

# **Cancer Patient Experience Survey**

2022 Results

## **Bradford Teaching Hospitals NHS Foundation Trust**

Published July 2023

## Executive Summary

### Questions Above Expected Range

	Case Mix Adjusted Scores			National Score
	2022 Score	Lower Expected Range	Upper Expected Range	
Q7. Patient felt the length of time waiting for diagnostic test results was about right	<b>85%</b>	72%	85%	<b>78%</b>

### Questions Below Expected Range

	Case Mix Adjusted Scores			National Score
	2022 Score	Lower Expected Range	Upper Expected Range	
Q5. Patient received all the information needed about the diagnostic test in advance	<b>88%</b>	88%	96%	<b>92%</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	<b>52%</b>	56%	76%	<b>66%</b>
Q41_2. Beforehand patient completely had enough understandable information about chemotherapy	<b>77%</b>	78%	92%	<b>85%</b>

## Introduction

The National Cancer Patient Experience Survey 2022 is the 12th 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 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 2022 survey involved 133 NHS Trusts. Out of 115,662 people, 61,268 people responded to the survey, yielding a response rate of 53%.

## 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 2022. The fieldwork for the survey was undertaken between November 2022 and February 2023.

As in the previous seven 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.

### 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 a Trust 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 Trust. 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%.

Please note that 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.

### Statistical significance

In the reporting of 2022 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 Trust, the results are not shown for that question for that Trust.

For Trusts 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** Trust has a score suppressed. If this happens, we will suppress another Trust's results (both the Trust level and subgroup 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 Trust.

The same rule applies to groups in each subgroup breakdown. For example, if only **one** Trust has the 85+ age group suppressed for Q25 we will need to suppress another Trust'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 Trust scored for each question in the survey compared with national results. It is aimed at helping individual Trusts 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 Trust.

Trusts 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 Trust performs better than what Trusts 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 Trust's size and demographics.

### **Comparability tables**

The comparability tables show the 2021 and 2022 unadjusted scores for this Trust for each scored question. If there is a statistically significant change from 2021 an arrow will be presented for the direction of change.

The adjusted 2022 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 type, Male/Female/Non-binary/Other, age, IMD quintile, long-term condition status and ethnicity breakdowns. Unadjusted scores for the same sub-group across different Trusts may not be comparable, as they do not account for the impact that differing patient populations might have on results.

### **Tumour type tables**

The tumour type 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 two columns representing the unadjusted scores of the last two years (2021 and 2022) for each scored question.

## **Further information**

This research was carried out in accordance with the international standard for organisations conducting social research (accreditation to ISO20252:2012; certificate number GB08/74322). The 2022 survey data has been produced and published in line with the Code of Practice for Official Statistics.

For more information on the methodology, please see the Technical Document. It can be viewed along with the 2022 questionnaire and survey guidance on the website at [www.ncpes.co.uk](http://www.ncpes.co.uk). For all other outputs at Trust level, please see the Excel tables and dashboards at [www.ncpes.co.uk](http://www.ncpes.co.uk).

## Response Rate

### Overall Response Rate

193 patients responded out of a total of 390 patients, resulting in a response rate of 49%.

	Sample Size	Adjusted Sample	Completed	Response Rate
Overall response rate	406	390	193	49%
National	123,632	115,662	61,268	53%

### Respondents by Survey Type

	Number of Respondents
Paper	151
Online	41
Phone	0
Translation Service	1
<b>Total</b>	<b>193</b>

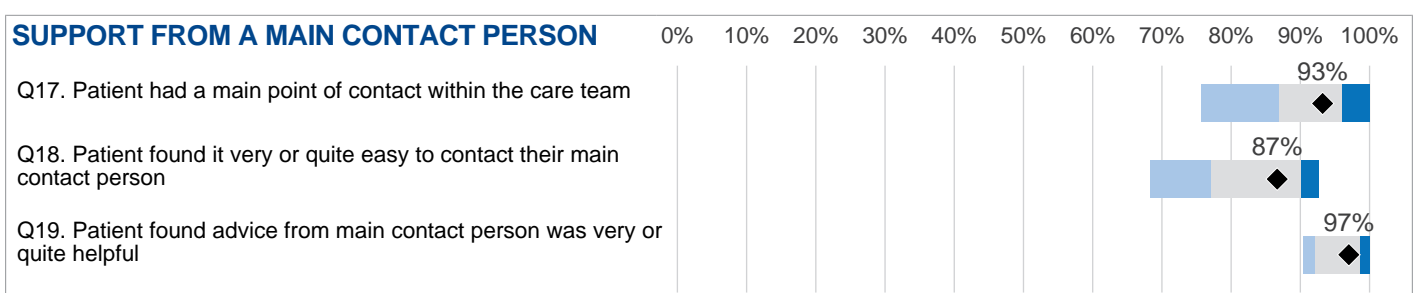
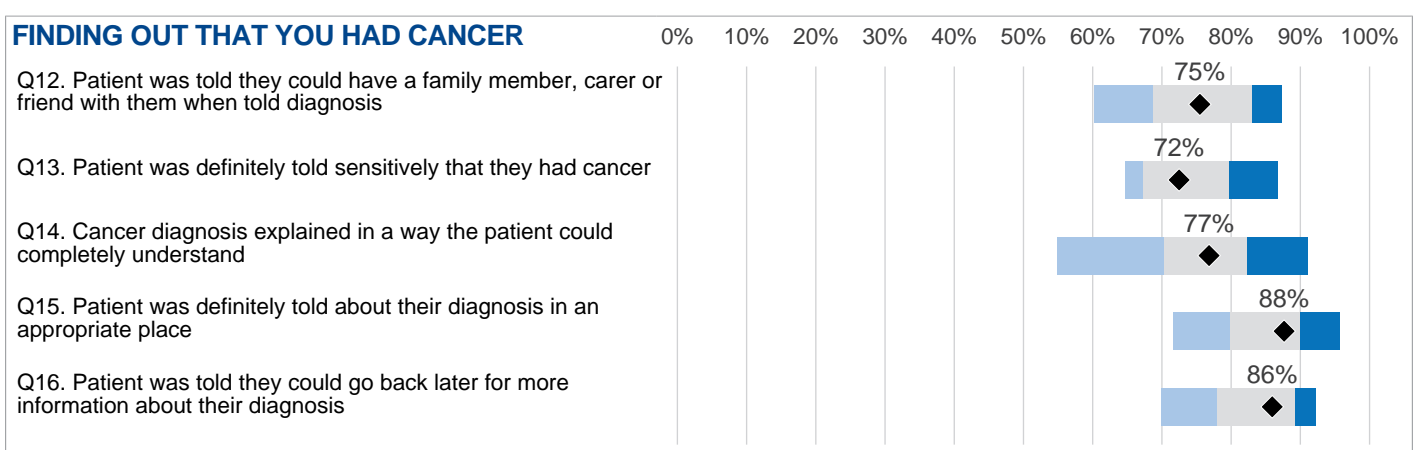
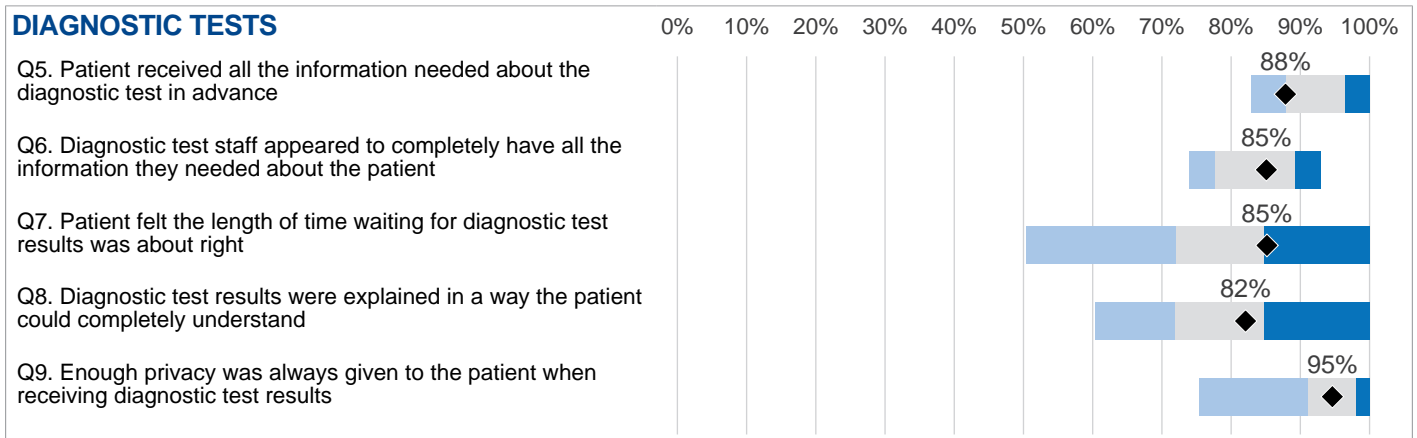
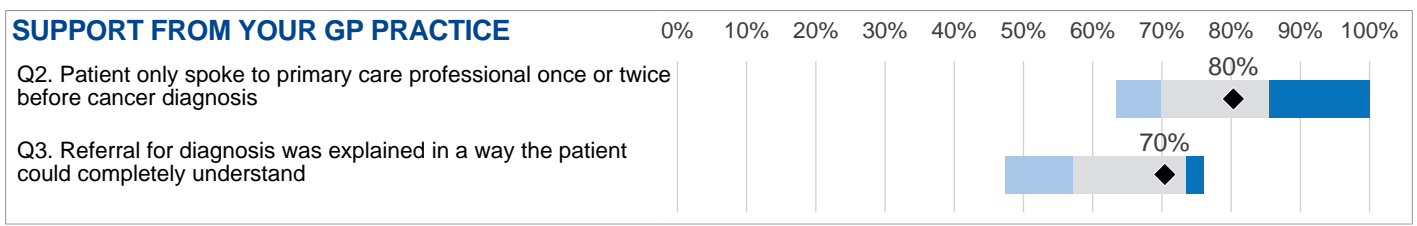
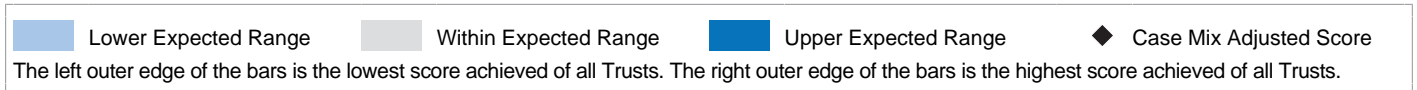
### Respondents by Tumour Group

	Number of Respondents
Brain / CNS	0
Breast	43
Colorectal / LGT	26
Gynaecological	10
Haematological	25
Head and Neck	6
Lung	8
Prostate	26
Sarcoma	*
Skin	*
Upper Gastro	11
Urological	12
Other	20
<b>Total</b>	<b>193</b>

## Respondents by Ethnicity

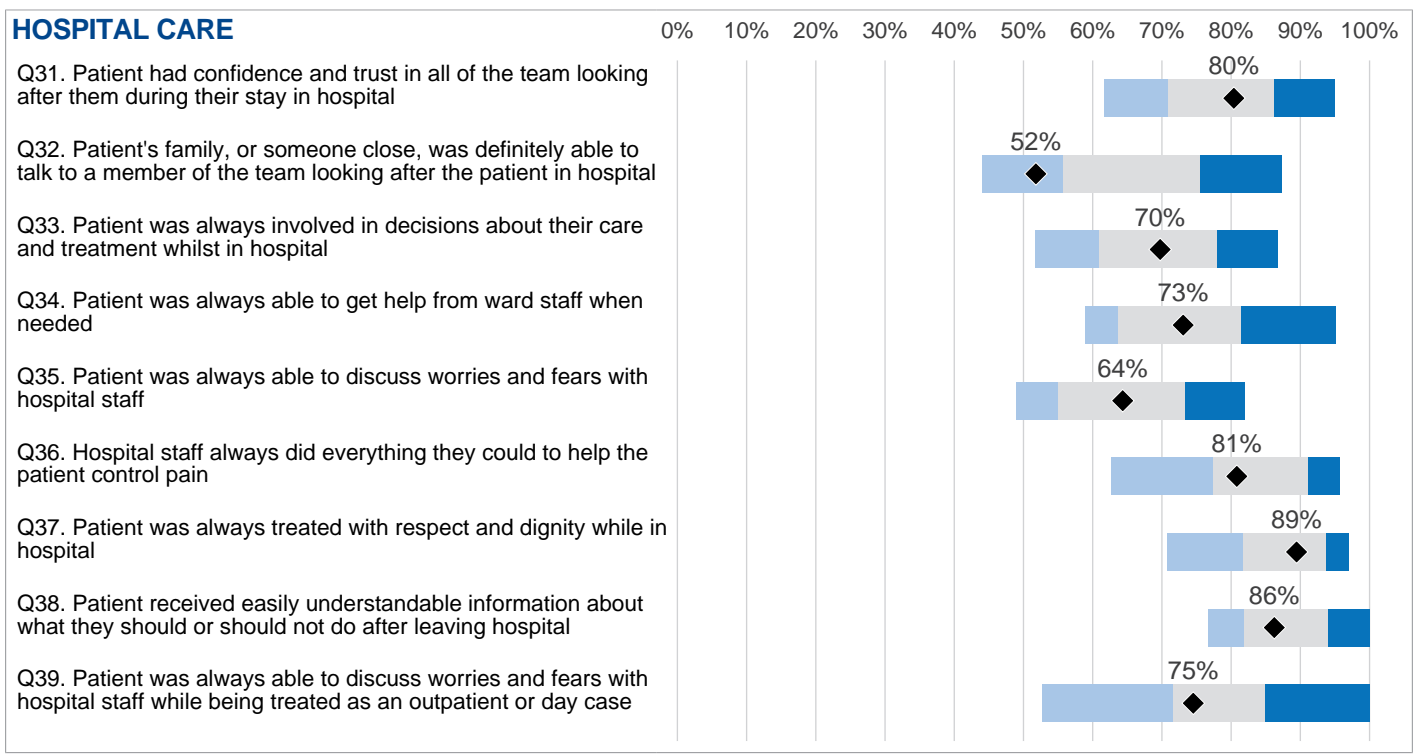
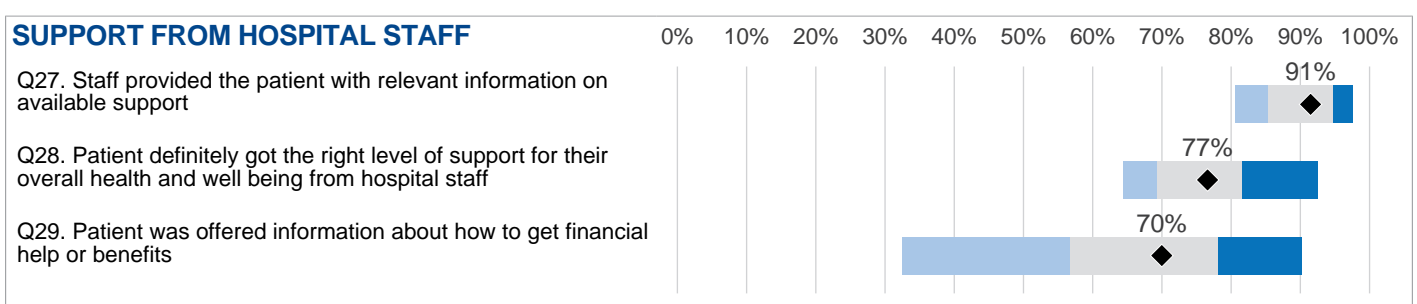
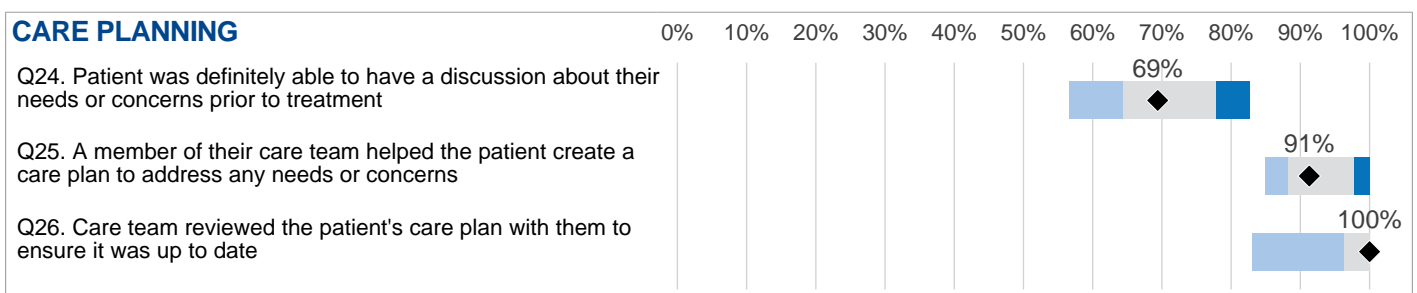
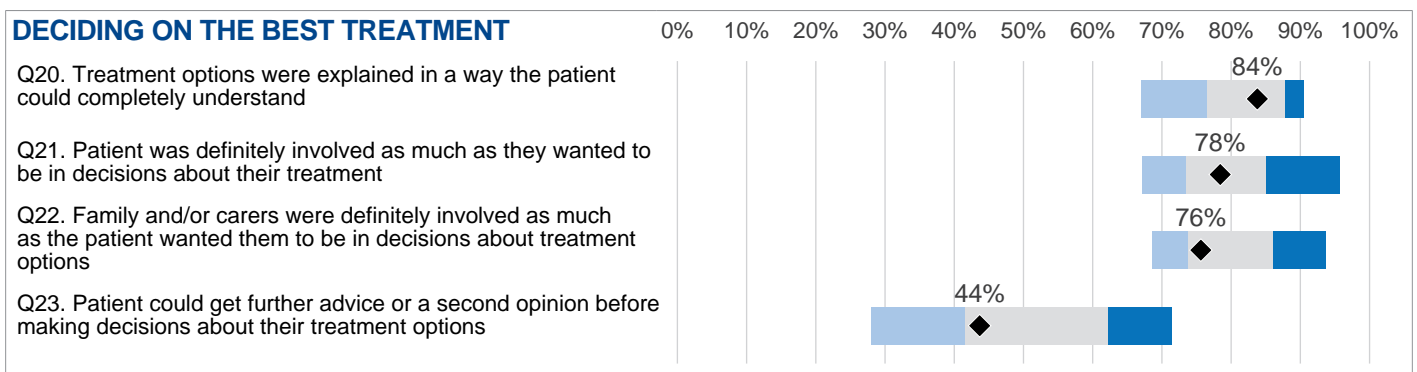
	Number of Respondents
<b>White</b>	
English / Welsh / Scottish / Northern Irish / British	155
Irish	*
Gypsy or Irish Traveller	*
Any other White background	*
<b>Mixed / Multiple Ethnicity</b>	
White and Black Caribbean	*
White and Black African	*
White and Asian	*
Any other Mixed / multiple ethnic background	*
<b>Asian or Asian British</b>	
Indian	7
Pakistani	6
Bangladeshi	*
Chinese	*
Any other Asian background	*
<b>Black / African / Caribbean / Black British</b>	
African	*
Caribbean	*
Any other Black / African / Caribbean background	*
<b>Other Ethnicity</b>	
Arab	*
Any other ethnic group	*
<b>Not given</b>	
Not given	11
<b>Total</b>	<b>193</b>

