

# **Cancer Patient Experience Survey**

2022 Results

## **NHS Hertfordshire and West Essex Integrated Care Board**

Published July 2023

## Executive Summary

NHS Hertfordshire and West Essex Integrated Care Board has no scores above expected range

### Questions Below Expected Range

	Case Mix Adjusted Scores			England Score
	2022 Score	Lower Expected Range	Upper Expected Range	
Q6. Diagnostic test staff appeared to completely have all the information they needed about the patient	<b>80%</b>	81%	86%	<b>83%</b>
Q13. Patient was definitely told sensitively that they had cancer	<b>70%</b>	70%	77%	<b>74%</b>
Q14. Cancer diagnosis explained in a way the patient could completely understand	<b>73%</b>	74%	79%	<b>76%</b>
Q21. Patient was definitely involved as much as they wanted to be in decisions about their treatment	<b>76%</b>	77%	82%	<b>79%</b>
Q24. Patient was definitely able to have a discussion about their needs or concerns prior to treatment	<b>68%</b>	68%	74%	<b>71%</b>
Q25. A member of their care team helped the patient create a care plan to address any needs or concerns	<b>90%</b>	91%	95%	<b>93%</b>
Q36. Hospital staff always did everything they could to help the patient control pain	<b>81%</b>	81%	88%	<b>84%</b>
Q41_3. Beforehand patient completely had enough understandable information about radiotherapy	<b>83%</b>	86%	91%	<b>88%</b>
Q41_4. Beforehand patient completely had enough understandable information about hormone therapy	<b>70%</b>	74%	84%	<b>79%</b>
Q42_1. Patient completely had enough understandable information about progress with surgery	<b>82%</b>	82%	87%	<b>85%</b>
Q42_2. Patient completely had enough understandable information about progress with chemotherapy	<b>74%</b>	76%	81%	<b>79%</b>
Q42_3. Patient completely had enough understandable information about progress with radiotherapy	<b>75%</b>	77%	84%	<b>81%</b>
Q42_4. Patient completely had enough understandable information about progress with hormone therapy	<b>65%</b>	67%	78%	<b>72%</b>
Q42_5. Patient completely had enough understandable information about progress with immunotherapy	<b>73%</b>	74%	85%	<b>79%</b>
Q44. Possible side effects from treatment were definitely explained in a way the patient could understand	<b>69%</b>	71%	77%	<b>74%</b>
Q45. Patient was always offered practical advice on dealing with any immediate side effects from treatment	<b>64%</b>	66%	73%	<b>69%</b>
Q46. Patient was given information that they could access about support in dealing with immediate side effects from treatment	<b>83%</b>	83%	89%	<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	<b>53%</b>	56%	62%	<b>59%</b>
Q48. Patient was definitely able to discuss options for managing the impact of any long-term side effects	<b>49%</b>	49%	57%	<b>53%</b>
Q49. Care team gave family, or someone close, all the information needed to help care for the patient at home	<b>54%</b>	54%	62%	<b>58%</b>
Q53. After treatment, the patient definitely could get enough emotional support at home from community or voluntary services	<b>24%</b>	25%	37%	<b>31%</b>
Q55. Patient was given enough information about the possibility and signs of cancer coming back or spreading	<b>57%</b>	59%	66%	<b>62%</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.

### How Alliance and ICB results are generated

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

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

### Case-mix adjustment

Both unadjusted and adjusted scores are presented in this report. Case-mix adjusted scores allow us to account for the impact that differing patient populations might have on results. By using the case-mix adjusted estimates we can obtain a greater understanding of how an ICB 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 ICB. 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 ICB, the results are not shown for that question for that ICB.

For ICBs 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** ICB has a score suppressed. If this happens, we will suppress another ICB's results (both the ICB 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 ICB.

The same rule applies to groups in each subgroup breakdown. For example, if only **one** ICB has the 85+ age group suppressed for Q25 we will need to suppress another ICB'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 ICB scored for each question in the survey compared with national results. It is aimed at helping individual ICBs to understand their performance and identify areas for local and regional 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 ICB.

ICBs 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 ICB performs better than what ICBs 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 ICB's size and demographics.

### Comparability tables

The comparability tables show the 2021 and 2022 unadjusted scores for this ICB 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 ICBs 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 ICB level, please see the Excel tables and dashboards at [www.ncpes.co.uk](http://www.ncpes.co.uk).

