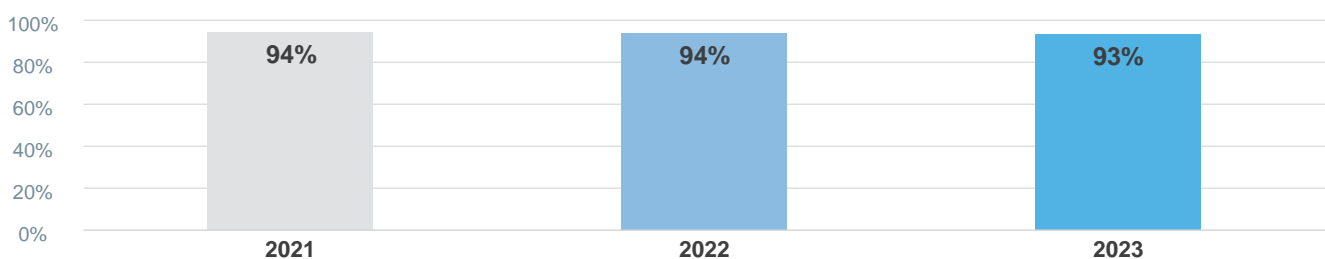
Q17. Patient had a main point of contact within the care team



Q18. Patient found it very or quite easy to contact their main contact person



Q19. Patient found advice from main contact person was very or quite helpful



## Year on year charts

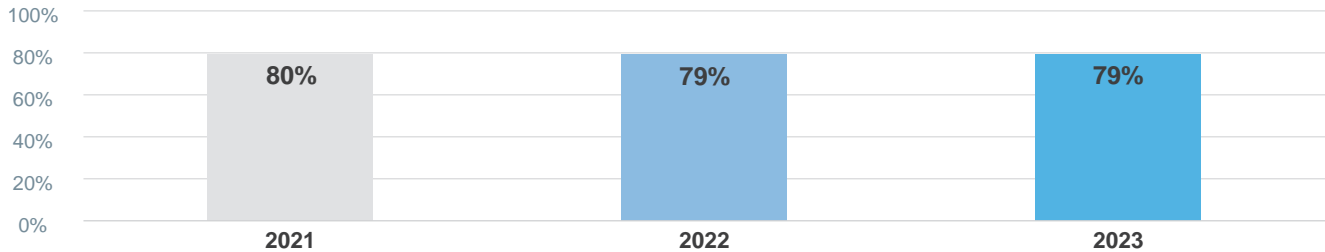
\* Indicates where a score is not available due to suppression or a low base size.

\*\* No score available for these years.

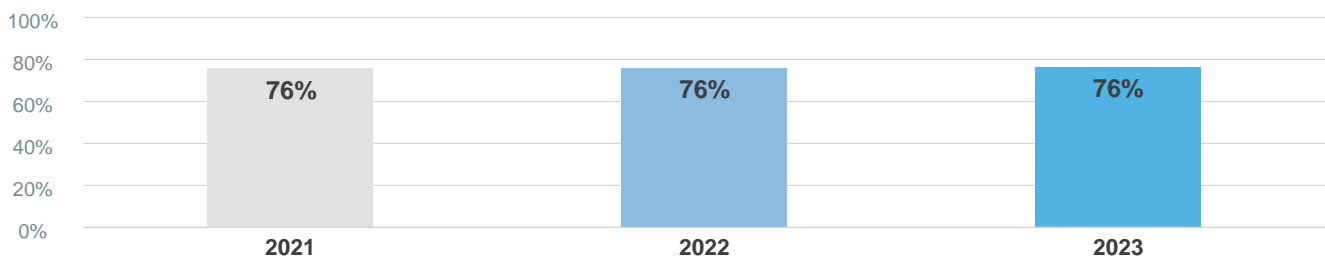
The scores are unadjusted and based on England scores only.

### DECIDING ON THE BEST TREATMENT

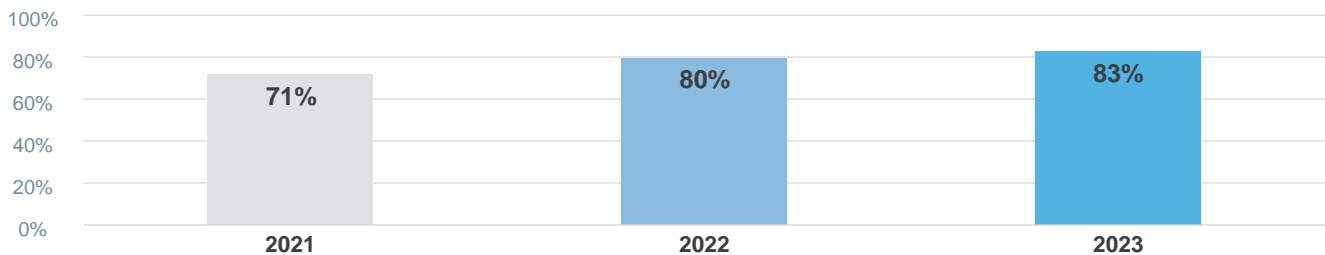
Q20. Treatment options were explained in a way the patient could completely understand



Q21. Patient was definitely involved as much as they wanted to be in decisions about their treatment



Q22. Family and/or carers were definitely involved as much as the patient wanted them to be in decisions about treatment options

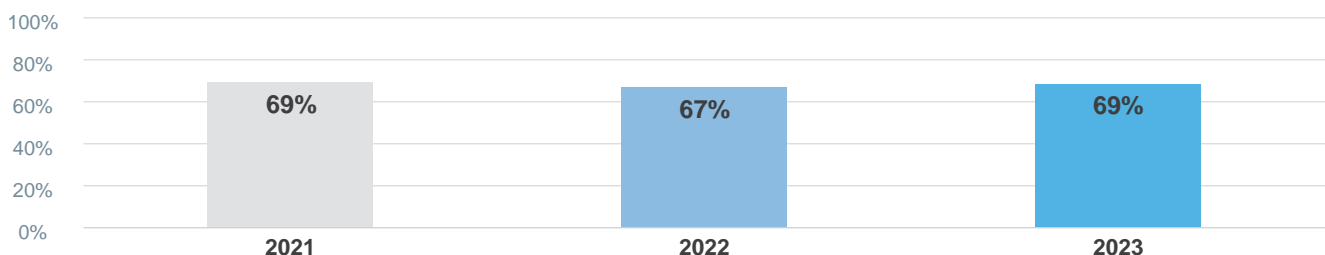


Q23. Patient could get further advice from a different healthcare professional before making decisions about their treatment options



### CARE PLANNING

Q24. Patient was definitely able to have a discussion about their needs or concerns prior to treatment