## Expected Range Charts

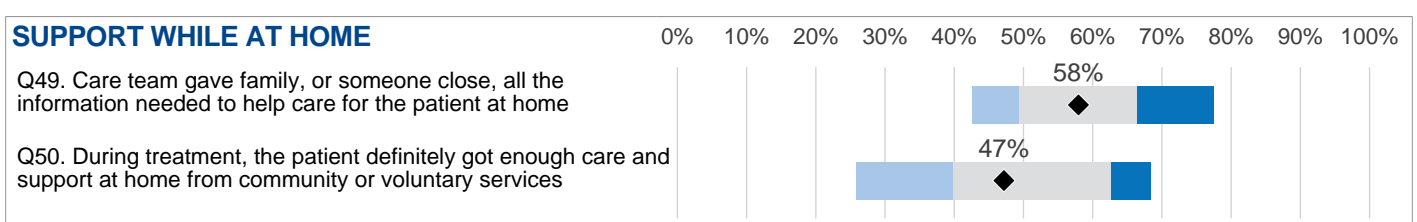
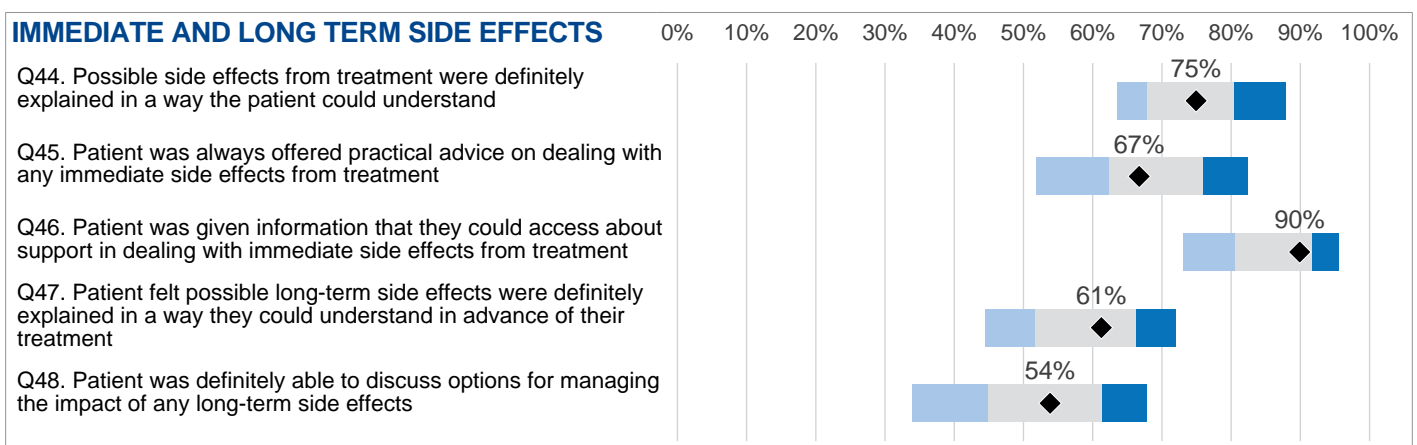
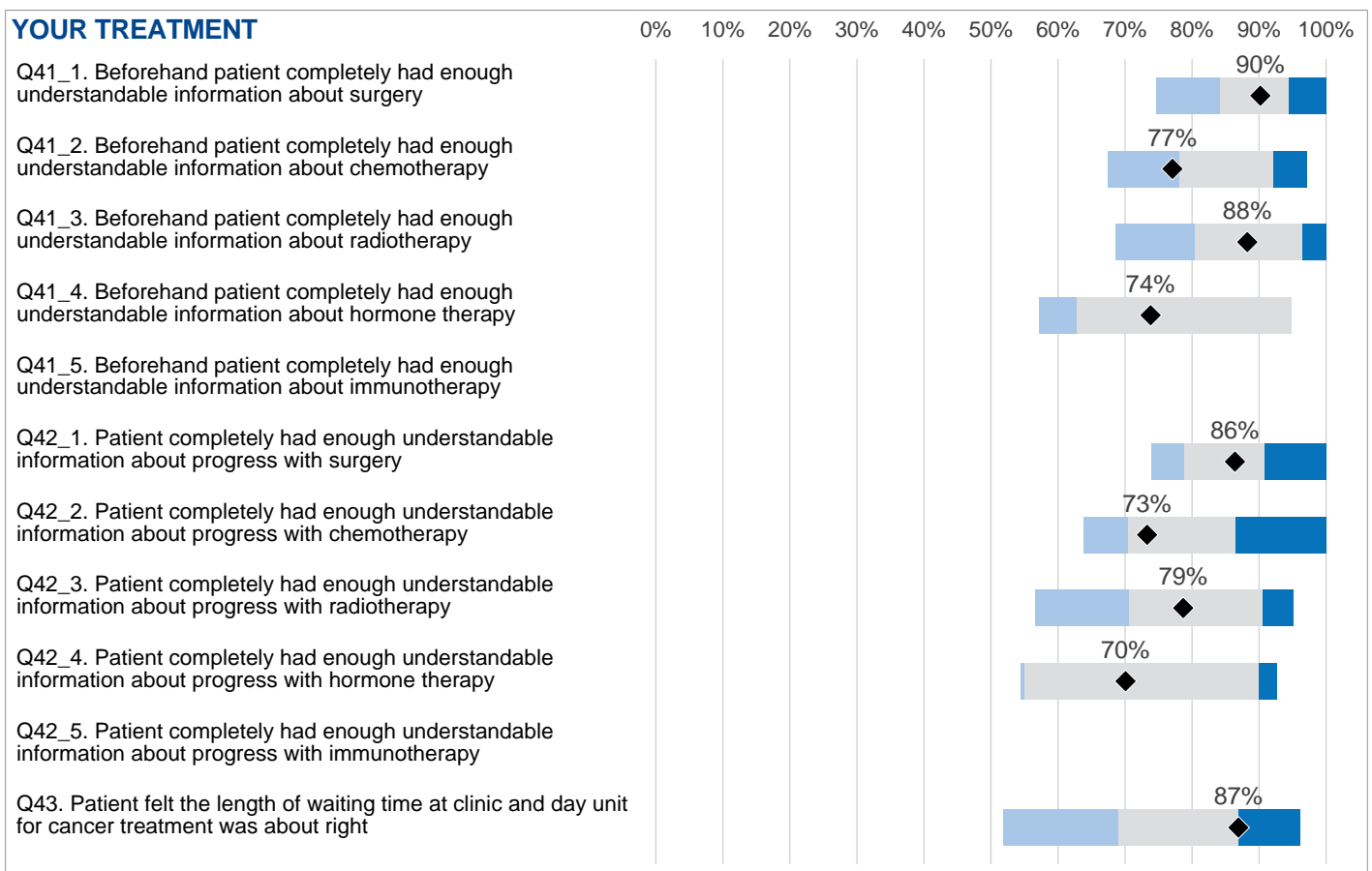




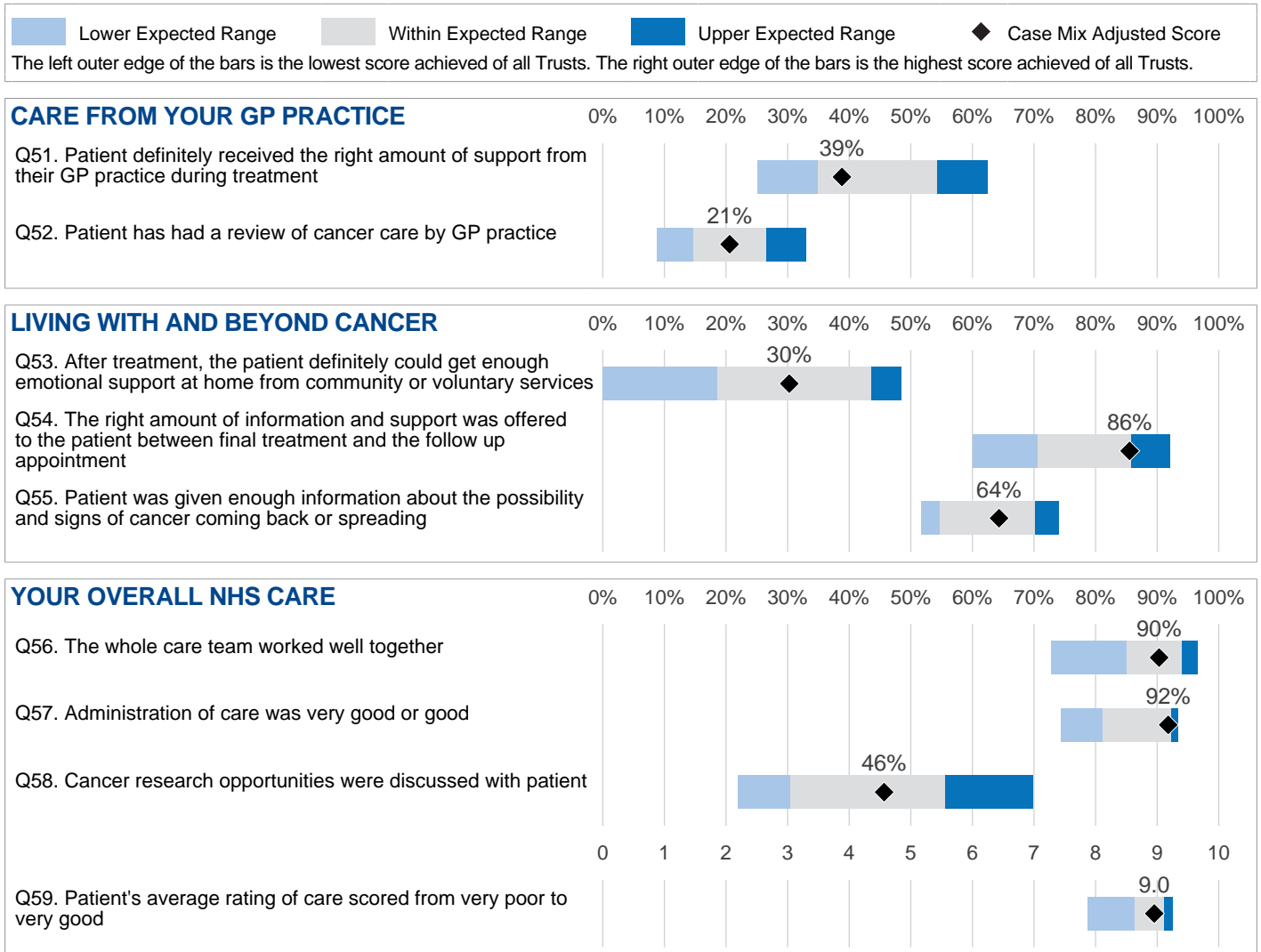
## Expected Range Charts



## Expected Range Charts



## Expected Range Charts



## Comparability tables

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

▲ or ▼

Change 2021-2022: Indicates where 2022 score is significantly higher or lower than 2021 score.

\*\* No score available for 2021.