## Response Rate

### Overall Response Rate

1,516 patients responded out of a total of 2,912 patients, resulting in a response rate of 52%.

	Sample Size	Adjusted Sample	Completed	Response Rate
Overall response rate	3,095	2,912	1,516	52%
National	123,632	115,662	61,268	53%

### Respondents by Survey Type

	Number of Respondents
Paper	1,207
Online	307
Phone	2
Translation Service	0
<b>Total</b>	<b>1,516</b>

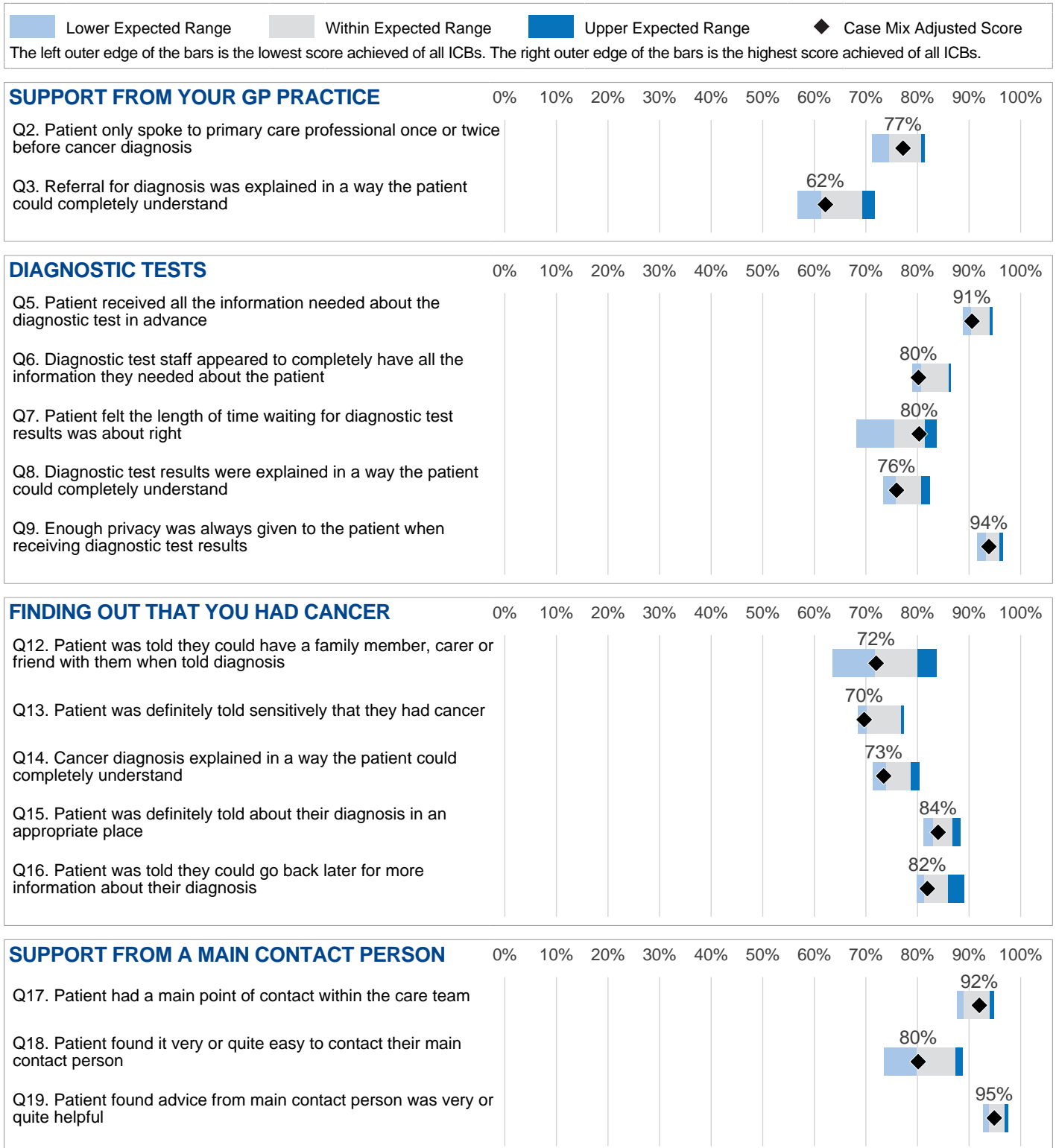
### Respondents by Tumour Group

	Number of Respondents
Brain / CNS	10
Breast	370
Colorectal / LGT	152
Gynaecological	103
Haematological	210
Head and Neck	51
Lung	69
Prostate	153
Sarcoma	8
Skin	55
Upper Gastro	75
Urological	96
Other	164
<b>Total</b>	<b>1,516</b>