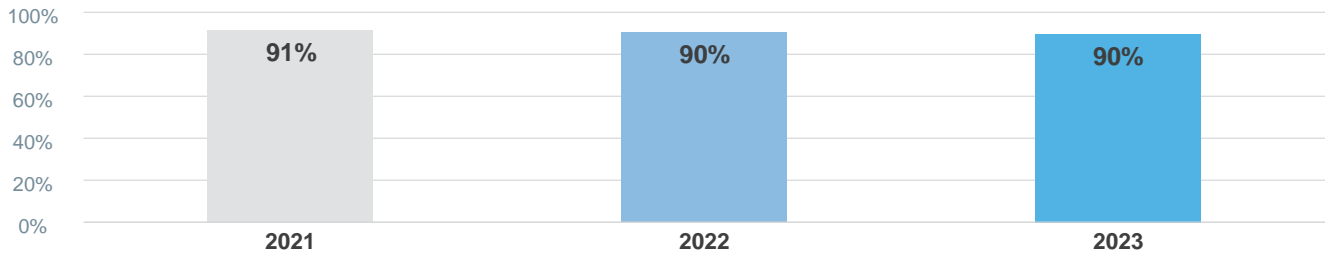
## Year on year charts

\* Indicates where a score is not available due to suppression or a low base size.

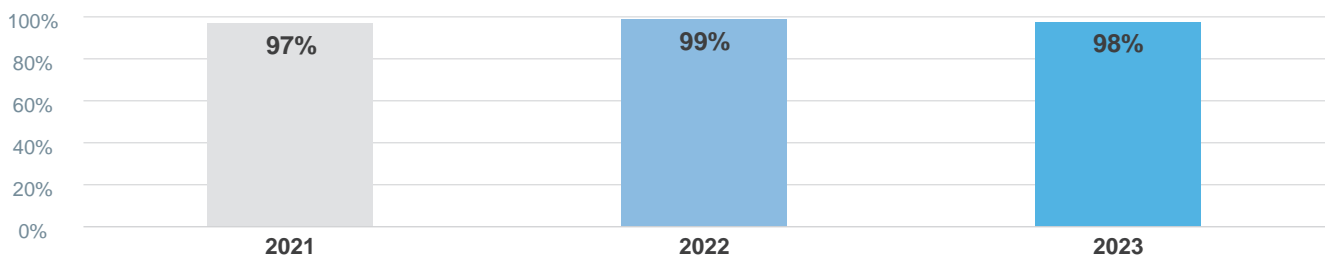
\*\* No score available for these years.

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Q25. A member of their care team helped the patient create a care plan to address any needs or concerns

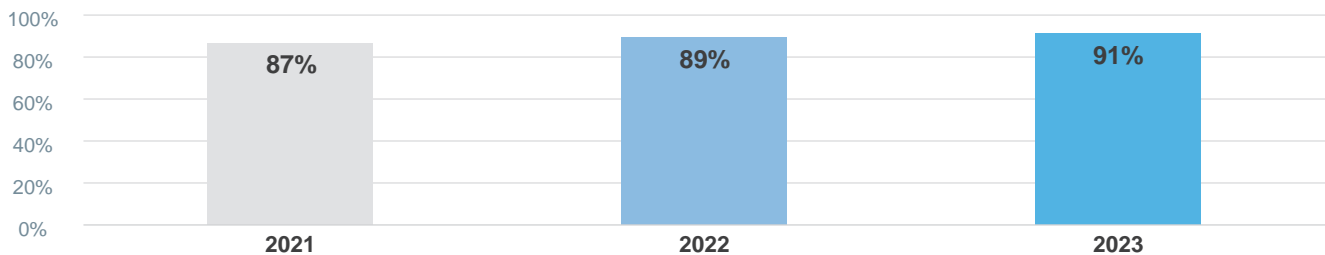


Q26. Care team reviewed the patient's care plan with them to ensure it was up to date

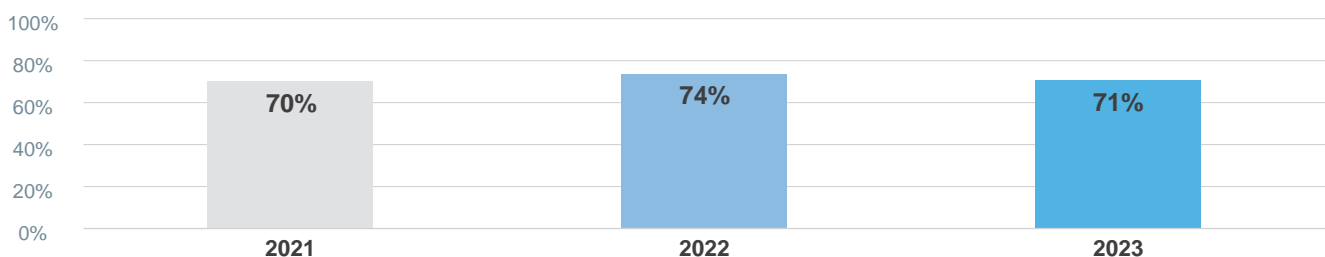


### SUPPORT FROM HOSPITAL STAFF

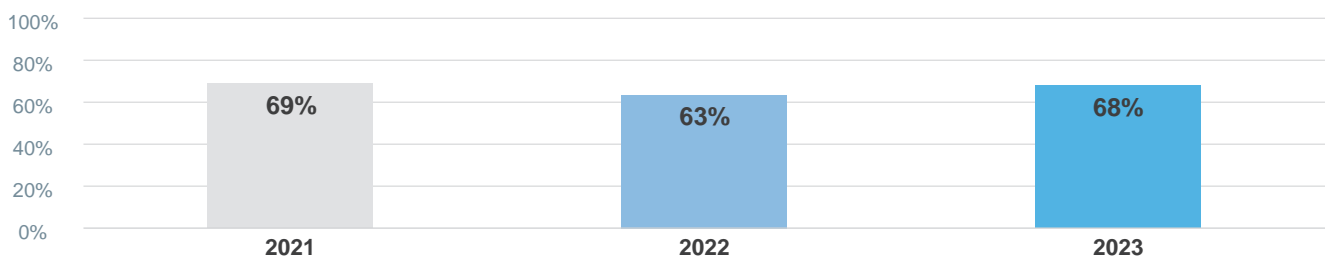
Q27. Staff provided the patient with relevant information on available support



Q28. Patient definitely got the right level of support for their overall health and well being from hospital staff



Q29. Patient was offered information about how to get financial help or benefits





## Year on year charts

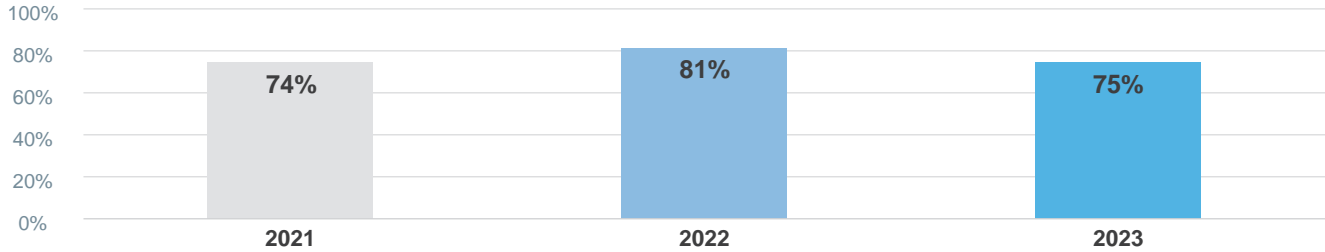
\* Indicates where a score is not available due to suppression or a low base size.