	Adjusted Score below Lower Expected Range
	Adjusted Score between Upper and Lower Expected Ranges
	Adjusted Score above Upper Expected Range

<b>SUPPORT FROM YOUR GP PRACTICE</b>	Unadjusted Scores					Case Mix Adjusted Scores			National Score
	2021 n	2021 Score	2022 n	2022 Score	Change 2021-2022	2022 Score	Lower Expected Range	Upper Expected Range	
Q2. Patient only spoke to primary care professional once or twice before cancer diagnosis	112	<b>79%</b>	110	<b>79%</b>		<b>80%</b>	70%	85%	<b>78%</b>
Q3. Referral for diagnosis was explained in a way the patient could completely understand	155	<b>66%</b>	131	<b>69%</b>		<b>70%</b>	57%	74%	<b>65%</b>

<b>DIAGNOSTIC TESTS</b>	Unadjusted Scores					Case Mix Adjusted Scores			National Score
	2021 n	2021 Score	2022 n	2022 Score	Change 2021-2022	2022 Score	Lower Expected Range	Upper Expected Range	
Q5. Patient received all the information needed about the diagnostic test in advance	205	<b>91%</b>	156	<b>88%</b>		<b>88%</b>	88%	96%	<b>92%</b>
Q6. Diagnostic test staff appeared to completely have all the information they needed about the patient	220	<b>83%</b>	161	<b>85%</b>		<b>85%</b>	78%	89%	<b>83%</b>
Q7. Patient felt the length of time waiting for diagnostic test results was about right	218	<b>84%</b>	162	<b>85%</b>		<b>85%</b>	72%	85%	<b>78%</b>
Q8. Diagnostic test results were explained in a way the patient could completely understand	220	<b>74%</b>	162	<b>82%</b>		<b>82%</b>	72%	85%	<b>78%</b>
Q9. Enough privacy was always given to the patient when receiving diagnostic test results	218	<b>92%</b>	164	<b>95%</b>		<b>95%</b>	91%	98%	<b>95%</b>

<b>FINDING OUT THAT YOU HAD CANCER</b>	Unadjusted Scores					Case Mix Adjusted Scores			National Score
	2021 n	2021 Score	2022 n	2022 Score	Change 2021-2022	2022 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	217	<b>73%</b>	180	<b>77%</b>		<b>75%</b>	69%	83%	<b>76%</b>
Q13. Patient was definitely told sensitively that they had cancer	236	<b>72%</b>	193	<b>73%</b>		<b>72%</b>	67%	80%	<b>74%</b>
Q14. Cancer diagnosis explained in a way the patient could completely understand	236	<b>72%</b>	192	<b>77%</b>		<b>77%</b>	70%	82%	<b>76%</b>
Q15. Patient was definitely told about their diagnosis in an appropriate place	236	<b>83%</b>	191	<b>88%</b>		<b>88%</b>	80%	90%	<b>85%</b>
Q16. Patient was told they could go back later for more information about their diagnosis	208	<b>84%</b>	164	<b>87%</b>		<b>86%</b>	78%	89%	<b>84%</b>

<b>SUPPORT FROM A MAIN CONTACT PERSON</b>	Unadjusted Scores					Case Mix Adjusted Scores			National Score
	2021 n	2021 Score	2022 n	2022 Score	Change 2021-2022	2022 Score	Lower Expected Range	Upper Expected Range	
Q17. Patient had a main point of contact within the care team	230	<b>92%</b>	183	<b>93%</b>		<b>93%</b>	87%	96%	<b>91%</b>
Q18. Patient found it very or quite easy to contact their main contact person	194	<b>87%</b>	154	<b>86%</b>		<b>87%</b>	77%	90%	<b>84%</b>
Q19. Patient found advice from main contact person was very or quite helpful	202	<b>97%</b>	165	<b>97%</b>		<b>97%</b>	92%	99%	<b>95%</b>

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\*\* No score available for 2021.

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	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			National Score
	2021 n	2021 Score	2022 n	2022 Score	Change 2021-2022	2022 Score	Lower Expected Range	Upper Expected Range	
Q20. Treatment options were explained in a way the patient could completely understand	227	80%	181	84%		84%	77%	88%	82%
Q21. Patient was definitely involved as much as they wanted to be in decisions about their treatment	237	74%	188	79%		78%	74%	85%	79%
Q22. Family and/or carers were definitely involved as much as the patient wanted them to be in decisions about treatment options	200	77%	170	75%		76%	74%	86%	80%
Q23. Patient could get further advice or a second opinion before making decisions about their treatment options	124	52%	92	47%		44%	42%	62%	52%

CARE PLANNING	Unadjusted Scores					Case Mix Adjusted Scores			National Score
	2021 n	2021 Score	2022 n	2022 Score	Change 2021-2022	2022 Score	Lower Expected Range	Upper Expected Range	
Q24. Patient was definitely able to have a discussion about their needs or concerns prior to treatment	212	68%	175	70%		69%	64%	78%	71%
Q25. A member of their care team helped the patient create a care plan to address any needs or concerns	136	90%	108	92%		91%	88%	98%	93%
Q26. Care team reviewed the patient's care plan with them to ensure it was up to date	107	99%	88	100%		100%	96%	100%	99%

SUPPORT FROM HOSPITAL STAFF	Unadjusted Scores					Case Mix Adjusted Scores			National Score
	2021 n	2021 Score	2022 n	2022 Score	Change 2021-2022	2022 Score	Lower Expected Range	Upper Expected Range	
Q27. Staff provided the patient with relevant information on available support	204	89%	173	92%		91%	85%	95%	90%
Q28. Patient definitely got the right level of support for their overall health and well being from hospital staff	234	69%	190	77%		77%	69%	82%	76%
Q29. Patient was offered information about how to get financial help or benefits	142	69%	116	70%		70%	57%	78%	67%

HOSPITAL CARE	Unadjusted Scores					Case Mix Adjusted Scores			National Score
	2021 n	2021 Score	2022 n	2022 Score	Change 2021-2022	2022 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	120	73%	114	82%		80%	71%	86%	79%
Q32. Patient's family, or someone close, was definitely able to talk to a member of the team looking after the patient in hospital	97	61%	88	53%		52%	56%	76%	66%
Q33. Patient was always involved in decisions about their care and treatment whilst in hospital	118	61%	111	71%		70%	61%	78%	70%
Q34. Patient was always able to get help from ward staff when needed	120	73%	111	75%		73%	64%	81%	73%
Q35. Patient was always able to discuss worries and fears with hospital staff	116	64%	106	66%		64%	55%	73%	64%
Q36. Hospital staff always did everything they could to help the patient control pain	105	82%	107	81%		81%	77%	91%	84%
Q37. Patient was always treated with respect and dignity while in hospital	120	88%	113	90%		89%	82%	94%	88%
Q38. Patient received easily understandable information about what they should or should not do after leaving hospital	116	84%	110	86%		86%	82%	94%	88%
Q39. Patient was always able to discuss worries and fears with hospital staff while being treated as an outpatient or day case	210	72%	154	75%		75%	72%	85%	78%

## Comparability tables

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\*\* No score available for 2021.

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	Adjusted Score below Lower Expected Range
	Adjusted Score between Upper and Lower Expected Ranges
	Adjusted Score above Upper Expected Range

YOUR TREATMENT	Unadjusted Scores					Case Mix Adjusted Scores			National Score
	2021 n	2021 Score	2022 n	2022 Score	Change 2021-2022	2022 Score	Lower Expected Range	Upper Expected Range	
Q41_1. Beforehand patient completely had enough understandable information about surgery	152	<b>86%</b>	137	<b>91%</b>		<b>90%</b>	84%	95%	<b>89%</b>
Q41_2. Beforehand patient completely had enough understandable information about chemotherapy	112	<b>83%</b>	99	<b>79%</b>		<b>77%</b>	78%	92%	<b>85%</b>
Q41_3. Beforehand patient completely had enough understandable information about radiotherapy	73	<b>96%</b>	61	<b>89%</b>		<b>88%</b>	80%	96%	<b>88%</b>
Q41_4. Beforehand patient completely had enough understandable information about hormone therapy	47	<b>81%</b>	25	<b>76%</b>		<b>74%</b>	63%	95%	<b>79%</b>
Q41_5. Beforehand patient completely had enough understandable information about immunotherapy	31	<b>71%</b>		*		*			<b>84%</b>
Q42_1. Patient completely had enough understandable information about progress with surgery	152	<b>82%</b>	137	<b>87%</b>		<b>86%</b>	79%	91%	<b>85%</b>
Q42_2. Patient completely had enough understandable information about progress with chemotherapy	112	<b>68%</b>	100	<b>75%</b>		<b>73%</b>	70%	87%	<b>79%</b>
Q42_3. Patient completely had enough understandable information about progress with radiotherapy	73	<b>81%</b>	60	<b>80%</b>		<b>79%</b>	71%	91%	<b>81%</b>
Q42_4. Patient completely had enough understandable information about progress with hormone therapy	46	<b>72%</b>	25	<b>72%</b>		<b>70%</b>	55%	90%	<b>72%</b>
Q42_5. Patient completely had enough understandable information about progress with immunotherapy	29	<b>66%</b>		*		*			<b>80%</b>
Q43. Patient felt the length of waiting time at clinic and day unit for cancer treatment was about right	230	<b>79%</b>	187	<b>87%</b>		<b>87%</b>	69%	87%	<b>78%</b>

IMMEDIATE AND LONG TERM SIDE EFFECTS	Unadjusted Scores					Case Mix Adjusted Scores			National Score
	2021 n	2021 Score	2022 n	2022 Score	Change 2021-2022	2022 Score	Lower Expected Range	Upper Expected Range	
Q44. Possible side effects from treatment were definitely explained in a way the patient could understand	223	<b>70%</b>	184	<b>76%</b>		<b>75%</b>	68%	81%	<b>74%</b>
Q45. Patient was always offered practical advice on dealing with any immediate side effects from treatment	215	<b>67%</b>	177	<b>68%</b>		<b>67%</b>	62%	76%	<b>69%</b>
Q46. Patient was given information that they could access about support in dealing with immediate side effects from treatment	180	<b>86%</b>	149	<b>90%</b>		<b>90%</b>	81%	92%	<b>86%</b>
Q47. Patient felt possible long-term side effects were definitely explained in a way they could understand in advance of their treatment	222	<b>62%</b>	172	<b>64%</b>		<b>61%</b>	52%	66%	<b>59%</b>
Q48. Patient was definitely able to discuss options for managing the impact of any long-term side effects	194	<b>51%</b>	159	<b>57%</b>		<b>54%</b>	45%	61%	<b>53%</b>

SUPPORT WHILE AT HOME	Unadjusted Scores					Case Mix Adjusted Scores			National Score
	2021 n	2021 Score	2022 n	2022 Score	Change 2021-2022	2022 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	162	<b>52%</b>	142	<b>60%</b>		<b>58%</b>	49%	66%	<b>58%</b>
Q50. During treatment, the patient definitely got enough care and support at home from community or voluntary services	99	<b>40%</b>	76	<b>47%</b>		<b>47%</b>	40%	63%	<b>51%</b>

## Comparability tables

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

▲ or ▼

Change 2021-2022: Indicates where 2022 score is significantly higher or lower than 2021 score.

\*\* No score available for 2021.

	Adjusted Score below Lower Expected Range
	Adjusted Score between Upper and Lower Expected Ranges
	Adjusted Score above Upper Expected Range

<b>CARE FROM YOUR GP PRACTICE</b>	Unadjusted Scores					Case Mix Adjusted Scores			National Score
	2021 n	2021 Score	2022 n	2022 Score	Change 2021-2022	2022 Score	Lower Expected Range	Upper Expected Range	
Q51. Patient definitely received the right amount of support from their GP practice during treatment	141	<b>46%</b>	121	<b>40%</b>		<b>39%</b>	35%	54%	<b>45%</b>
Q52. Patient has had a review of cancer care by GP practice	223	<b>25%</b>	184	<b>22%</b>		<b>21%</b>	15%	27%	<b>21%</b>

<b>LIVING WITH AND BEYOND CANCER</b>	Unadjusted Scores					Case Mix Adjusted Scores			National Score
	2021 n	2021 Score	2022 n	2022 Score	Change 2021-2022	2022 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	58	<b>21%</b>	53	<b>32%</b>		<b>30%</b>	19%	44%	<b>31%</b>
Q54. The right amount of information and support was offered to the patient between final treatment and the follow up appointment	114	<b>77%</b>	112	<b>86%</b>		<b>86%</b>	71%	86%	<b>78%</b>
Q55. Patient was given enough information about the possibility and signs of cancer coming back or spreading	205	<b>63%</b>	154	<b>65%</b>		<b>64%</b>	55%	70%	<b>62%</b>

<b>YOUR OVERALL NHS CARE</b>	Unadjusted Scores					Case Mix Adjusted Scores			National Score
	2021 n	2021 Score	2022 n	2022 Score	Change 2021-2022	2022 Score	Lower Expected Range	Upper Expected Range	
Q56. The whole care team worked well together	227	<b>88%</b>	183	<b>91%</b>		<b>90%</b>	85%	94%	<b>90%</b>
Q57. Administration of care was very good or good	232	<b>84%</b>	190	<b>92%</b>		<b>92%</b>	81%	92%	<b>87%</b>
Q58. Cancer research opportunities were discussed with patient	131	<b>37%</b>	111	<b>47%</b>		<b>46%</b>	30%	56%	<b>43%</b>
Q59. Patient's average rating of care scored from very poor to very good	227	<b>8.6</b>	188	<b>8.9</b>		<b>9.0</b>	8.6	9.1	<b>8.9</b>



## Tumour type tables

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

	Tumour Type													
	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	*	96%	53%	*	31%	*	*	94%	*	*	*	*	92%	<b>79%</b>
Q3. Referral for diagnosis was explained in a way the patient could completely understand	*	96%	38%	60%	67%	*	*	70%	*	*	*	*	69%	<b>69%</b>

	Tumour Type													
	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	*	95%	82%	90%	73%	*	*	90%	*	*	73%	90%	93%	<b>88%</b>
Q6. Diagnostic test staff appeared to completely have all the information they needed about the patient	*	92%	87%	80%	71%	*	*	100%	*	*	64%	90%	73%	<b>85%</b>
Q7. Patient felt the length of time waiting for diagnostic test results was about right	*	89%	91%	*	71%	*	*	100%	*	*	100%	70%	81%	<b>85%</b>
Q8. Diagnostic test results were explained in a way the patient could completely understand	*	97%	74%	60%	63%	*	*	75%	*	*	100%	100%	80%	<b>82%</b>
Q9. Enough privacy was always given to the patient when receiving diagnostic test results	*	97%	87%	90%	94%	*	*	100%	*	*	82%	100%	94%	<b>95%</b>

	Tumour Type													
	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	*	90%	64%	80%	71%	*	*	87%	*	*	73%	73%	68%	<b>77%</b>
Q13. Patient was definitely told sensitively that they had cancer	*	95%	54%	80%	56%	*	*	69%	*	*	64%	75%	65%	<b>73%</b>
Q14. Cancer diagnosis explained in a way the patient could completely understand	*	91%	77%	60%	58%	*	*	69%	*	*	82%	92%	65%	<b>77%</b>
Q15. Patient was definitely told about their diagnosis in an appropriate place	*	91%	88%	*	75%	*	*	96%	*	*	73%	100%	80%	<b>88%</b>
Q16. Patient was told they could go back later for more information about their diagnosis	*	97%	76%	*	94%	*	*	77%	*	*	82%	91%	69%	<b>87%</b>