## Respondents by Ethnicity

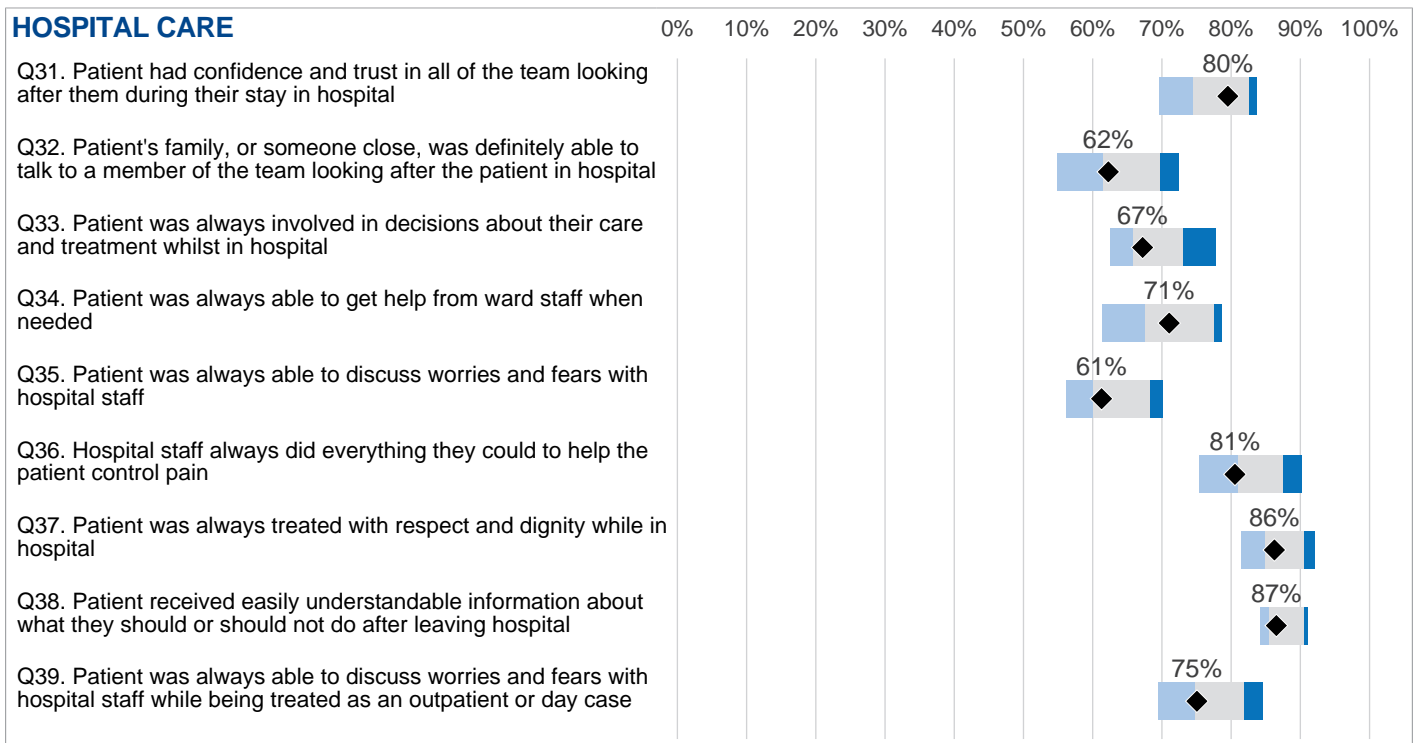
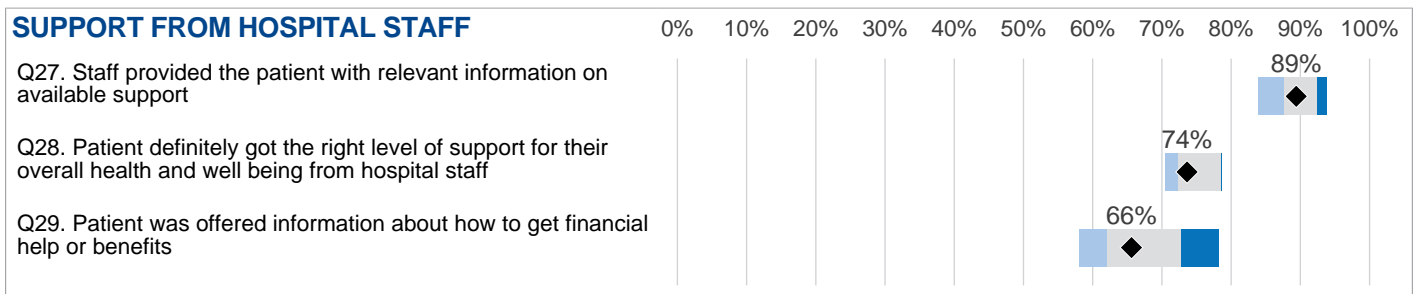
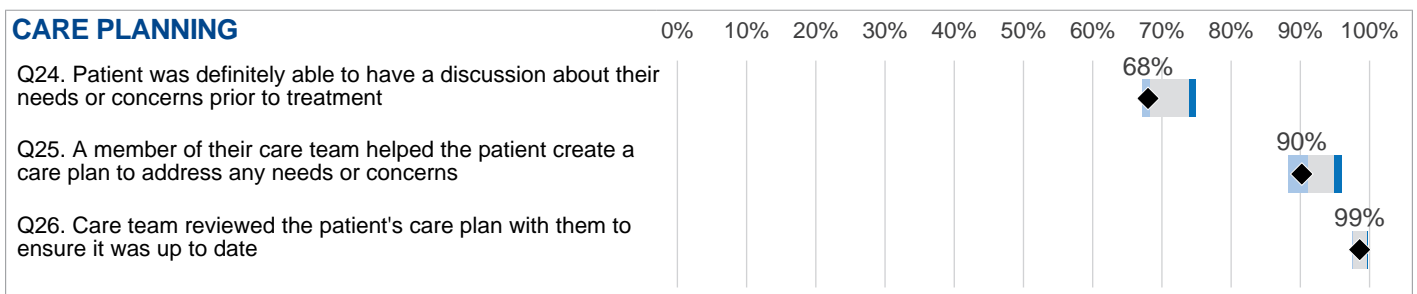
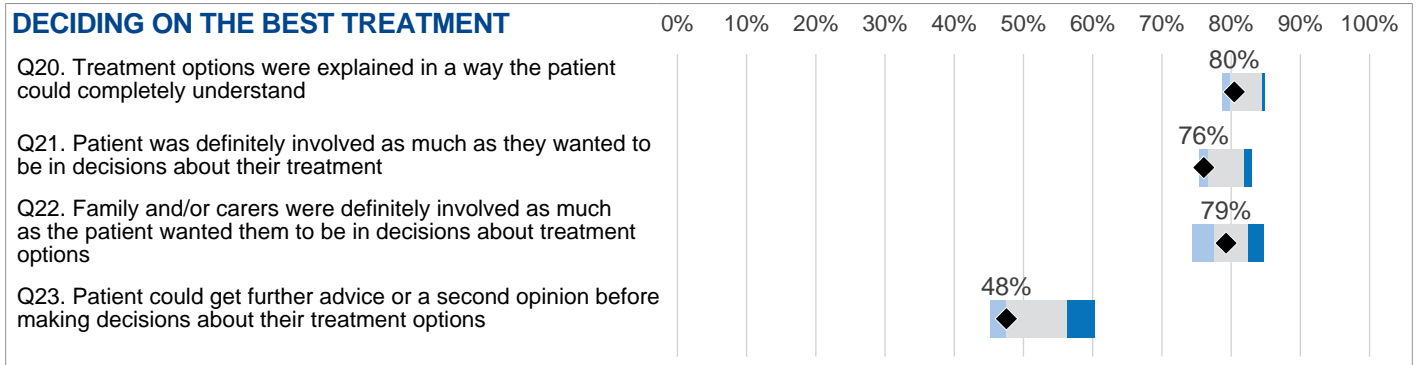
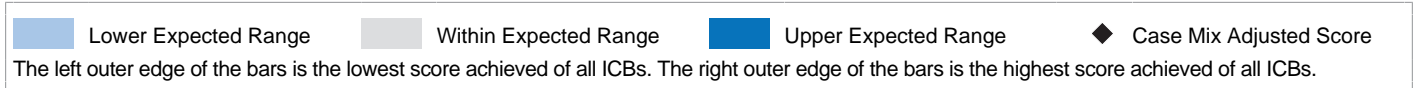
	Number of Respondents
<b>White</b>	
English / Welsh / Scottish / Northern Irish / British	1,256
Irish	19
Gypsy or Irish Traveller	*
Any other White background	49
<b>Mixed / Multiple Ethnicity</b>	
White and Black Caribbean	*
White and Black African	*
White and Asian	*
Any other Mixed / multiple ethnic background	*
<b>Asian or Asian British</b>	
Indian	17
Pakistani	*
Bangladeshi	*
Chinese	*
Any other Asian background	7
<b>Black / African / Caribbean / Black British</b>	
African	14
Caribbean	11
Any other Black / African / Caribbean background	*
<b>Other Ethnicity</b>	
Arab	*
Any other ethnic group	*
<b>Not given</b>	
Not given	112
<b>Total</b>	<b>1,516</b>