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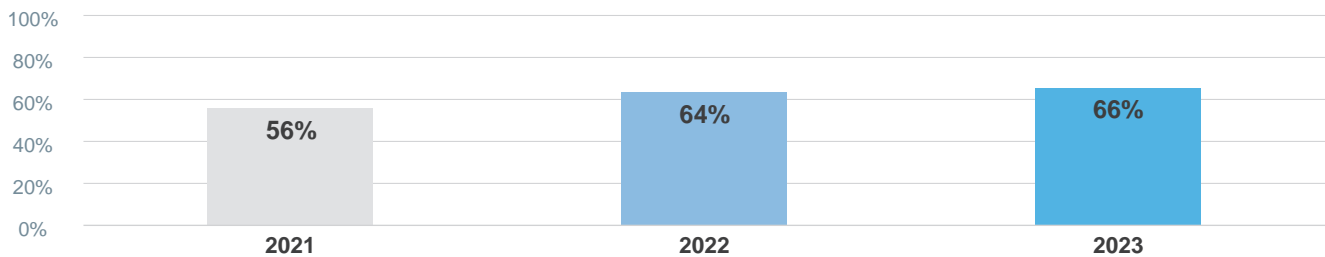
The scores are unadjusted and based on England scores only.

### HOSPITAL CARE

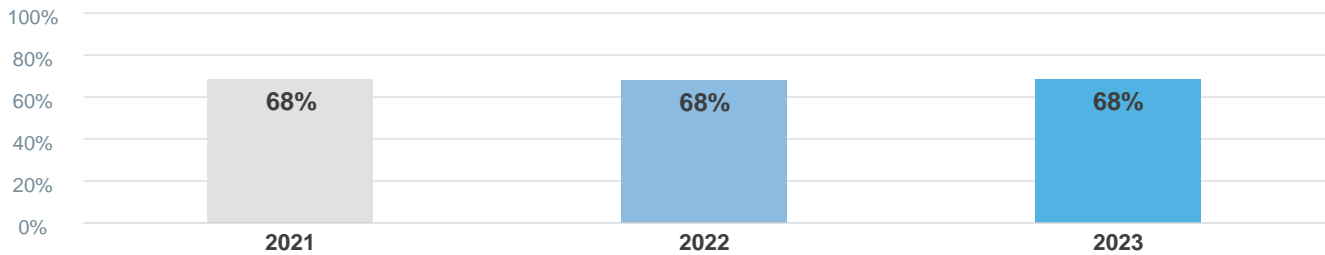
Q31. Patient had confidence and trust in all of the team looking after them during their stay in hospital



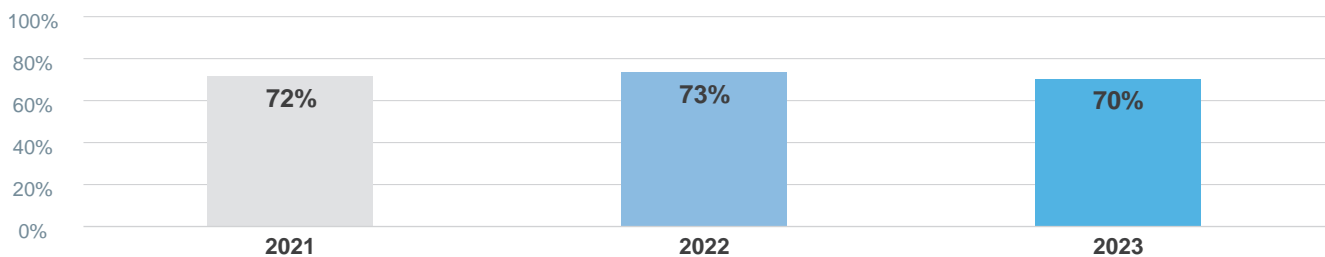
Q32. Patient's family, or someone close, was definitely able to talk to a member of the team looking after the patient in hospital



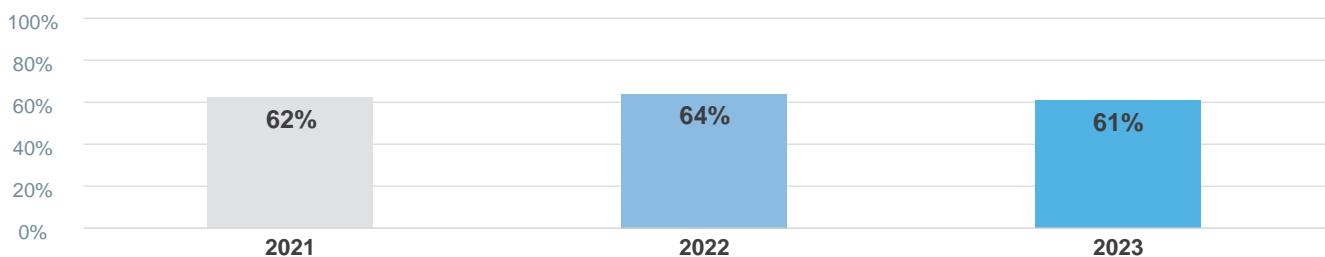
Q33. Patient was always involved in decisions about their care and treatment whilst in hospital