## Tumour type tables

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

	Tumour Type													All Cancers
	Brain / CNS	Breast	Colorectal / LGT	Gynaecological	Haematological	Head and Neck	Lung	Prostate	Sarcoma	Skin	Upper Gastro	Urological	Other	
Q17. Patient had a main point of contact within the care team	*	100%	96%	*	96%	*	*	92%	*	*	90%	90%	74%	<b>93%</b>
Q18. Patient found it very or quite easy to contact their main contact person	*	95%	85%	*	79%	*	*	95%	*	*	*	*	85%	<b>86%</b>
Q19. Patient found advice from main contact person was very or quite helpful	*	98%	96%	*	96%	*	*	100%	*	*	*	*	93%	<b>97%</b>

	Tumour Type													All Cancers
	Brain / CNS	Breast	Colorectal / LGT	Gynaecological	Haematological	Head and Neck	Lung	Prostate	Sarcoma	Skin	Upper Gastro	Urological	Other	
Q20. Treatment options were explained in a way the patient could completely understand	*	93%	83%	*	64%	*	*	85%	*	*	73%	100%	84%	<b>84%</b>
Q21. Patient was definitely involved as much as they wanted to be in decisions about their treatment	*	98%	80%	80%	52%	*	*	85%	*	*	73%	67%	68%	<b>79%</b>
Q22. Family and/or carers were definitely involved as much as the patient wanted them to be in decisions about treatment options	*	87%	68%	*	54%	*	*	83%	*	*	70%	*	63%	<b>75%</b>
Q23. Patient could get further advice or a second opinion before making decisions about their treatment options	*	63%	47%	*	36%	*	*	62%	*	*	*	*	*	<b>47%</b>

	Tumour Type													All Cancers
	Brain / CNS	Breast	Colorectal / LGT	Gynaecological	Haematological	Head and Neck	Lung	Prostate	Sarcoma	Skin	Upper Gastro	Urological	Other	
Q24. Patient was definitely able to have a discussion about their needs or concerns prior to treatment	*	85%	60%	*	52%	*	*	63%	*	*	80%	90%	69%	<b>70%</b>
Q25. A member of their care team helped the patient create a care plan to address any needs or concerns	*	100%	94%	*	93%	*	*	80%	*	*	*	*	*	<b>92%</b>
Q26. Care team reviewed the patient's care plan with them to ensure it was up to date	*	100%	100%	*	*	*	*	100%	*	*	*	*	*	<b>100%</b>

	Tumour Type													All Cancers
	Brain / CNS	Breast	Colorectal / LGT	Gynaecological	Haematological	Head and Neck	Lung	Prostate	Sarcoma	Skin	Upper Gastro	Urological	Other	
Q27. Staff provided the patient with relevant information on available support	*	98%	88%	100%	85%	*	*	88%	*	*	100%	82%	94%	<b>92%</b>
Q28. Patient definitely got the right level of support for their overall health and well being from hospital staff	*	88%	72%	90%	64%	*	*	69%	*	*	73%	67%	79%	<b>77%</b>
Q29. Patient was offered information about how to get financial help or benefits	*	86%	41%	*	84%	*	*	69%	*	*	*	*	*	<b>70%</b>

## Tumour type tables

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

	Tumour Type													
	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	*	88%	72%	*	77%	*	*	78%	*	*	82%	91%	*	<b>82%</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%	38%	*	50%	*	*	50%	*	*	36%	*	*	<b>53%</b>
Q33. Patient was always involved in decisions about their care and treatment whilst in hospital	*	88%	59%	*	46%	*	*	83%	*	*	64%	90%	*	<b>71%</b>
Q34. Patient was always able to get help from ward staff when needed	*	82%	47%	*	83%	*	*	78%	*	*	73%	90%	*	<b>75%</b>
Q35. Patient was always able to discuss worries and fears with hospital staff	*	69%	44%	*	83%	*	*	71%	*	*	60%	*	*	<b>66%</b>
Q36. Hospital staff always did everything they could to help the patient control pain	*	94%	76%	*	*	*	*	78%	*	*	90%	80%	*	<b>81%</b>
Q37. Patient was always treated with respect and dignity while in hospital	*	76%	89%	*	92%	*	*	89%	*	*	91%	90%	*	<b>90%</b>
Q38. Patient received easily understandable information about what they should or should not do after leaving hospital	*	87%	89%	*	83%	*	*	83%	*	*	70%	91%	*	<b>86%</b>
Q39. Patient was always able to discuss worries and fears with hospital staff while being treated as an outpatient or day case	*	80%	61%	*	74%	*	*	76%	*	*	*	*	59%	<b>75%</b>

	Tumour Type													
	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	*	97%	86%	*	*	*	*	82%	*	*	*	100%	80%	<b>91%</b>
Q41_2. Beforehand patient completely had enough understandable information about chemotherapy	*	90%	65%	*	73%	*	*	*	*	*	90%	*	*	<b>79%</b>
Q41_3. Beforehand patient completely had enough understandable information about radiotherapy	*	97%	*	*	*	*	*	*	*	*	*	*	*	<b>89%</b>
Q41_4. Beforehand patient completely had enough understandable information about hormone therapy	*	82%	*	*	*	*	*	*	*	*	*	*	*	<b>76%</b>
Q41_5. Beforehand patient completely had enough understandable information about immunotherapy	*	*	*	*	*	*	*	*	*	*	*	*	*	*
Q42_1. Patient completely had enough understandable information about progress with surgery	*	92%	82%	*	*	*	*	82%	*	*	*	100%	73%	<b>87%</b>
Q42_2. Patient completely had enough understandable information about progress with chemotherapy	*	77%	59%	*	77%	*	*	*	*	*	80%	*	*	<b>75%</b>
Q42_3. Patient completely had enough understandable information about progress with radiotherapy	*	87%	*	*	*	*	*	*	*	*	*	*	*	<b>80%</b>
Q42_4. Patient completely had enough understandable information about progress with hormone therapy	*	82%	*	*	*	*	*	*	*	*	*	*	*	<b>72%</b>
Q42_5. Patient completely had enough understandable information about progress with immunotherapy	*	*	*	*	*	*	*	*	*	*	*	*	*	*
Q43. Patient felt the length of waiting time at clinic and day unit for cancer treatment was about right	*	88%	92%	*	74%	*	*	92%	*	*	73%	92%	85%	<b>87%</b>

## Tumour type tables

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

	Tumour Type													
	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	*	91%	71%	*	68%	*	*	77%	*	*	73%	92%	56%	<b>76%</b>
Q45. Patient was always offered practical advice on dealing with any immediate side effects from treatment	*	93%	55%	*	55%	*	*	56%	*	*	64%	73%	53%	<b>68%</b>
Q46. Patient was given information that they could access about support in dealing with immediate side effects from treatment	*	100%	80%	*	90%	*	*	88%	*	*	80%	91%	86%	<b>90%</b>
Q47. Patient felt possible long-term side effects were definitely explained in a way they could understand in advance of their treatment	*	88%	50%	*	45%	*	*	74%	*	*	55%	58%	41%	<b>64%</b>
Q48. Patient was definitely able to discuss options for managing the impact of any long-term side effects	*	76%	55%	*	42%	*	*	55%	*	*	50%	64%	33%	<b>57%</b>

	Tumour Type													
	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	*	67%	45%	*	63%	*	*	63%	*	*	50%	64%	54%	<b>60%</b>
Q50. During treatment, the patient definitely got enough care and support at home from community or voluntary services	*	54%	46%	*	50%	*	*	*	*	*	*	*	*	<b>47%</b>

	Tumour Type													
	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	*	50%	33%	*	18%	*	*	35%	*	*	*	*	50%	<b>40%</b>
Q52. Patient has had a review of cancer care by GP practice	*	20%	12%	40%	8%	*	*	33%	*	*	18%	27%	16%	<b>22%</b>

## Tumour type tables

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

	Tumour Type													
	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	*	*	27%	*	*	*	*	*	*	*	*	*	*	<b>32%</b>
Q54. The right amount of information and support was offered to the patient between final treatment and the follow up appointment	*	96%	72%	*	*	*	*	83%	*	*	*	*	*	<b>86%</b>
Q55. Patient was given enough information about the possibility and signs of cancer coming back or spreading	*	70%	52%	*	58%	*	*	68%	*	*	*	*	65%	<b>65%</b>

	Tumour Type													
	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	*	95%	88%	*	91%	*	*	91%	*	*	73%	92%	100%	<b>91%</b>
Q57. Administration of care was very good or good	*	95%	85%	90%	88%	*	*	88%	*	*	82%	100%	100%	<b>92%</b>
Q58. Cancer research opportunities were discussed with patient	*	50%	47%	*	67%	*	*	53%	*	*	*	*	45%	<b>47%</b>
Q59. Patient's average rating of care scored from very poor to very good	*	9.3	8.6	9.1	8.7	*	*	9.1	*	*	8.0	8.6	9.2	<b>8.9</b>

## Age group tables

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

<b>SUPPORT FROM YOUR GP PRACTICE</b>	Age								
	16 - 24	25 - 34	35 - 44	45 - 54	55 - 64	65 - 74	75 - 84	85+	All
Q2. Patient only spoke to primary care professional once or twice before cancer diagnosis	*	*	*	92%	75%	80%	77%	*	<b>79%</b>
Q3. Referral for diagnosis was explained in a way the patient could completely understand	*	*	*	80%	82%	65%	61%	*	<b>69%</b>

<b>DIAGNOSTIC TESTS</b>	Age								
	16 - 24	25 - 34	35 - 44	45 - 54	55 - 64	65 - 74	75 - 84	85+	All
Q5. Patient received all the information needed about the diagnostic test in advance	*	*	*	76%	94%	86%	88%	*	<b>88%</b>
Q6. Diagnostic test staff appeared to completely have all the information they needed about the patient	*	*	*	76%	83%	89%	86%	*	<b>85%</b>
Q7. Patient felt the length of time waiting for diagnostic test results was about right	*	*	*	71%	84%	89%	89%	*	<b>85%</b>
Q8. Diagnostic test results were explained in a way the patient could completely understand	*	*	*	76%	83%	81%	86%	*	<b>82%</b>
Q9. Enough privacy was always given to the patient when receiving diagnostic test results	*	*	*	88%	94%	96%	97%	*	<b>95%</b>

<b>FINDING OUT THAT YOU HAD CANCER</b>	Age								
	16 - 24	25 - 34	35 - 44	45 - 54	55 - 64	65 - 74	75 - 84	85+	All
Q12. Patient was told they could have a family member, carer or friend with them when told diagnosis	*	*	*	72%	78%	77%	75%	*	<b>77%</b>
Q13. Patient was definitely told sensitively that they had cancer	*	*	*	79%	77%	72%	64%	*	<b>73%</b>
Q14. Cancer diagnosis explained in a way the patient could completely understand	*	*	*	68%	77%	77%	83%	*	<b>77%</b>
Q15. Patient was definitely told about their diagnosis in an appropriate place	*	*	*	79%	89%	89%	88%	*	<b>88%</b>
Q16. Patient was told they could go back later for more information about their diagnosis	*	*	*	82%	90%	87%	84%	*	<b>87%</b>

<b>SUPPORT FROM A MAIN CONTACT PERSON</b>	Age								
	16 - 24	25 - 34	35 - 44	45 - 54	55 - 64	65 - 74	75 - 84	85+	All
Q17. Patient had a main point of contact within the care team	*	*	*	95%	96%	94%	90%	*	<b>93%</b>
Q18. Patient found it very or quite easy to contact their main contact person	*	*	*	71%	93%	89%	79%	*	<b>86%</b>
Q19. Patient found advice from main contact person was very or quite helpful	*	*	*	100%	98%	98%	94%	*	<b>97%</b>

<b>DECIDING ON THE BEST TREATMENT</b>	Age								
	16 - 24	25 - 34	35 - 44	45 - 54	55 - 64	65 - 74	75 - 84	85+	All
Q20. Treatment options were explained in a way the patient could completely understand	*	*	*	76%	83%	82%	89%	*	<b>84%</b>
Q21. Patient was definitely involved as much as they wanted to be in decisions about their treatment	*	*	*	63%	83%	75%	88%	*	<b>79%</b>
Q22. Family and/or carers were definitely involved as much as the patient wanted them to be in decisions about treatment options	*	*	*	72%	79%	73%	74%	*	<b>75%</b>
Q23. Patient could get further advice or a second opinion before making decisions about their treatment options	*	*	*	*	44%	50%	50%	*	<b>47%</b>

## Age group tables

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

<b>CARE PLANNING</b>	Age								<b>All</b>
	16 - 24	25 - 34	35 - 44	45 - 54	55 - 64	65 - 74	75 - 84	85+	
Q24. Patient was definitely able to have a discussion about their needs or concerns prior to treatment	*	*	*	68%	72%	64%	77%	*	<b>70%</b>
Q25. A member of their care team helped the patient create a care plan to address any needs or concerns	*	*	*	85%	100%	85%	96%	*	<b>92%</b>
Q26. Care team reviewed the patient's care plan with them to ensure it was up to date	*	*	*	100%	100%	100%	100%	*	<b>100%</b>

<b>SUPPORT FROM HOSPITAL STAFF</b>	Age								<b>All</b>
	16 - 24	25 - 34	35 - 44	45 - 54	55 - 64	65 - 74	75 - 84	85+	
Q27. Staff provided the patient with relevant information on available support	*	*	*	89%	96%	87%	94%	*	<b>92%</b>
Q28. Patient definitely got the right level of support for their overall health and well being from hospital staff	*	*	*	63%	81%	75%	80%	*	<b>77%</b>
Q29. Patient was offered information about how to get financial help or benefits	*	*	*	81%	79%	63%	50%	*	<b>70%</b>

<b>HOSPITAL CARE</b>	Age								<b>All</b>
	16 - 24	25 - 34	35 - 44	45 - 54	55 - 64	65 - 74	75 - 84	85+	
Q31. Patient had confidence and trust in all of the team looking after them during their stay in hospital	*	*	*	67%	84%	83%	83%	*	<b>82%</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	*	*	*	50%	56%	47%	67%	*	<b>53%</b>
Q33. Patient was always involved in decisions about their care and treatment whilst in hospital	*	*	*	75%	73%	66%	74%	*	<b>71%</b>
Q34. Patient was always able to get help from ward staff when needed	*	*	*	83%	68%	75%	78%	*	<b>75%</b>
Q35. Patient was always able to discuss worries and fears with hospital staff	*	*	*	50%	69%	64%	71%	*	<b>66%</b>
Q36. Hospital staff always did everything they could to help the patient control pain	*	*	*	83%	77%	80%	90%	*	<b>81%</b>
Q37. Patient was always treated with respect and dignity while in hospital	*	*	*	83%	91%	85%	100%	*	<b>90%</b>
Q38. Patient received easily understandable information about what they should or should not do after leaving hospital	*	*	*	83%	84%	90%	82%	*	<b>86%</b>
Q39. Patient was always able to discuss worries and fears with hospital staff while being treated as an outpatient or day case	*	*	*	76%	77%	68%	78%	*	<b>75%</b>