## Expected Range Charts

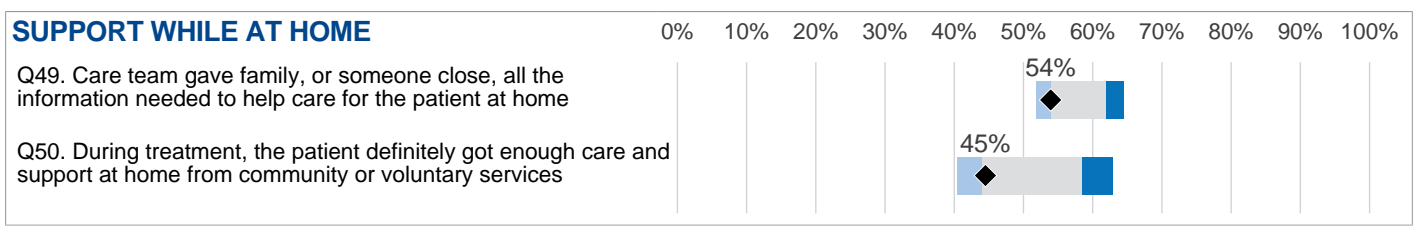
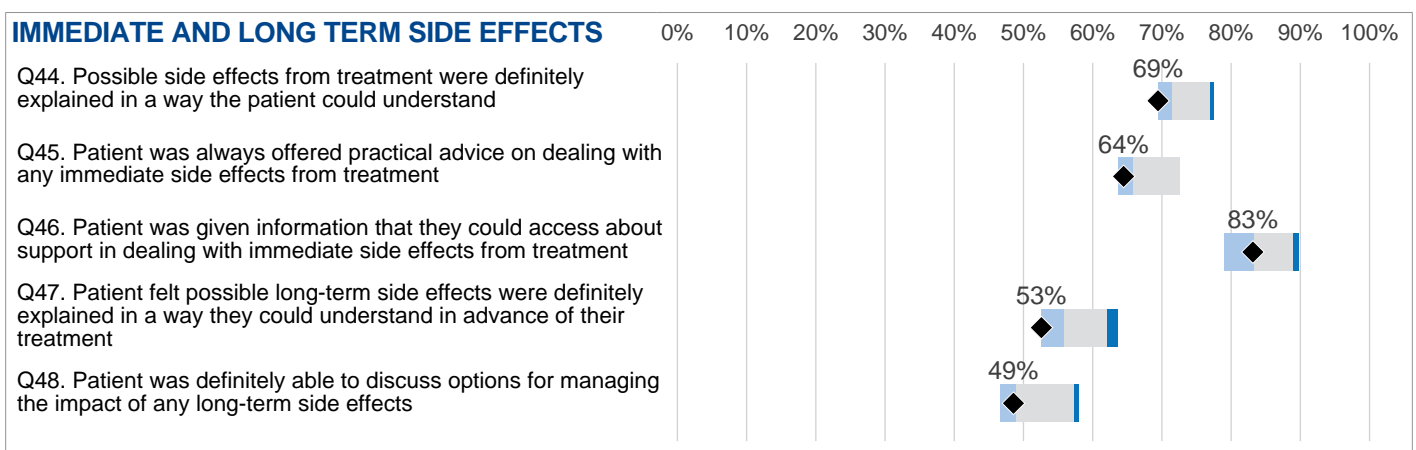