Q34. Patient was always able to get help from ward staff when needed



Q35. Patient was always able to discuss worries and fears with hospital staff

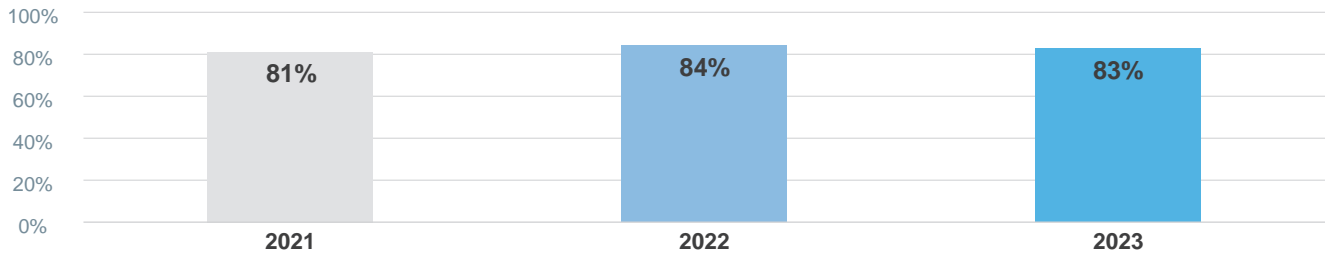


## Year on year charts

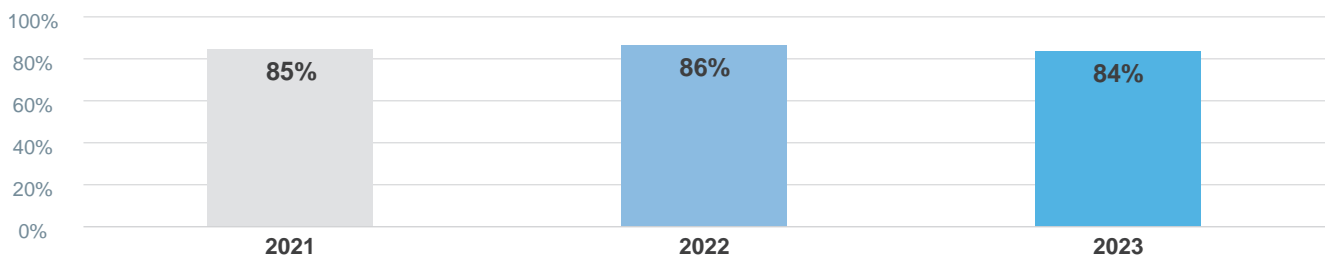
\* Indicates where a score is not available due to suppression or a low base size. \*\* No score available for these years.

The scores are unadjusted and based on England scores only.

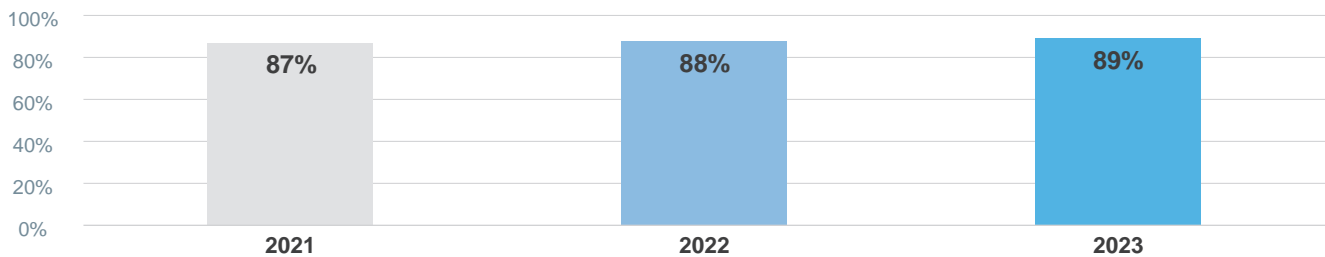
Q36. Hospital staff always did everything they could to help the patient control pain



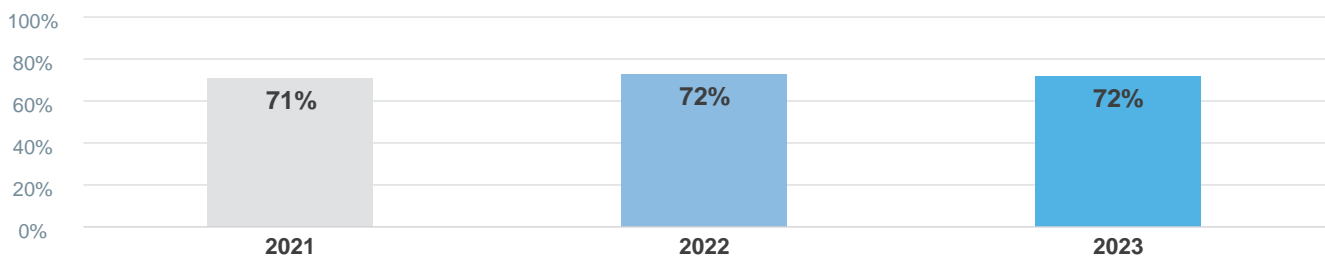
Q37. Patient was always treated with respect and dignity while in hospital



Q38. Patient received easily understandable information about what they should or should not do after leaving hospital

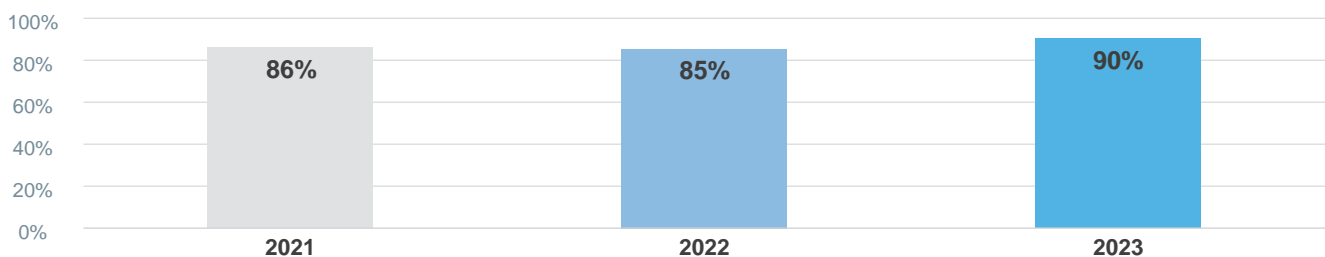


Q39. Patient was always able to discuss worries and fears with hospital staff while being treated as an outpatient or day case



### YOUR TREATMENT

Q41\_1. Beforehand patient completely had enough understandable information about surgery



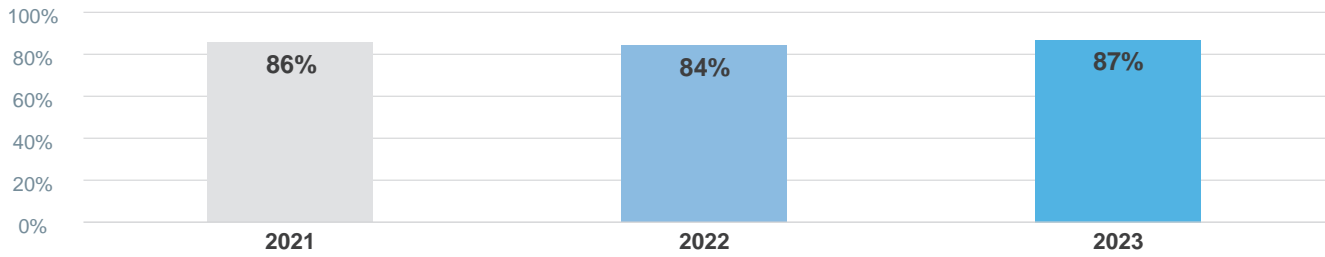
## Year on year charts

\* Indicates where a score is not available due to suppression or a low base size.