## Age group tables

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

YOUR TREATMENT	Age								All
	16 - 24	25 - 34	35 - 44	45 - 54	55 - 64	65 - 74	75 - 84	85+	
Q41_1. Beforehand patient completely had enough understandable information about surgery	*	*	*	94%	87%	91%	93%	*	<b>91%</b>
Q41_2. Beforehand patient completely had enough understandable information about chemotherapy	*	*	*	82%	74%	90%	65%	*	<b>79%</b>
Q41_3. Beforehand patient completely had enough understandable information about radiotherapy	*	*	*	90%	85%	89%	90%	*	<b>89%</b>
Q41_4. Beforehand patient completely had enough understandable information about hormone therapy	*	*	*	*	*	*	*	*	<b>76%</b>
Q41_5. Beforehand patient completely had enough understandable information about immunotherapy	*	*	*	*	*	*	*	*	*
Q42_1. Patient completely had enough understandable information about progress with surgery	*	*	*	88%	87%	87%	86%	*	<b>87%</b>
Q42_2. Patient completely had enough understandable information about progress with chemotherapy	*	*	*	67%	79%	72%	71%	*	<b>75%</b>
Q42_3. Patient completely had enough understandable information about progress with radiotherapy	*	*	*	*	85%	83%	80%	*	<b>80%</b>
Q42_4. Patient completely had enough understandable information about progress with hormone therapy	*	*	*	*	*	*	*	*	<b>72%</b>
Q42_5. Patient completely had enough understandable information about progress with immunotherapy	*	*	*	*	*	*	*	*	*
Q43. Patient felt the length of waiting time at clinic and day unit for cancer treatment was about right	*	*	*	83%	93%	83%	88%	*	<b>87%</b>

IMMEDIATE AND LONG TERM SIDE EFFECTS	Age								All
	16 - 24	25 - 34	35 - 44	45 - 54	55 - 64	65 - 74	75 - 84	85+	
Q44. Possible side effects from treatment were definitely explained in a way the patient could understand	*	*	*	84%	82%	72%	72%	*	<b>76%</b>
Q45. Patient was always offered practical advice on dealing with any immediate side effects from treatment	*	*	*	58%	68%	68%	71%	*	<b>68%</b>
Q46. Patient was given information that they could access about support in dealing with immediate side effects from treatment	*	*	*	83%	96%	90%	83%	*	<b>90%</b>
Q47. Patient felt possible long-term side effects were definitely explained in a way they could understand in advance of their treatment	*	*	*	74%	70%	58%	65%	*	<b>64%</b>
Q48. Patient was definitely able to discuss options for managing the impact of any long-term side effects	*	*	*	61%	65%	42%	67%	*	<b>57%</b>

SUPPORT WHILE AT HOME	Age								All
	16 - 24	25 - 34	35 - 44	45 - 54	55 - 64	65 - 74	75 - 84	85+	
Q49. Care team gave family, or someone close, all the information needed to help care for the patient at home	*	*	*	44%	67%	49%	75%	*	<b>60%</b>
Q50. During treatment, the patient definitely got enough care and support at home from community or voluntary services	*	*	*	*	48%	48%	53%	*	<b>47%</b>

CARE FROM YOUR GP PRACTICE	Age								All
	16 - 24	25 - 34	35 - 44	45 - 54	55 - 64	65 - 74	75 - 84	85+	
Q51. Patient definitely received the right amount of support from their GP practice during treatment	*	*	*	40%	38%	44%	33%	*	<b>40%</b>
Q52. Patient has had a review of cancer care by GP practice	*	*	*	12%	26%	23%	15%	*	<b>22%</b>



## Age group tables

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

	Age								All
	16 - 24	25 - 34	35 - 44	45 - 54	55 - 64	65 - 74	75 - 84	85+	
Q53. After treatment, the patient definitely could get enough emotional support at home from community or voluntary services	*	*	*	*	31%	30%	40%	*	<b>32%</b>
Q54. The right amount of information and support was offered to the patient between final treatment and the follow up appointment	*	*	*	55%	94%	84%	96%	*	<b>86%</b>
Q55. Patient was given enough information about the possibility and signs of cancer coming back or spreading	*	*	*	63%	66%	60%	74%	*	<b>65%</b>

	Age								All
	16 - 24	25 - 34	35 - 44	45 - 54	55 - 64	65 - 74	75 - 84	85+	
Q56. The whole care team worked well together	*	*	*	83%	91%	89%	95%	*	<b>91%</b>
Q57. Administration of care was very good or good	*	*	*	89%	93%	91%	93%	*	<b>92%</b>
Q58. Cancer research opportunities were discussed with patient	*	*	*	23%	57%	45%	43%	*	<b>47%</b>
Q59. Patient's average rating of care scored from very poor to very good	*	*	*	8.2	9.1	9.0	8.9	*	<b>8.9</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	77%	80%	*	*	*	*	<b>79%</b>
Q3. Referral for diagnosis was explained in a way the patient could completely understand	77%	64%	*	*	*	*	<b>69%</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	87%	88%	*	*	*	*	<b>88%</b>
Q6. Diagnostic test staff appeared to completely have all the information they needed about the patient	84%	86%	*	*	*	*	<b>85%</b>
Q7. Patient felt the length of time waiting for diagnostic test results was about right	80%	89%	*	*	*	*	<b>85%</b>
Q8. Diagnostic test results were explained in a way the patient could completely understand	82%	81%	*	*	*	*	<b>82%</b>
Q9. Enough privacy was always given to the patient when receiving diagnostic test results	93%	96%	*	*	*	*	<b>95%</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	74%	78%	*	*	*	*	<b>77%</b>
Q13. Patient was definitely told sensitively that they had cancer	81%	64%	*	*	*	*	<b>73%</b>
Q14. Cancer diagnosis explained in a way the patient could completely understand	75%	80%	*	*	*	*	<b>77%</b>
Q15. Patient was definitely told about their diagnosis in an appropriate place	84%	91%	*	*	*	*	<b>88%</b>
Q16. Patient was told they could go back later for more information about their diagnosis	89%	84%	*	*	*	*	<b>87%</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	97%	90%	*	*	*	*	<b>93%</b>
Q18. Patient found it very or quite easy to contact their main contact person	88%	84%	*	*	*	*	<b>86%</b>
Q19. Patient found advice from main contact person was very or quite helpful	97%	97%	*	*	*	*	<b>97%</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						
	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	84%	84%	*	*	*	*	<b>84%</b>
Q21. Patient was definitely involved as much as they wanted to be in decisions about their treatment	77%	80%	*	*	*	*	<b>79%</b>
Q22. Family and/or carers were definitely involved as much as the patient wanted them to be in decisions about treatment options	74%	76%	*	*	*	*	<b>75%</b>
Q23. Patient could get further advice or a second opinion before making decisions about their treatment options	42%	55%	*	*	*	*	<b>47%</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	70%	70%	*	*	*	*	<b>70%</b>
Q25. A member of their care team helped the patient create a care plan to address any needs or concerns	95%	89%	*	*	*	*	<b>92%</b>
Q26. Care team reviewed the patient's care plan with them to ensure it was up to date	100%	100%	*	*	*	*	<b>100%</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	93%	91%	*	*	*	*	<b>92%</b>
Q28. Patient definitely got the right level of support for their overall health and well being from hospital staff	80%	76%	*	*	*	*	<b>77%</b>
Q29. Patient was offered information about how to get financial help or benefits	75%	69%	*	*	*	*	<b>70%</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	78%	85%	*	*	*	*	<b>82%</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	58%	51%	*	*	*	*	<b>53%</b>
Q33. Patient was always involved in decisions about their care and treatment whilst in hospital	70%	72%	*	*	*	*	<b>71%</b>
Q34. Patient was always able to get help from ward staff when needed	77%	73%	*	*	*	*	<b>75%</b>
Q35. Patient was always able to discuss worries and fears with hospital staff	67%	65%	*	*	*	*	<b>66%</b>
Q36. Hospital staff always did everything they could to help the patient control pain	83%	81%	*	*	*	*	<b>81%</b>
Q37. Patient was always treated with respect and dignity while in hospital	89%	92%	*	*	*	*	<b>90%</b>
Q38. Patient received easily understandable information about what they should or should not do after leaving hospital	88%	86%	*	*	*	*	<b>86%</b>
Q39. Patient was always able to discuss worries and fears with hospital staff while being treated as an outpatient or day case	73%	76%	*	*	*	*	<b>75%</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	93%	90%	*	*	*	*	<b>91%</b>
Q41_2. Beforehand patient completely had enough understandable information about chemotherapy	78%	80%	*	*	*	*	<b>79%</b>
Q41_3. Beforehand patient completely had enough understandable information about radiotherapy	88%	94%	*	*	*	*	<b>89%</b>
Q41_4. Beforehand patient completely had enough understandable information about hormone therapy	74%	*	*	*	*	*	<b>76%</b>
Q41_5. Beforehand patient completely had enough understandable information about immunotherapy	*	*	*	*	*	*	*
Q42_1. Patient completely had enough understandable information about progress with surgery	87%	89%	*	*	*	*	<b>87%</b>
Q42_2. Patient completely had enough understandable information about progress with chemotherapy	71%	80%	*	*	*	*	<b>75%</b>
Q42_3. Patient completely had enough understandable information about progress with radiotherapy	80%	81%	*	*	*	*	<b>80%</b>
Q42_4. Patient completely had enough understandable information about progress with hormone therapy	68%	*	*	*	*	*	<b>72%</b>
Q42_5. Patient completely had enough understandable information about progress with immunotherapy	*	*	*	*	*	*	*
Q43. Patient felt the length of waiting time at clinic and day unit for cancer treatment was about right	88%	86%	*	*	*	*	<b>87%</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	74%	79%	*	*	*	*	<b>76%</b>
Q45. Patient was always offered practical advice on dealing with any immediate side effects from treatment	69%	67%	*	*	*	*	<b>68%</b>
Q46. Patient was given information that they could access about support in dealing with immediate side effects from treatment	91%	90%	*	*	*	*	<b>90%</b>
Q47. Patient felt possible long-term side effects were definitely explained in a way they could understand in advance of their treatment	68%	61%	*	*	*	*	<b>64%</b>
Q48. Patient was definitely able to discuss options for managing the impact of any long-term side effects	64%	51%	*	*	*	*	<b>57%</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	59%	63%	*	*	*	*	<b>60%</b>
Q50. During treatment, the patient definitely got enough care and support at home from community or voluntary services	46%	49%	*	*	*	*	<b>47%</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	43%	38%	*	*	*	*	<b>40%</b>
Q52. Patient has had a review of cancer care by GP practice	23%	22%	*	*	*	*	<b>22%</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	37%	25%	*	*	*	*	<b>32%</b>
Q54. The right amount of information and support was offered to the patient between final treatment and the follow up appointment	89%	83%	*	*	*	*	<b>86%</b>
Q55. Patient was given enough information about the possibility and signs of cancer coming back or spreading	66%	65%	*	*	*	*	<b>65%</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	94%	88%	*	*	*	*	<b>91%</b>
Q57. Administration of care was very good or good	93%	91%	*	*	*	*	<b>92%</b>
Q58. Cancer research opportunities were discussed with patient	40%	53%	*	*	*	*	<b>47%</b>
Q59. Patient's average rating of care scored from very poor to very good	9.1	8.8	*	*	*	*	<b>8.9</b>

## Ethnicity tables

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

	Ethnicity						All
	White	Mixed	Asian	Black	Other	Not given	
<b>SUPPORT FROM YOUR GP PRACTICE</b>							
Q2. Patient only spoke to primary care professional once or twice before cancer diagnosis	84%	*	*	*	*	*	<b>79%</b>
Q3. Referral for diagnosis was explained in a way the patient could completely understand	72%	*	42%	*	*	*	<b>69%</b>

	Ethnicity						All
	White	Mixed	Asian	Black	Other	Not given	
<b>DIAGNOSTIC TESTS</b>							
Q5. Patient received all the information needed about the diagnostic test in advance	89%	*	77%	*	*	*	<b>88%</b>
Q6. Diagnostic test staff appeared to completely have all the information they needed about the patient	89%	*	64%	*	*	*	<b>85%</b>
Q7. Patient felt the length of time waiting for diagnostic test results was about right	87%	*	64%	*	*	*	<b>85%</b>
Q8. Diagnostic test results were explained in a way the patient could completely understand	82%	*	64%	*	*	*	<b>82%</b>
Q9. Enough privacy was always given to the patient when receiving diagnostic test results	95%	*	86%	*	*	*	<b>95%</b>

	Ethnicity						All
	White	Mixed	Asian	Black	Other	Not given	
<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%	*	67%	*	*	90%	<b>77%</b>
Q13. Patient was definitely told sensitively that they had cancer	75%	*	50%	*	*	73%	<b>73%</b>
Q14. Cancer diagnosis explained in a way the patient could completely understand	78%	*	63%	*	*	82%	<b>77%</b>
Q15. Patient was definitely told about their diagnosis in an appropriate place	87%	*	87%	*	*	100%	<b>88%</b>
Q16. Patient was told they could go back later for more information about their diagnosis	86%	*	93%	*	*	*	<b>87%</b>

	Ethnicity						All
	White	Mixed	Asian	Black	Other	Not given	
<b>SUPPORT FROM A MAIN CONTACT PERSON</b>							
Q17. Patient had a main point of contact within the care team	94%	*	100%	*	*	*	<b>93%</b>
Q18. Patient found it very or quite easy to contact their main contact person	89%	*	50%	*	*	*	<b>86%</b>
Q19. Patient found advice from main contact person was very or quite helpful	97%	*	93%	*	*	*	<b>97%</b>

	Ethnicity						All
	White	Mixed	Asian	Black	Other	Not given	
<b>DECIDING ON THE BEST TREATMENT</b>							
Q20. Treatment options were explained in a way the patient could completely understand	86%	*	60%	*	*	82%	<b>84%</b>
Q21. Patient was definitely involved as much as they wanted to be in decisions about their treatment	80%	*	56%	*	*	91%	<b>79%</b>
Q22. Family and/or carers were definitely involved as much as the patient wanted them to be in decisions about treatment options	76%	*	63%	*	*	*	<b>75%</b>
Q23. Patient could get further advice or a second opinion before making decisions about their treatment options	49%	*	36%	*	*	*	<b>47%</b>

## Ethnicity tables

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

<b>CARE PLANNING</b>	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	74%	*	31%	*	*	80%	<b>70%</b>
Q25. A member of their care team helped the patient create a care plan to address any needs or concerns	92%	*	90%	*	*	*	<b>92%</b>
Q26. Care team reviewed the patient's care plan with them to ensure it was up to date	100%	*	*	*	*	*	<b>100%</b>

<b>SUPPORT FROM HOSPITAL STAFF</b>	Ethnicity						
	White	Mixed	Asian	Black	Other	Not given	All
Q27. Staff provided the patient with relevant information on available support	93%	*	87%	*	*	90%	<b>92%</b>
Q28. Patient definitely got the right level of support for their overall health and well being from hospital staff	78%	*	75%	*	*	73%	<b>77%</b>
Q29. Patient was offered information about how to get financial help or benefits	73%	*	64%	*	*	*	<b>70%</b>