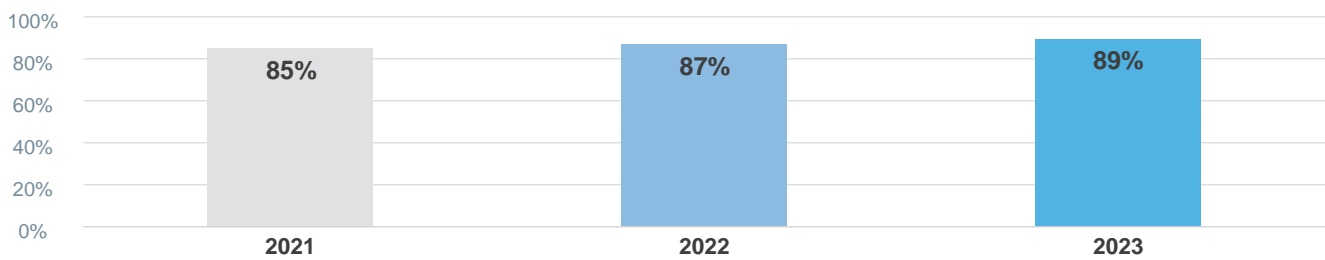
\*\* No score available for these years.

The scores are unadjusted and based on England scores only.

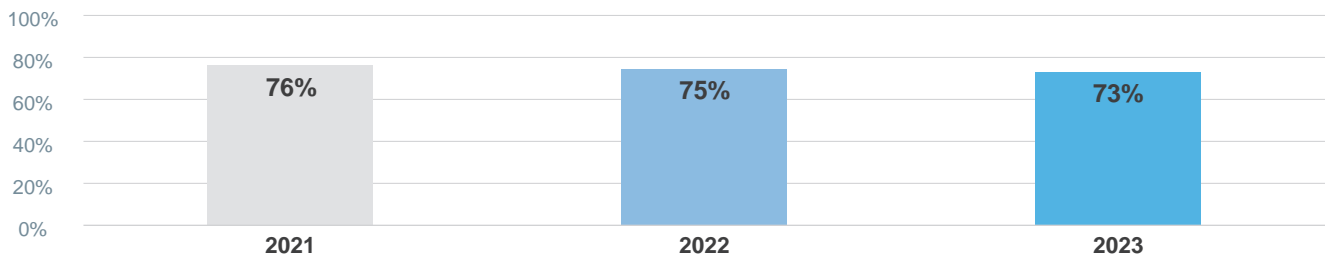
Q41\_2. Beforehand patient completely had enough understandable information about chemotherapy



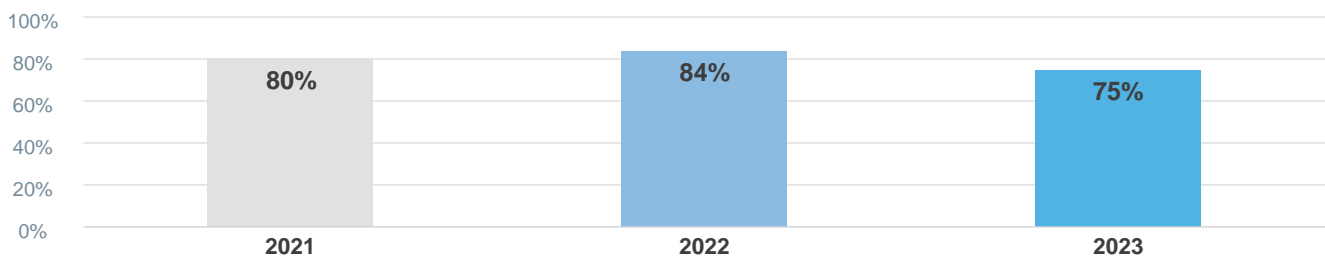
Q41\_3. Beforehand patient completely had enough understandable information about radiotherapy



Q41\_4. Beforehand patient completely had enough understandable information about hormone therapy



Q41\_5. Beforehand patient completely had enough understandable information about immunotherapy



Q42\_1. Patient completely had enough understandable information about their response to surgery



## Year on year charts

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\*\* No score available for these years.

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Q42\_2. Patient completely had enough understandable information about their response to chemotherapy



Q42\_3. Patient completely had enough understandable information about their response to radiotherapy



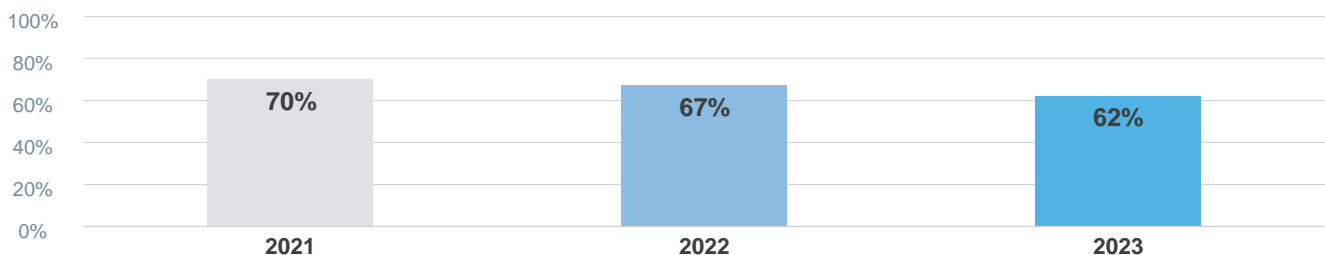
Q42\_4. Patient completely had enough understandable information about their response to hormone therapy



Q42\_5. Patient completely had enough understandable information about their response to immunotherapy



Q43. Patient felt the length of waiting time at clinic and day unit for cancer treatment was about right



## Year on year charts

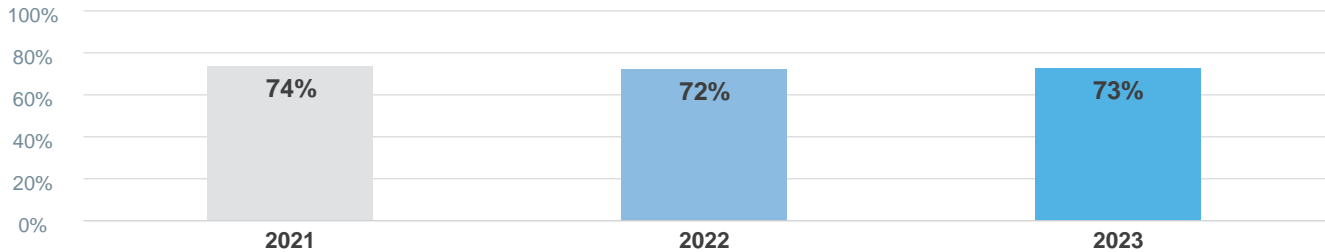
\* Indicates where a score is not available due to suppression or a low base size.