<b>HOSPITAL CARE</b>	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	81%	*	75%	*	*	*	<b>82%</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	55%	*	33%	*	*	*	<b>53%</b>
Q33. Patient was always involved in decisions about their care and treatment whilst in hospital	72%	*	58%	*	*	*	<b>71%</b>
Q34. Patient was always able to get help from ward staff when needed	74%	*	67%	*	*	*	<b>75%</b>
Q35. Patient was always able to discuss worries and fears with hospital staff	68%	*	50%	*	*	*	<b>66%</b>
Q36. Hospital staff always did everything they could to help the patient control pain	82%	*	67%	*	*	*	<b>81%</b>
Q37. Patient was always treated with respect and dignity while in hospital	89%	*	92%	*	*	*	<b>90%</b>
Q38. Patient received easily understandable information about what they should or should not do after leaving hospital	87%	*	75%	*	*	*	<b>86%</b>
Q39. Patient was always able to discuss worries and fears with hospital staff while being treated as an outpatient or day case	78%	*	50%	*	*	82%	<b>75%</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	94%	*	64%	*	*	*	<b>91%</b>
Q41_2. Beforehand patient completely had enough understandable information about chemotherapy	82%	*	50%	*	*	*	<b>79%</b>
Q41_3. Beforehand patient completely had enough understandable information about radiotherapy	90%	*	*	*	*	*	<b>89%</b>
Q41_4. Beforehand patient completely had enough understandable information about hormone therapy	80%	*	*	*	*	*	<b>76%</b>
Q41_5. Beforehand patient completely had enough understandable information about immunotherapy	*	*	*	*	*	*	*
Q42_1. Patient completely had enough understandable information about progress with surgery	89%	*	64%	*	*	*	<b>87%</b>
Q42_2. Patient completely had enough understandable information about progress with chemotherapy	77%	*	60%	*	*	*	<b>75%</b>
Q42_3. Patient completely had enough understandable information about progress with radiotherapy	82%	*	*	*	*	*	<b>80%</b>
Q42_4. Patient completely had enough understandable information about progress with hormone therapy	75%	*	*	*	*	*	<b>72%</b>
Q42_5. Patient completely had enough understandable information about progress with immunotherapy	*	*	*	*	*	*	*
Q43. Patient felt the length of waiting time at clinic and day unit for cancer treatment was about right	90%	*	69%	*	*	91%	<b>87%</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	77%	*	63%	*	*	90%	<b>76%</b>
Q45. Patient was always offered practical advice on dealing with any immediate side effects from treatment	69%	*	43%	*	*	70%	<b>68%</b>
Q46. Patient was given information that they could access about support in dealing with immediate side effects from treatment	92%	*	73%	*	*	*	<b>90%</b>
Q47. Patient felt possible long-term side effects were definitely explained in a way they could understand in advance of their treatment	66%	*	44%	*	*	73%	<b>64%</b>
Q48. Patient was definitely able to discuss options for managing the impact of any long-term side effects	61%	*	27%	*	*	*	<b>57%</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	61%	*	53%	*	*	*	<b>60%</b>
Q50. During treatment, the patient definitely got enough care and support at home from community or voluntary services	45%	*	50%	*	*	*	<b>47%</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%	*	36%	*	*	*	<b>40%</b>
Q52. Patient has had a review of cancer care by GP practice	23%	*	27%	*	*	10%	<b>22%</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	33%	*	*	*	*	*	<b>32%</b>
Q54. The right amount of information and support was offered to the patient between final treatment and the follow up appointment	89%	*	50%	*	*	*	<b>86%</b>
Q55. Patient was given enough information about the possibility and signs of cancer coming back or spreading	67%	*	43%	*	*	*	<b>65%</b>

<b>YOUR OVERALL NHS CARE</b>	Ethnicity						
	White	Mixed	Asian	Black	Other	Not given	All
Q56. The whole care team worked well together	90%	*	94%	*	*	*	<b>91%</b>
Q57. Administration of care was very good or good	91%	*	94%	*	*	100%	<b>92%</b>
Q58. Cancer research opportunities were discussed with patient	44%	*	*	*	*	*	<b>47%</b>
Q59. Patient's average rating of care scored from very poor to very good	9.0	*	8.3	*	*	9.7	<b>8.9</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)	Non-England	All
<b>SUPPORT FROM YOUR GP PRACTICE</b>							
Q2. Patient only spoke to primary care professional once or twice before cancer diagnosis	73%	67%	85%	87%	92%	*	<b>79%</b>
Q3. Referral for diagnosis was explained in a way the patient could completely understand	63%	48%	83%	72%	94%	*	<b>69%</b>

	IMD Quintile						
	1 (most deprived)	2	3	4	5 (least deprived)	Non-England	All
<b>DIAGNOSTIC TESTS</b>							
Q5. Patient received all the information needed about the diagnostic test in advance	80%	90%	90%	93%	94%	*	<b>88%</b>
Q6. Diagnostic test staff appeared to completely have all the information they needed about the patient	78%	87%	84%	90%	95%	*	<b>85%</b>
Q7. Patient felt the length of time waiting for diagnostic test results was about right	82%	81%	90%	87%	89%	*	<b>85%</b>
Q8. Diagnostic test results were explained in a way the patient could completely understand	72%	78%	94%	90%	84%	*	<b>82%</b>
Q9. Enough privacy was always given to the patient when receiving diagnostic test results	90%	97%	97%	94%	100%	*	<b>95%</b>

	IMD Quintile						
	1 (most deprived)	2	3	4	5 (least deprived)	Non-England	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	71%	82%	76%	83%	74%	*	<b>77%</b>
Q13. Patient was definitely told sensitively that they had cancer	70%	68%	76%	74%	83%	*	<b>73%</b>
Q14. Cancer diagnosis explained in a way the patient could completely understand	67%	85%	82%	77%	83%	*	<b>77%</b>
Q15. Patient was definitely told about their diagnosis in an appropriate place	86%	85%	88%	89%	96%	*	<b>88%</b>
Q16. Patient was told they could go back later for more information about their diagnosis	84%	82%	91%	93%	84%	*	<b>87%</b>

	IMD Quintile						
	1 (most deprived)	2	3	4	5 (least deprived)	Non-England	All
<b>SUPPORT FROM A MAIN CONTACT PERSON</b>							
Q17. Patient had a main point of contact within the care team	95%	90%	97%	94%	90%	*	<b>93%</b>
Q18. Patient found it very or quite easy to contact their main contact person	83%	84%	89%	83%	100%	*	<b>86%</b>
Q19. Patient found advice from main contact person was very or quite helpful	92%	97%	100%	100%	100%	*	<b>97%</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)	Non-England	All
<b>DECIDING ON THE BEST TREATMENT</b>							
Q20. Treatment options were explained in a way the patient could completely understand	85%	79%	85%	88%	83%	*	<b>84%</b>
Q21. Patient was definitely involved as much as they wanted to be in decisions about their treatment	68%	80%	85%	86%	83%	*	<b>79%</b>
Q22. Family and/or carers were definitely involved as much as the patient wanted them to be in decisions about treatment options	64%	72%	71%	93%	90%	*	<b>75%</b>
Q23. Patient could get further advice or a second opinion before making decisions about their treatment options	45%	36%	67%	33%	60%	*	<b>47%</b>

	IMD Quintile						
	1 (most deprived)	2	3	4	5 (least deprived)	Non-England	All
<b>CARE PLANNING</b>							
Q24. Patient was definitely able to have a discussion about their needs or concerns prior to treatment	65%	58%	78%	87%	71%	*	<b>70%</b>
Q25. A member of their care team helped the patient create a care plan to address any needs or concerns	95%	91%	88%	89%	93%	*	<b>92%</b>
Q26. Care team reviewed the patient's care plan with them to ensure it was up to date	100%	100%	100%	100%	100%	*	<b>100%</b>

	IMD Quintile						
	1 (most deprived)	2	3	4	5 (least deprived)	Non-England	All
<b>SUPPORT FROM HOSPITAL STAFF</b>							
Q27. Staff provided the patient with relevant information on available support	87%	91%	94%	97%	95%	*	<b>92%</b>
Q28. Patient definitely got the right level of support for their overall health and well being from hospital staff	78%	71%	85%	71%	86%	*	<b>77%</b>
Q29. Patient was offered information about how to get financial help or benefits	65%	73%	71%	67%	*	*	<b>70%</b>

	IMD Quintile						
	1 (most deprived)	2	3	4	5 (least deprived)	Non-England	All
<b>HOSPITAL CARE</b>							
Q31. Patient had confidence and trust in all of the team looking after them during their stay in hospital	74%	80%	88%	86%	86%	*	<b>82%</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	57%	67%	47%	38%	57%	*	<b>53%</b>
Q33. Patient was always involved in decisions about their care and treatment whilst in hospital	74%	65%	63%	75%	85%	*	<b>71%</b>
Q34. Patient was always able to get help from ward staff when needed	71%	70%	87%	75%	71%	*	<b>75%</b>
Q35. Patient was always able to discuss worries and fears with hospital staff	62%	72%	64%	67%	71%	*	<b>66%</b>
Q36. Hospital staff always did everything they could to help the patient control pain	76%	74%	90%	85%	85%	*	<b>81%</b>
Q37. Patient was always treated with respect and dignity while in hospital	94%	90%	83%	85%	100%	*	<b>90%</b>
Q38. Patient received easily understandable information about what they should or should not do after leaving hospital	86%	85%	90%	90%	79%	*	<b>86%</b>
Q39. Patient was always able to discuss worries and fears with hospital staff while being treated as an outpatient or day case	63%	79%	85%	76%	79%	*	<b>75%</b>

## IMD quintile tables

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

<b>YOUR TREATMENT</b>	IMD Quintile						
	1 (most deprived)	2	3	4	5 (least deprived)	Non-England	All
Q41_1. Beforehand patient completely had enough understandable information about surgery	83%	91%	100%	96%	85%	*	<b>91%</b>
Q41_2. Beforehand patient completely had enough understandable information about chemotherapy	73%	70%	87%	*	*	*	<b>79%</b>
Q41_3. Beforehand patient completely had enough understandable information about radiotherapy	76%	92%	93%	*	*	*	<b>89%</b>
Q41_4. Beforehand patient completely had enough understandable information about hormone therapy	70%	*	*	*	*	*	<b>76%</b>
Q41_5. Beforehand patient completely had enough understandable information about immunotherapy	*	*	*	*	*	*	*
Q42_1. Patient completely had enough understandable information about progress with surgery	80%	83%	96%	96%	80%	*	<b>87%</b>
Q42_2. Patient completely had enough understandable information about progress with chemotherapy	68%	80%	78%	80%	*	*	<b>75%</b>
Q42_3. Patient completely had enough understandable information about progress with radiotherapy	76%	91%	79%	*	*	*	<b>80%</b>
Q42_4. Patient completely had enough understandable information about progress with hormone therapy	60%	*	*	*	*	*	<b>72%</b>
Q42_5. Patient completely had enough understandable information about progress with immunotherapy	*	*	*	*	*	*	*
Q43. Patient felt the length of waiting time at clinic and day unit for cancer treatment was about right	84%	83%	94%	89%	91%	*	<b>87%</b>

<b>IMMEDIATE AND LONG TERM SIDE EFFECTS</b>	IMD Quintile						
	1 (most deprived)	2	3	4	5 (least deprived)	Non-England	All
Q44. Possible side effects from treatment were definitely explained in a way the patient could understand	73%	72%	82%	81%	75%	*	<b>76%</b>
Q45. Patient was always offered practical advice on dealing with any immediate side effects from treatment	61%	66%	71%	80%	68%	*	<b>68%</b>
Q46. Patient was given information that they could access about support in dealing with immediate side effects from treatment	85%	85%	97%	93%	100%	*	<b>90%</b>
Q47. Patient felt possible long-term side effects were definitely explained in a way they could understand in advance of their treatment	60%	64%	81%	58%	56%	*	<b>64%</b>
Q48. Patient was definitely able to discuss options for managing the impact of any long-term side effects	55%	52%	61%	54%	69%	*	<b>57%</b>

<b>SUPPORT WHILE AT HOME</b>	IMD Quintile						
	1 (most deprived)	2	3	4	5 (least deprived)	Non-England	All
Q49. Care team gave family, or someone close, all the information needed to help care for the patient at home	60%	68%	43%	67%	60%	*	<b>60%</b>
Q50. During treatment, the patient definitely got enough care and support at home from community or voluntary services	43%	44%	50%	70%	*	*	<b>47%</b>

<b>CARE FROM YOUR GP PRACTICE</b>	IMD Quintile						
	1 (most deprived)	2	3	4	5 (least deprived)	Non-England	All
Q51. Patient definitely received the right amount of support from their GP practice during treatment	35%	26%	50%	50%	36%	*	<b>40%</b>
Q52. Patient has had a review of cancer care by GP practice	24%	13%	24%	26%	23%	*	<b>22%</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)	Non-England	All
Q53. After treatment, the patient definitely could get enough emotional support at home from community or voluntary services	30%	20%	*	*	*	*	<b>32%</b>
Q54. The right amount of information and support was offered to the patient between final treatment and the follow up appointment	71%	90%	95%	88%	94%	*	<b>86%</b>
Q55. Patient was given enough information about the possibility and signs of cancer coming back or spreading	63%	71%	59%	76%	50%	*	<b>65%</b>

	IMD Quintile						
	1 (most deprived)	2	3	4	5 (least deprived)	Non-England	All
Q56. The whole care team worked well together	91%	90%	88%	91%	95%	*	<b>91%</b>
Q57. Administration of care was very good or good	92%	93%	94%	89%	95%	*	<b>92%</b>
Q58. Cancer research opportunities were discussed with patient	44%	58%	32%	57%	44%	*	<b>47%</b>
Q59. Patient's average rating of care scored from very poor to very good	8.8	8.9	9.2	8.8	9.3	*	<b>8.9</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	76%	80%	*	79%
Q3. Referral for diagnosis was explained in a way the patient could completely understand	68%	70%	*	69%

	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	89%	84%	*	88%
Q6. Diagnostic test staff appeared to completely have all the information they needed about the patient	82%	91%	*	85%
Q7. Patient felt the length of time waiting for diagnostic test results was about right	84%	88%	*	85%
Q8. Diagnostic test results were explained in a way the patient could completely understand	81%	81%	*	82%
Q9. Enough privacy was always given to the patient when receiving diagnostic test results	93%	97%	*	95%

	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	80%	73%	70%	77%
Q13. Patient was definitely told sensitively that they had cancer	74%	72%	73%	73%
Q14. Cancer diagnosis explained in a way the patient could completely understand	78%	75%	82%	77%
Q15. Patient was definitely told about their diagnosis in an appropriate place	87%	88%	100%	88%
Q16. Patient was told they could go back later for more information about their diagnosis	88%	84%	*	87%

	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	95%	91%	90%	93%
Q18. Patient found it very or quite easy to contact their main contact person	85%	88%	*	86%
Q19. Patient found advice from main contact person was very or quite helpful	98%	95%	*	97%

	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	84%	84%	82%	84%
Q21. Patient was definitely involved as much as they wanted to be in decisions about their treatment	80%	76%	82%	79%
Q22. Family and/or carers were definitely involved as much as the patient wanted them to be in decisions about treatment options	74%	75%	*	75%
Q23. Patient could get further advice or a second opinion before making decisions about their treatment options	46%	52%	*	47%

## 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	73%	66%	70%	<b>70%</b>
Q25. A member of their care team helped the patient create a care plan to address any needs or concerns	94%	91%	*	<b>92%</b>
Q26. Care team reviewed the patient's care plan with them to ensure it was up to date	100%	100%	*	<b>100%</b>

<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	94%	90%	80%	<b>92%</b>
Q28. Patient definitely got the right level of support for their overall health and well being from hospital staff	75%	82%	73%	<b>77%</b>
Q29. Patient was offered information about how to get financial help or benefits	68%	79%	*	<b>70%</b>

<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	83%	80%	*	<b>82%</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	53%	53%	*	<b>53%</b>
Q33. Patient was always involved in decisions about their care and treatment whilst in hospital	70%	70%	*	<b>71%</b>
Q34. Patient was always able to get help from ward staff when needed	73%	77%	*	<b>75%</b>
Q35. Patient was always able to discuss worries and fears with hospital staff	62%	68%	*	<b>66%</b>
Q36. Hospital staff always did everything they could to help the patient control pain	79%	83%	*	<b>81%</b>
Q37. Patient was always treated with respect and dignity while in hospital	88%	93%	*	<b>90%</b>
Q38. Patient received easily understandable information about what they should or should not do after leaving hospital	85%	87%	*	<b>86%</b>
Q39. Patient was always able to discuss worries and fears with hospital staff while being treated as an outpatient or day case	73%	75%	*	<b>75%</b>

## Long term condition status tables

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

<b>YOUR TREATMENT</b>	Long term condition status			
	Yes	No	Not given	All
Q41_1. Beforehand patient completely had enough understandable information about surgery	91%	91%	*	<b>91%</b>
Q41_2. Beforehand patient completely had enough understandable information about chemotherapy	74%	86%	*	<b>79%</b>
Q41_3. Beforehand patient completely had enough understandable information about radiotherapy	88%	94%	*	<b>89%</b>
Q41_4. Beforehand patient completely had enough understandable information about hormone therapy	79%	*	*	<b>76%</b>
Q41_5. Beforehand patient completely had enough understandable information about immunotherapy	*	*	*	*
Q42_1. Patient completely had enough understandable information about progress with surgery	87%	87%	*	<b>87%</b>
Q42_2. Patient completely had enough understandable information about progress with chemotherapy	75%	78%	*	<b>75%</b>
Q42_3. Patient completely had enough understandable information about progress with radiotherapy	80%	80%	*	<b>80%</b>
Q42_4. Patient completely had enough understandable information about progress with hormone therapy	71%	*	*	<b>72%</b>
Q42_5. Patient completely had enough understandable information about progress with immunotherapy	*	*	*	*
Q43. Patient felt the length of waiting time at clinic and day unit for cancer treatment was about right	87%	88%	82%	<b>87%</b>

<b>IMMEDIATE AND LONG TERM SIDE EFFECTS</b>	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	75%	77%	80%	<b>76%</b>
Q45. Patient was always offered practical advice on dealing with any immediate side effects from treatment	66%	68%	80%	<b>68%</b>
Q46. Patient was given information that they could access about support in dealing with immediate side effects from treatment	90%	91%	*	<b>90%</b>
Q47. Patient felt possible long-term side effects were definitely explained in a way they could understand in advance of their treatment	66%	60%	64%	<b>64%</b>
Q48. Patient was definitely able to discuss options for managing the impact of any long-term side effects	56%	57%	*	<b>57%</b>

<b>SUPPORT WHILE AT HOME</b>	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	68%	47%	*	<b>60%</b>
Q50. During treatment, the patient definitely got enough care and support at home from community or voluntary services	43%	50%	*	<b>47%</b>

<b>CARE FROM YOUR GP PRACTICE</b>	Long term condition status			
	Yes	No	Not given	All
Q51. Patient definitely received the right amount of support from their GP practice during treatment	35%	49%	*	<b>40%</b>
Q52. Patient has had a review of cancer care by GP practice	18%	28%	18%	<b>22%</b>



## 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	<b>All</b>
Q53. After treatment, the patient definitely could get enough emotional support at home from community or voluntary services	27%	35%	*	<b>32%</b>
Q54. The right amount of information and support was offered to the patient between final treatment and the follow up appointment	83%	88%	*	<b>86%</b>
Q55. Patient was given enough information about the possibility and signs of cancer coming back or spreading	62%	70%	*	<b>65%</b>

<b>YOUR OVERALL NHS CARE</b>	Long term condition status			
	Yes	No	Not given	<b>All</b>
Q56. The whole care team worked well together	91%	91%	*	<b>91%</b>
Q57. Administration of care was very good or good	93%	90%	100%	<b>92%</b>
Q58. Cancer research opportunities were discussed with patient	48%	44%	*	<b>47%</b>
Q59. Patient's average rating of care scored from very poor to very good	8.8	9.1	9.5	<b>8.9</b>

## Year on Year Charts

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

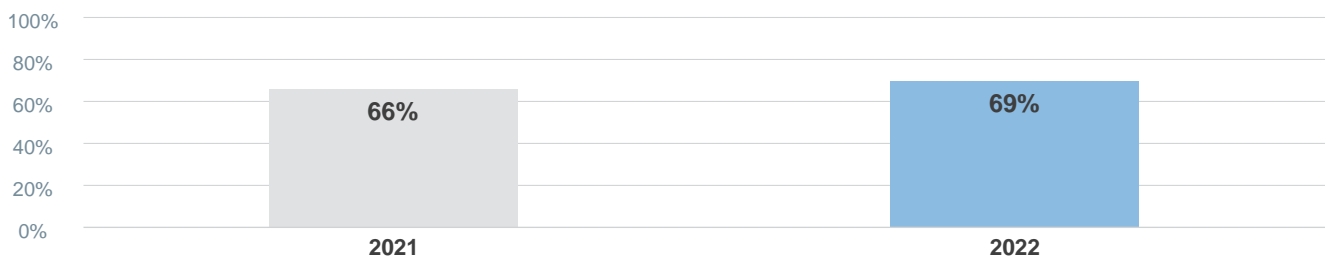
The scores are unadjusted and based on England scores only.

### 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



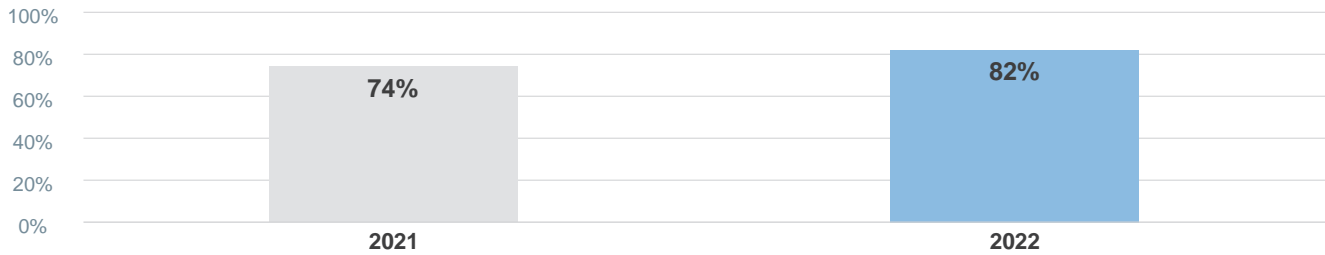
Q7. Patient felt the length of time waiting for diagnostic test results was about right



## Year on Year Charts

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

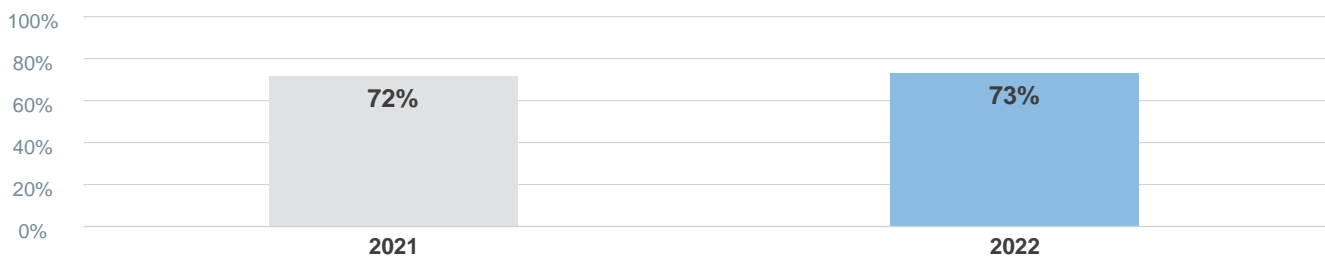


### FINDING OUT THAT YOU HAD CANCER

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.

The scores are unadjusted and based on England scores only.

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

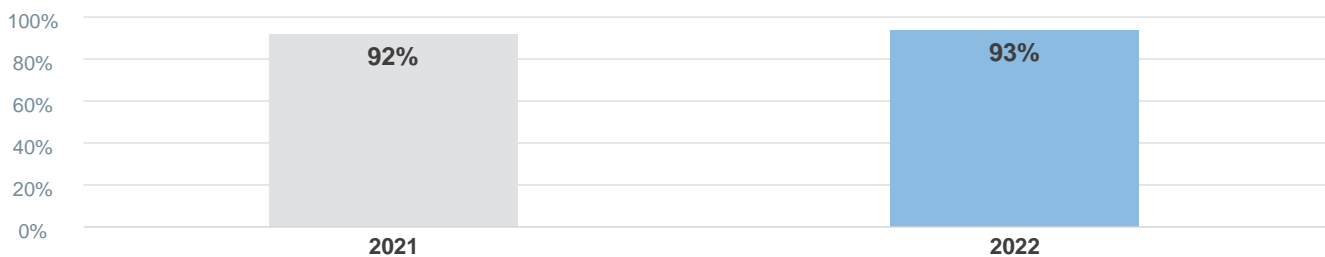


Q16. Patient was told they could go back later for more information about their diagnosis

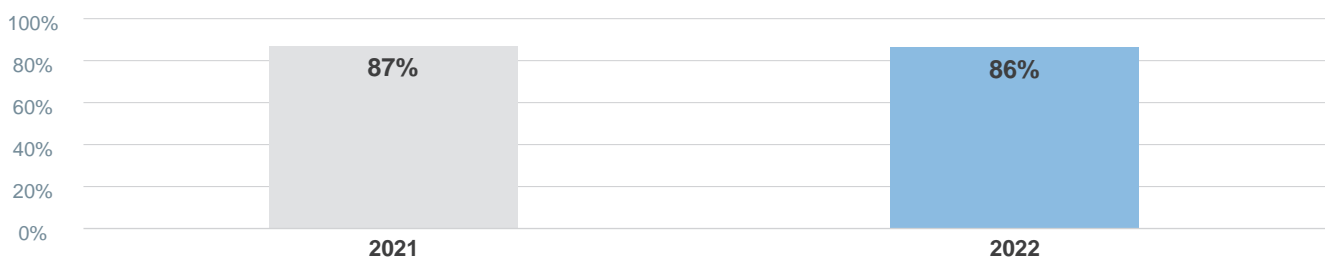


### SUPPORT FROM A MAIN CONTACT PERSON

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.

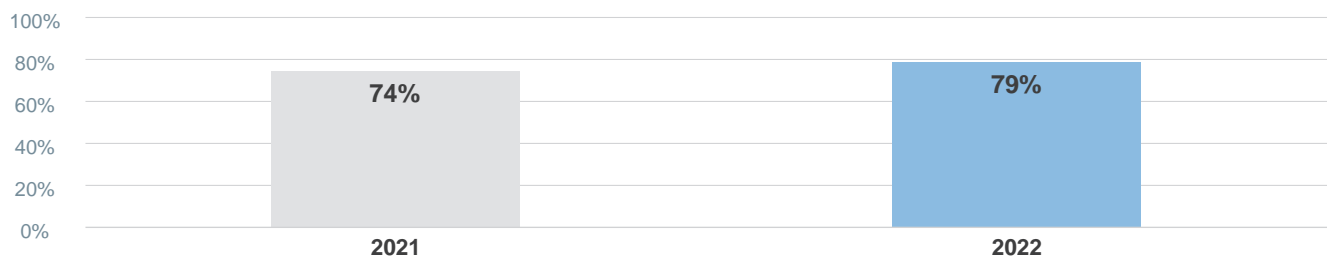
The scores are unadjusted and based on England scores only.

### DECIDING ON THE BEST TREATMENT

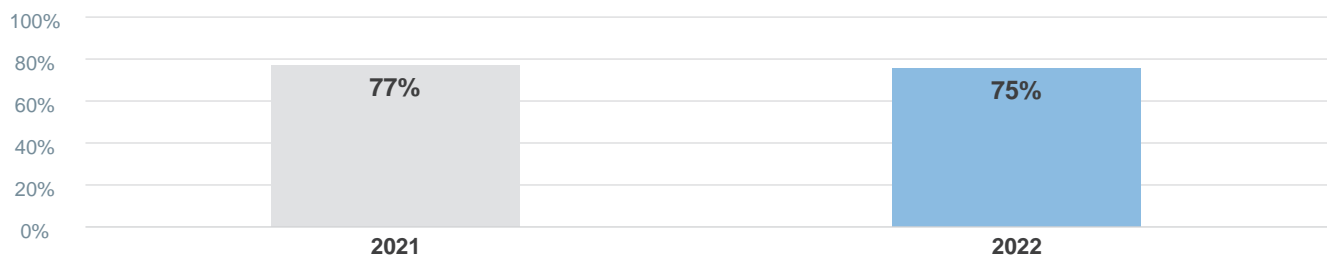
Q20. Treatment options were explained in a way the patient could completely understand



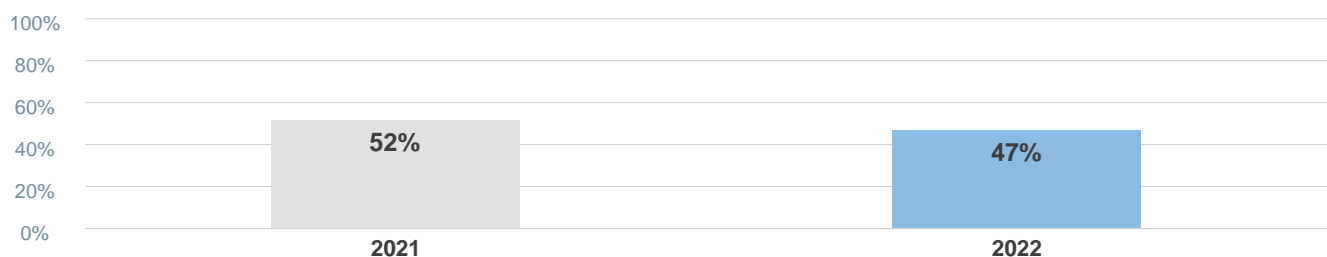
Q21. Patient was definitely involved as much as they wanted to be in decisions about their treatment



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

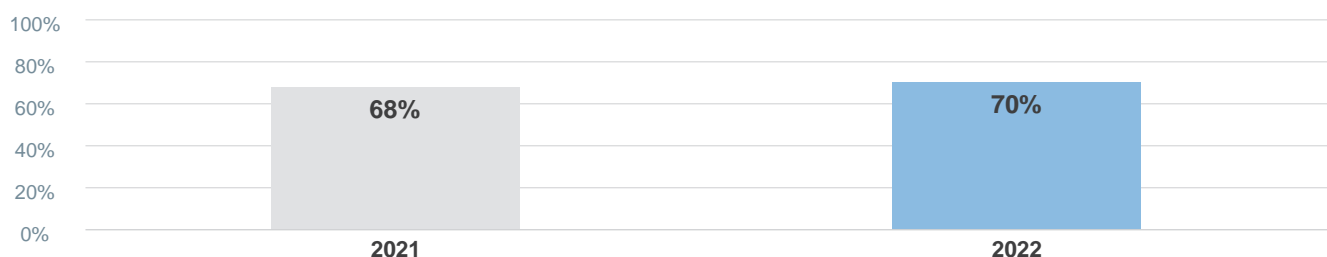


Q23. Patient could get further advice or a second opinion before making decisions about their treatment options



### 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. The scores are unadjusted and based on England scores only.

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

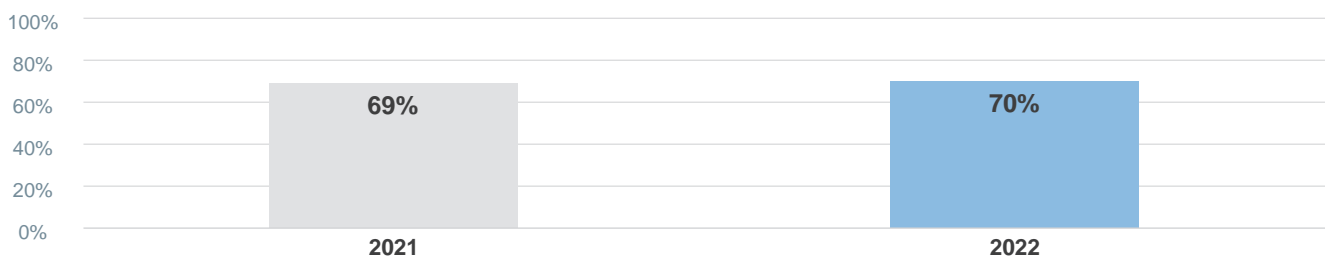
Q27. Staff provided the patient with relevant information on available support



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.