\*\* No score available for these years.

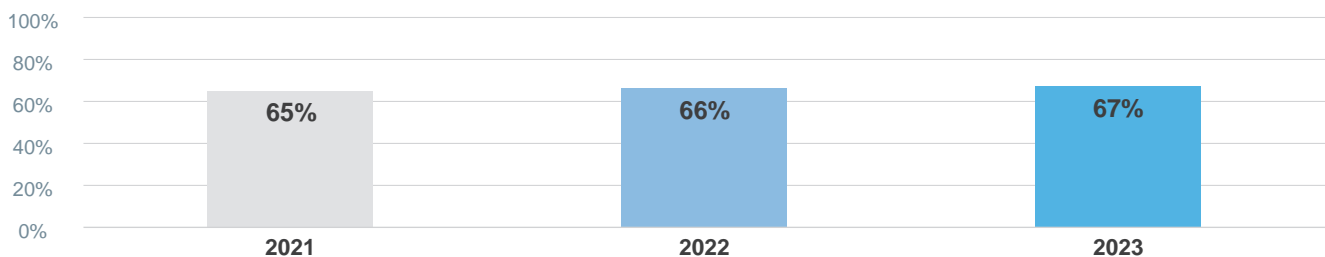
The scores are unadjusted and based on England scores only.

### IMMEDIATE AND LONG TERM SIDE EFFECTS

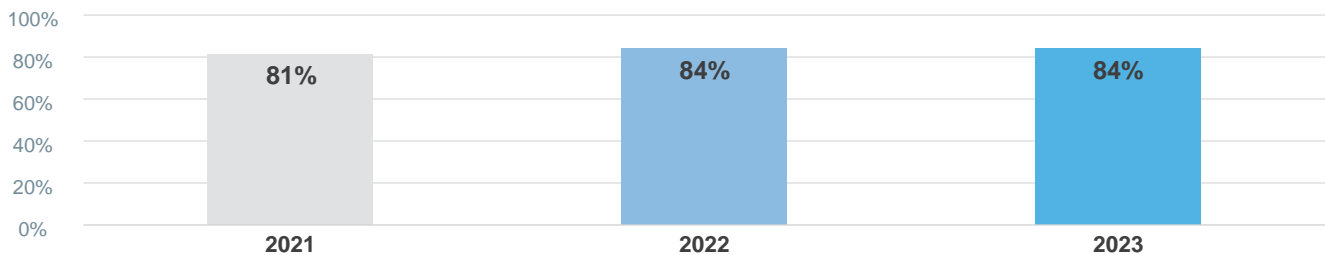
Q44. Possible side effects from treatment were definitely explained in a way the patient could understand



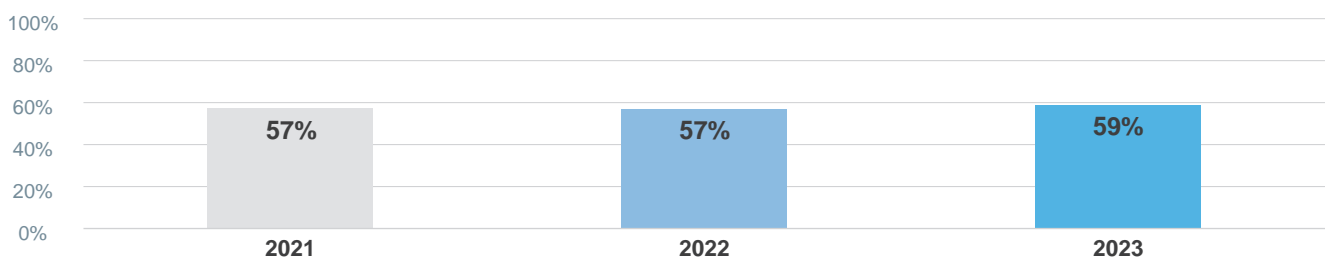
Q45. Patient was always offered practical advice on dealing with any immediate side effects from treatment



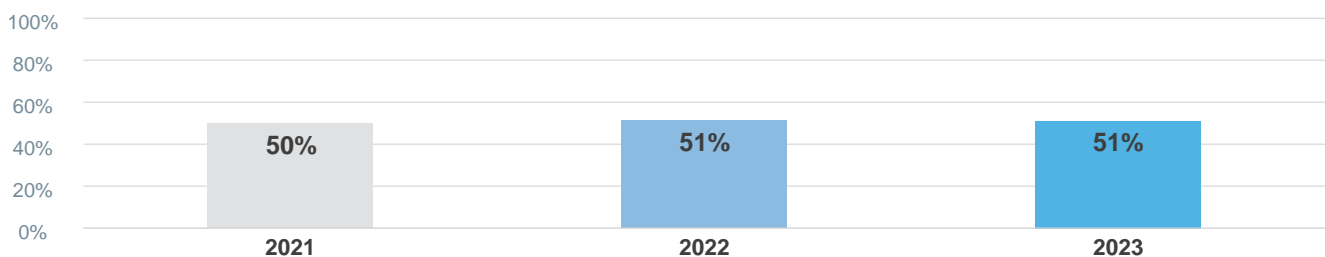
Q46. Patient was given information that they could access about support in dealing with immediate side effects from treatment



Q47. Patient felt possible long-term side effects were definitely explained in a way they could understand in advance of their treatment



Q48. Patient was definitely able to discuss options for managing the impact of any long-term side effects



## Year on year charts

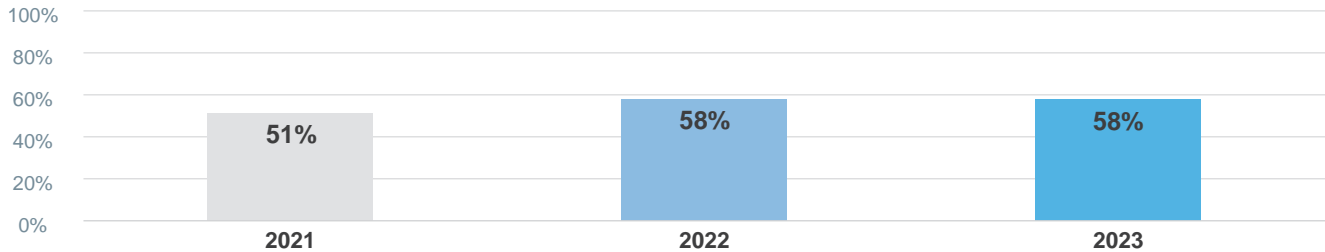
\* Indicates where a score is not available due to suppression or a low base size.

\*\* No score available for these years.