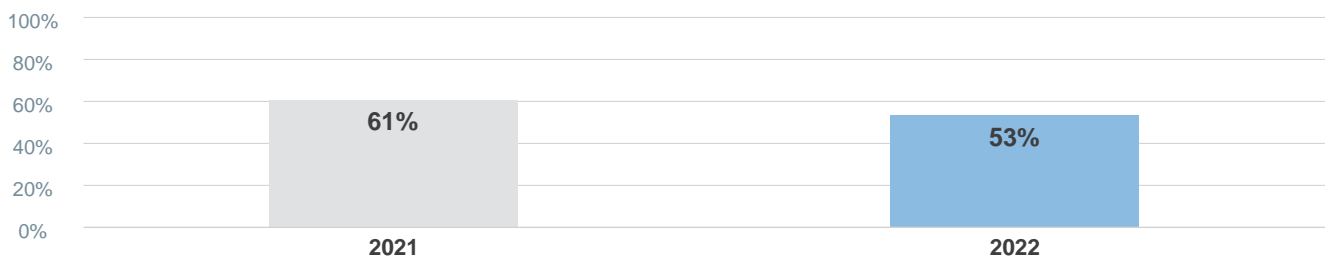
The scores are unadjusted and based on England scores only.

### HOSPITAL CARE

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



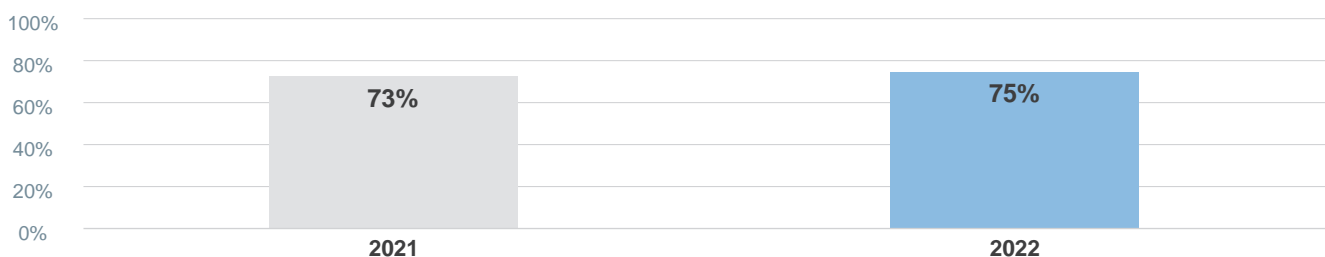
Q32. Patient's family, or someone close, was definitely able to talk to a member of the team looking after the patient in hospital



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



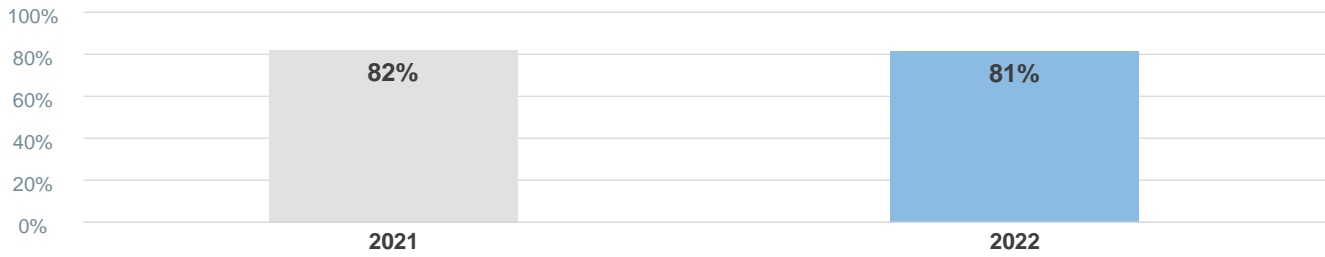
Q35. Patient was always able to discuss worries and fears with hospital staff



## Year on Year Charts

\* Indicates where a score is not available due to suppression or a low base size. The scores are unadjusted and based on England scores only.

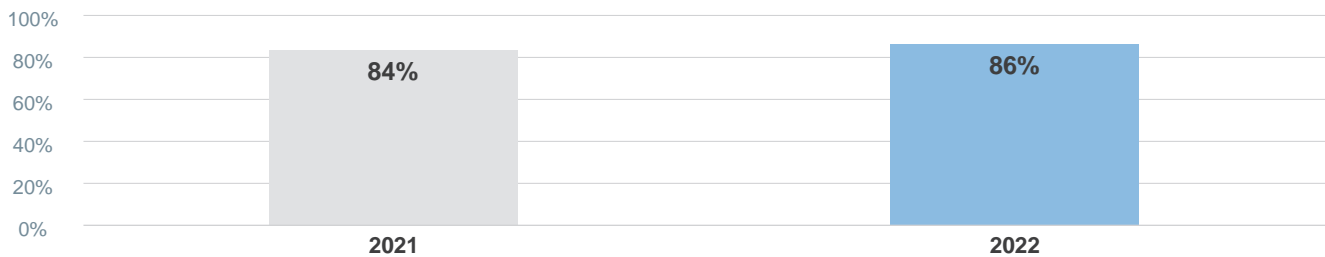
**Q36. Hospital staff always did everything they could to help the patient control pain**



**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**

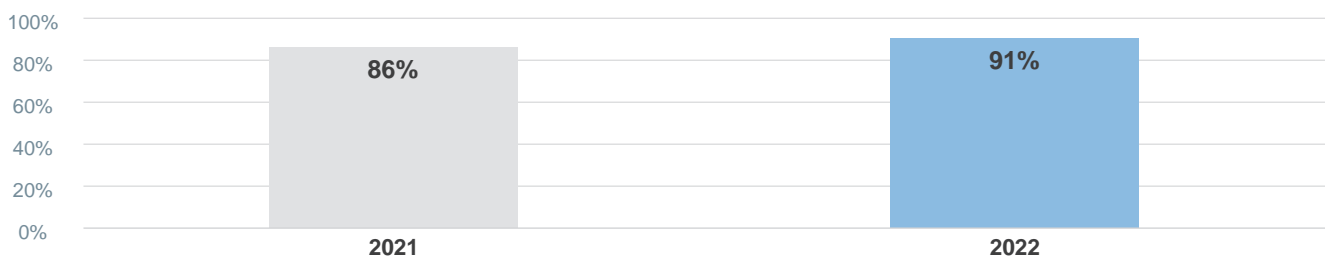


**Q39. Patient was always able to discuss worries and fears with hospital staff while being treated as an outpatient or day case**



### 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. 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 progress with surgery



## Year on Year Charts

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**Q42\_2. Patient completely had enough understandable information about progress with chemotherapy**



**Q42\_3. Patient completely had enough understandable information about progress with radiotherapy**



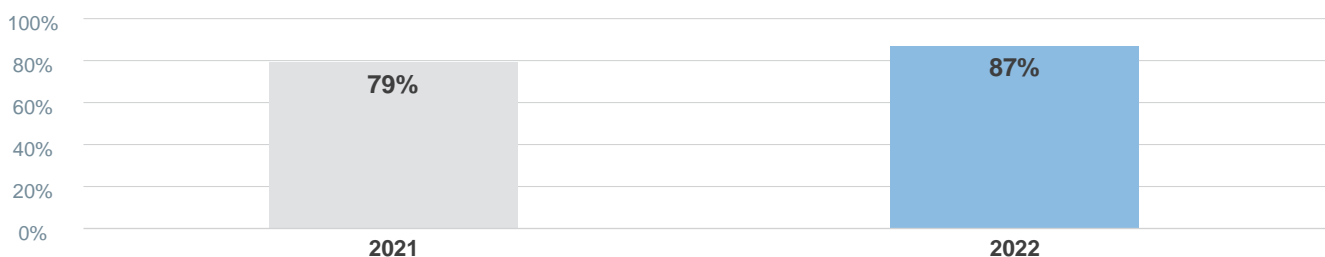
**Q42\_4. Patient completely had enough understandable information about progress with hormone therapy**



**Q42\_5. Patient completely had enough understandable information about progress with immunotherapy**



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



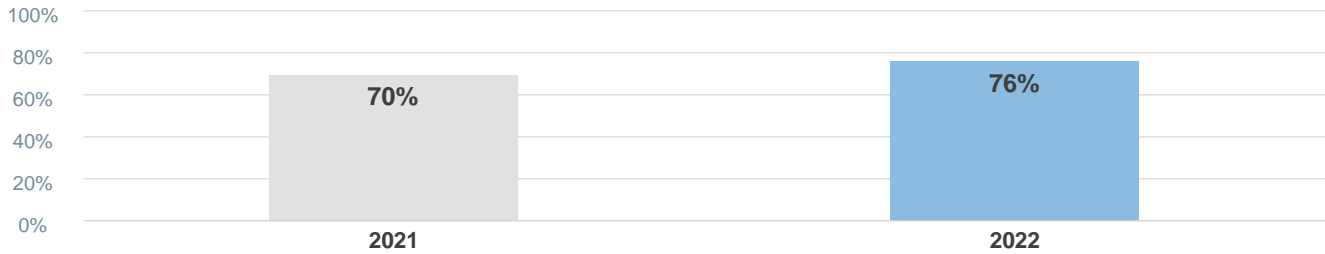
## Year on Year Charts

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### 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



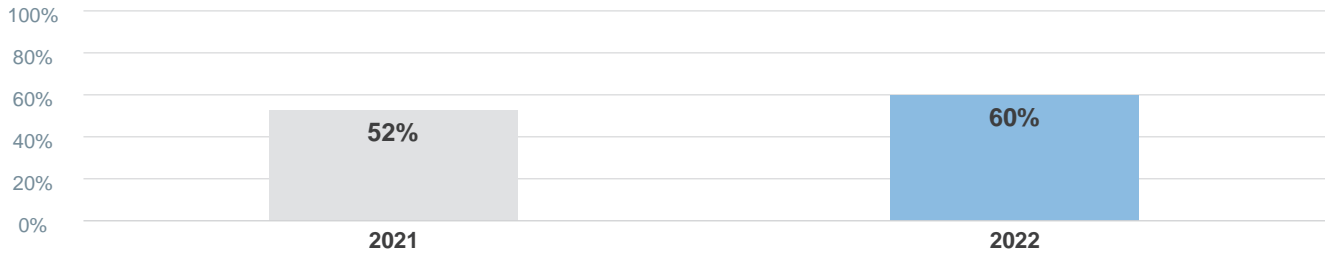
## Year on Year Charts

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### 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

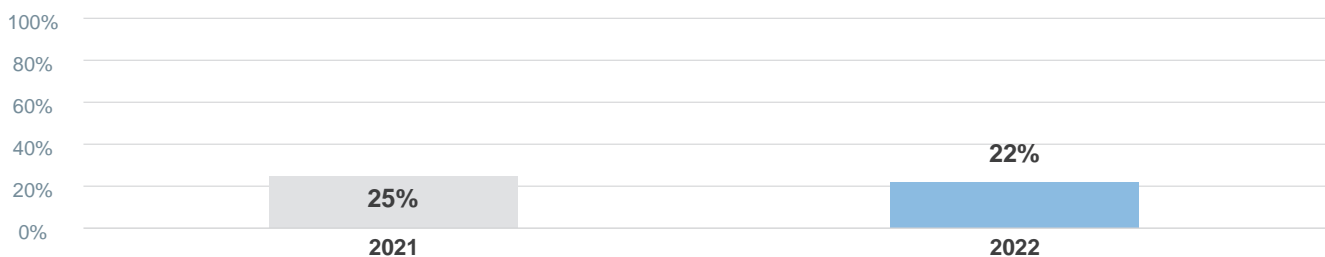


### CARE FROM YOUR GP PRACTICE

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

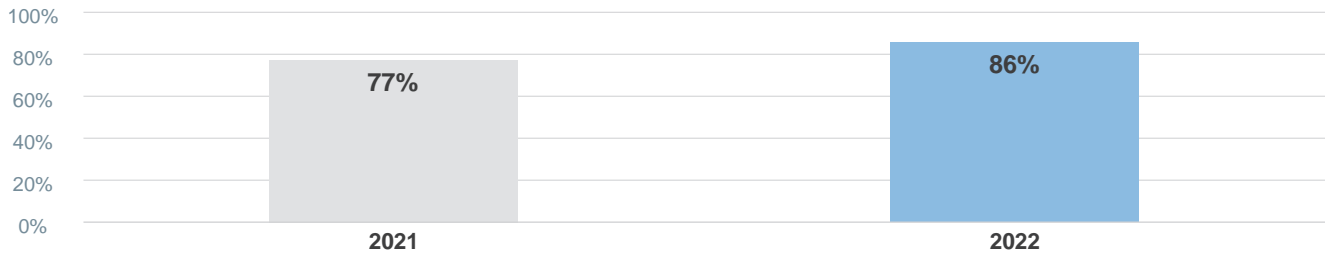
Q53. After treatment, the patient definitely could get enough emotional support at home from community or voluntary services



## Year on Year Charts

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**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

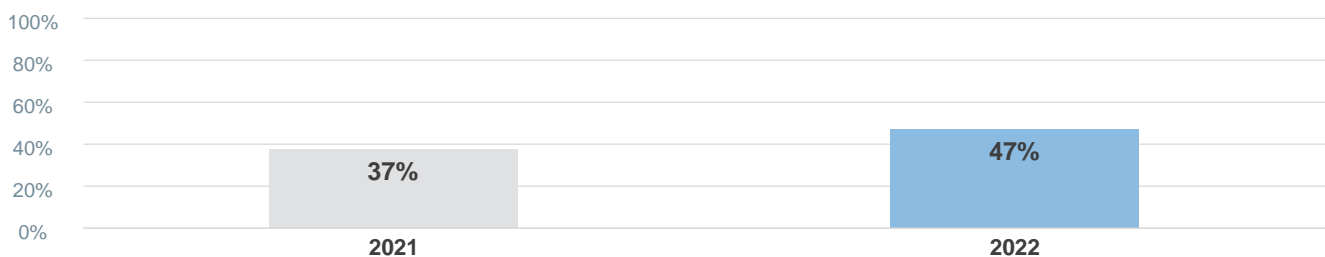
**Q56. The whole care team worked well together**



**Q57. Administration of care was very good or good**



**Q58. Cancer research opportunities were discussed with patient**



## Year on Year Charts

\* Indicates where a score is not available due to suppression or a low base size. The scores are unadjusted and based on England scores only.

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