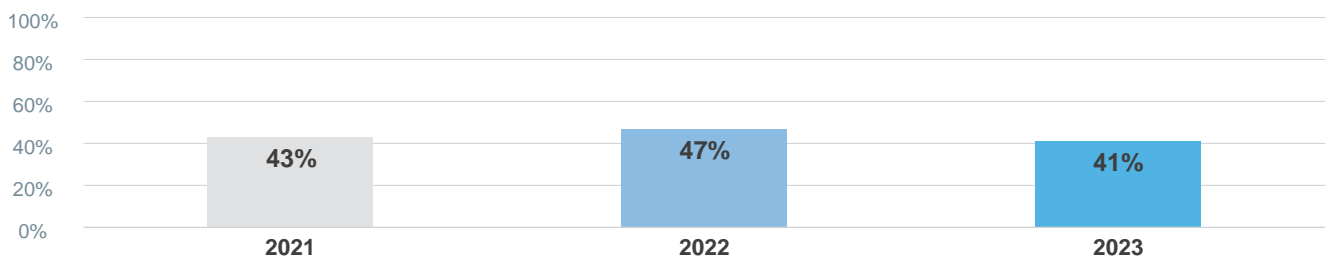
The scores are unadjusted and based on England scores only.

### SUPPORT WHILE AT HOME

Q49. Care team gave family, or someone close, all the information needed to help care for the patient at home

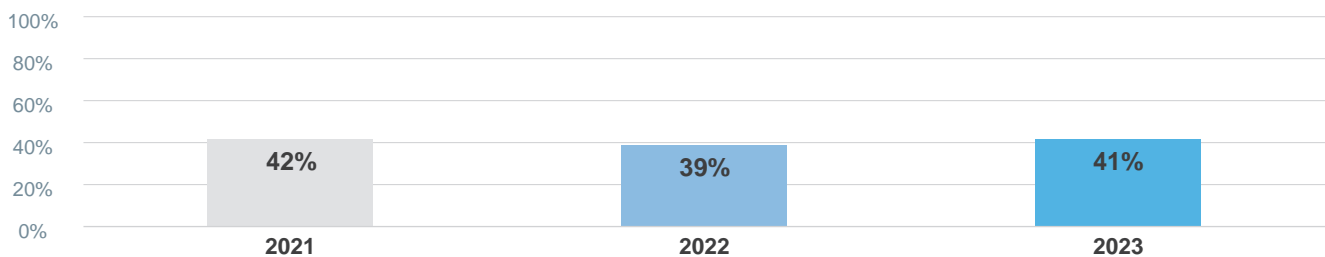


Q50. During treatment, the patient definitely got enough care and support at home from community or voluntary services

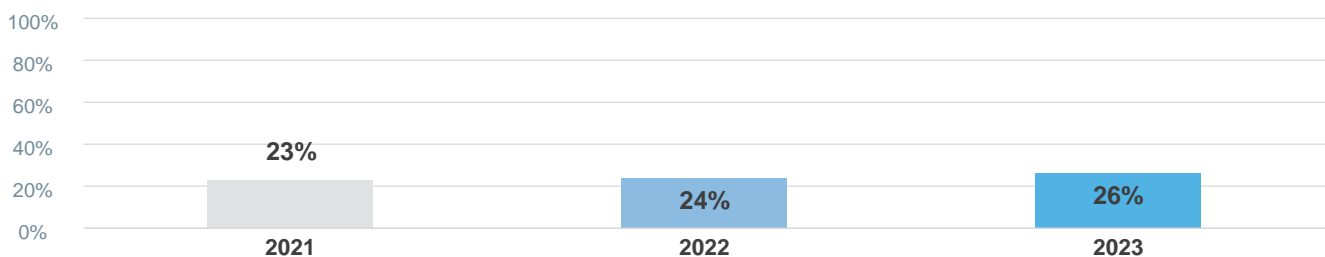


### CARE FROM YOUR GP PRACTICE

Q51. Patient definitely received the right amount of support from their GP practice during treatment

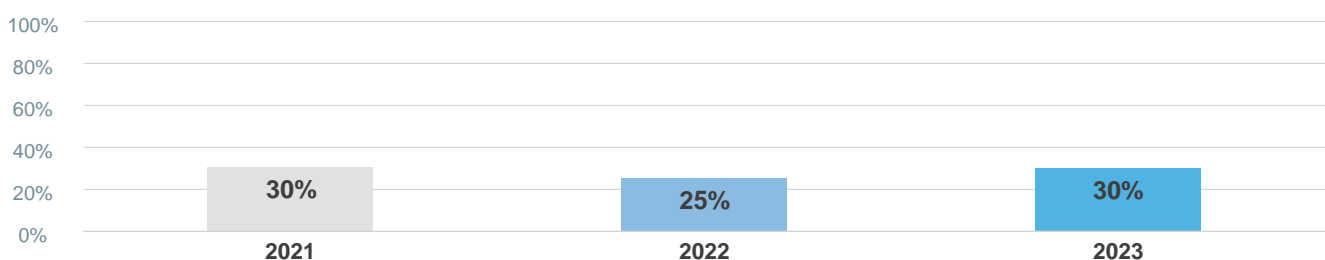


Q52. Patient has had a review of cancer care by GP practice



### LIVING WITH AND BEYOND CANCER

Q53. After treatment, the patient definitely could get enough emotional support at home from community or voluntary services



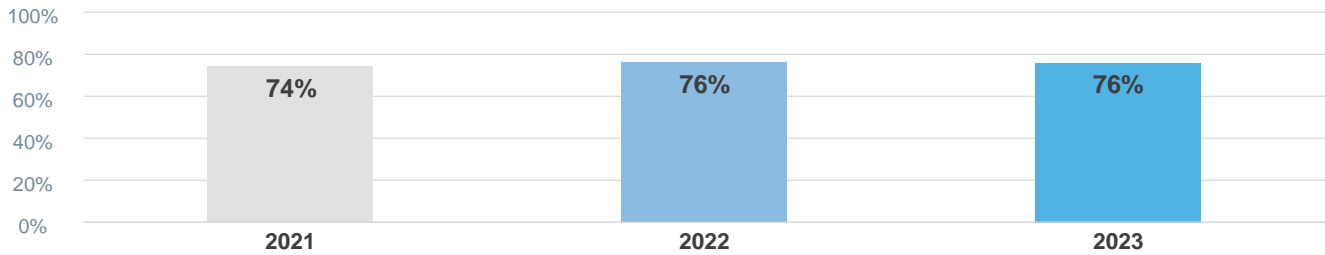
## Year on year charts

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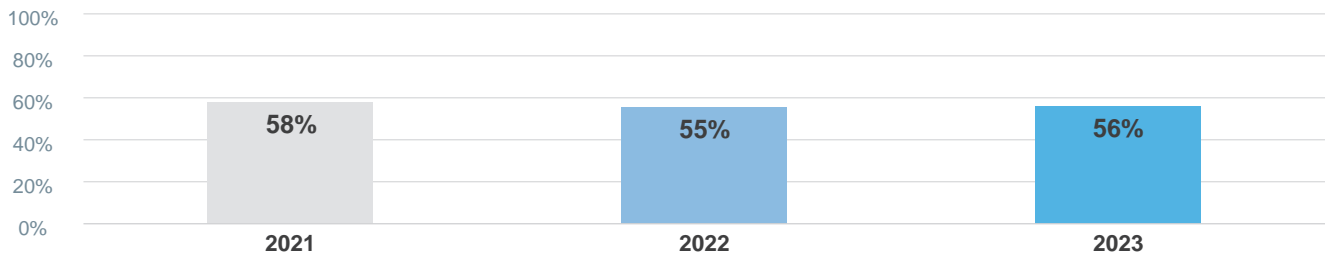
\*\* No score available for these years.

The scores are unadjusted and based on England scores only.

Q54. The right amount of information and support was offered to the patient between final treatment and the follow up appointment

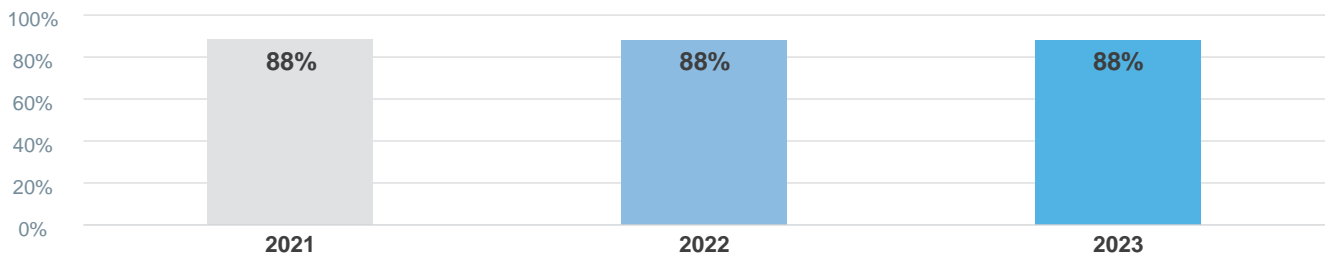


Q55. Patient was given enough information about the possibility and signs of cancer coming back or spreading

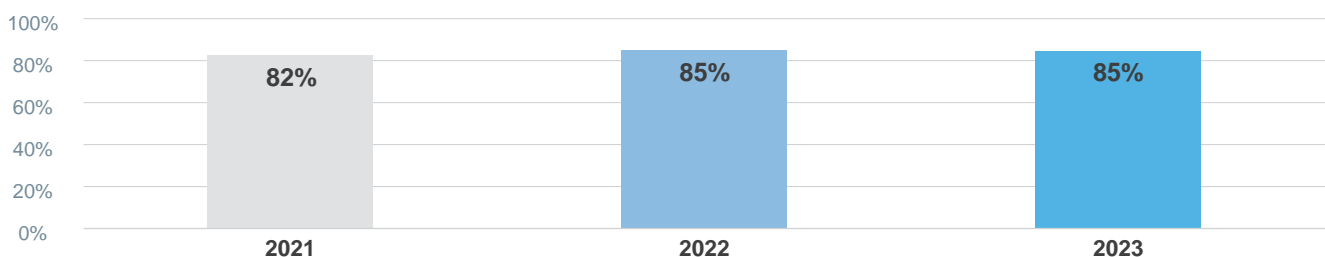


### YOUR OVERALL NHS CARE

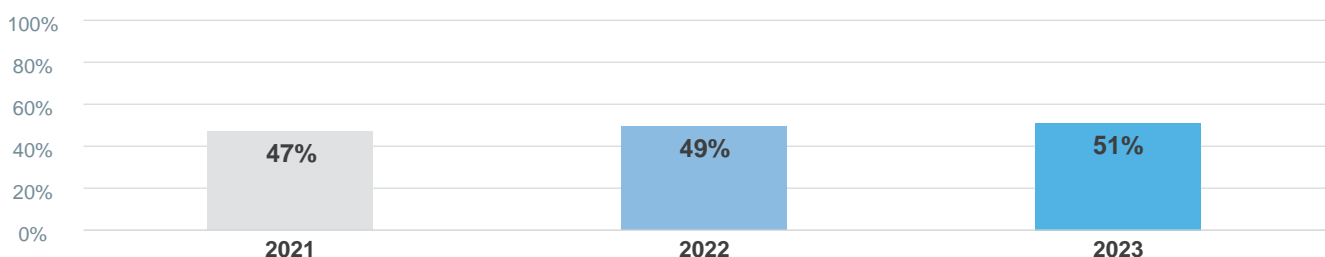
Q56. The whole care team worked well together



Q57. Administration of care was very good or good



Q58. Cancer research opportunities were discussed with patient



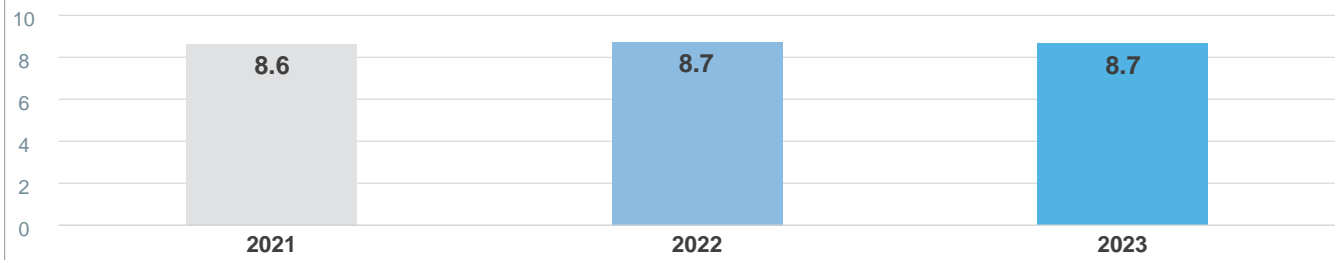
## Year on year charts

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\*\* No score available for these years.




The scores are unadjusted and based on England scores only.

Q59. Patient's average rating of care scored from very poor to very good








## Trust expected range summary

Data labels relate to the number of scores that fell below, within and above the expected range		Number of scores below the lower expected range
		Number of scores between the upper and lower expected ranges
		Number of scores above the upper expected range

Trust		Expected range classification
RQX	Homerton Healthcare NHS Foundation Trust	
RF4	Barking, Havering and Redbridge University Hospitals NHS Trust	
R1H	Barts Health NHS Trust	

## ICB expected range summary

<p>Data labels relate to the number of scores that fell below, within and above the expected range</p>		<p>Number of scores below the lower expected range</p>
		<p>Number of scores between the upper and lower expected ranges</p>
		<p>Number of scores above the upper expected range</p>

ICB		Expected range classification	
QMF	NHS North East London Integrated Care Board	15	46