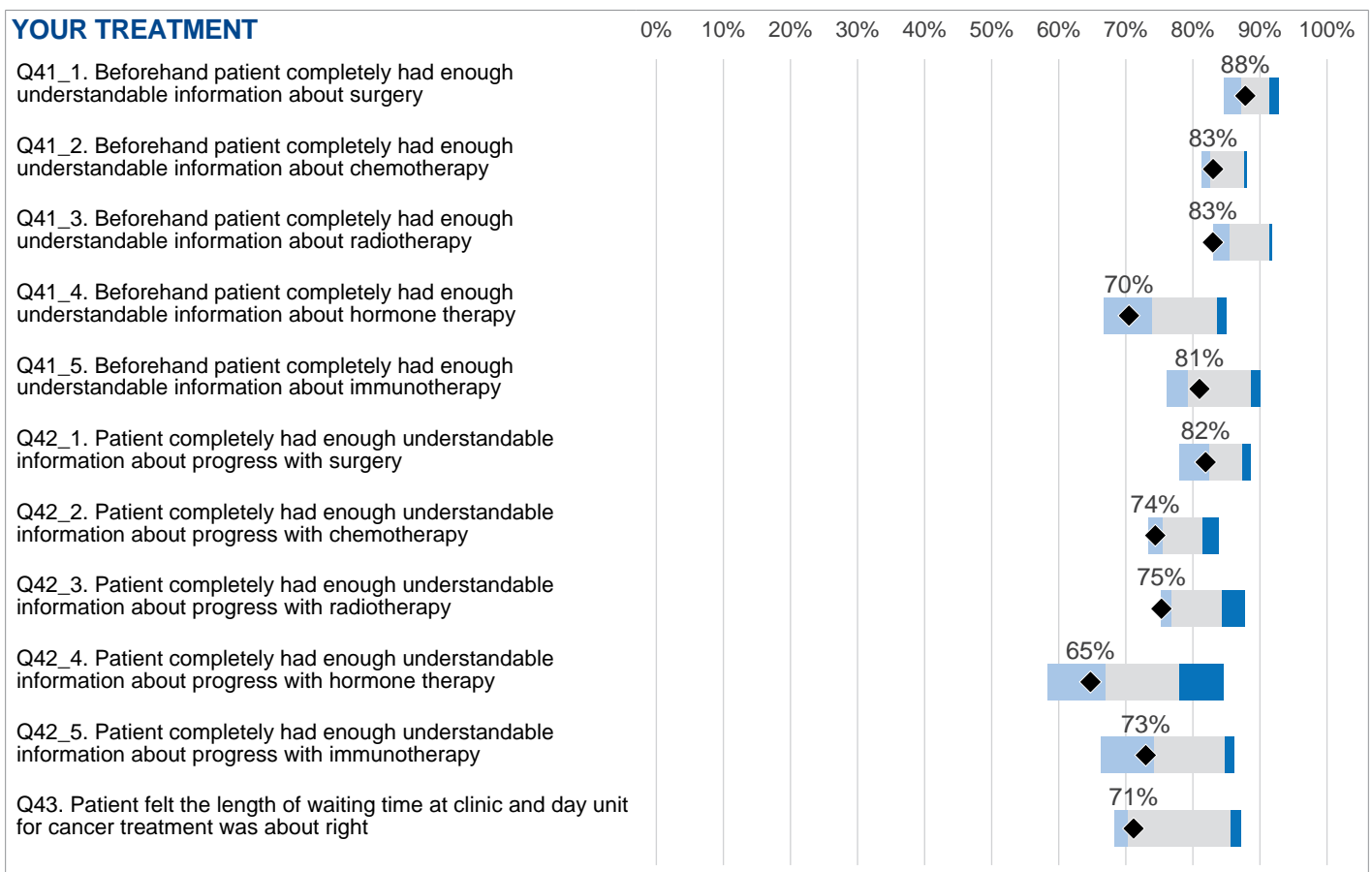
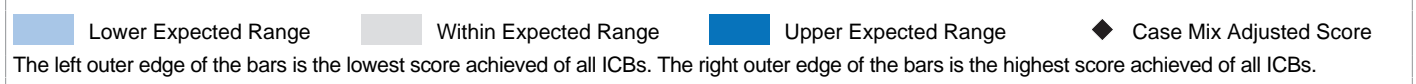




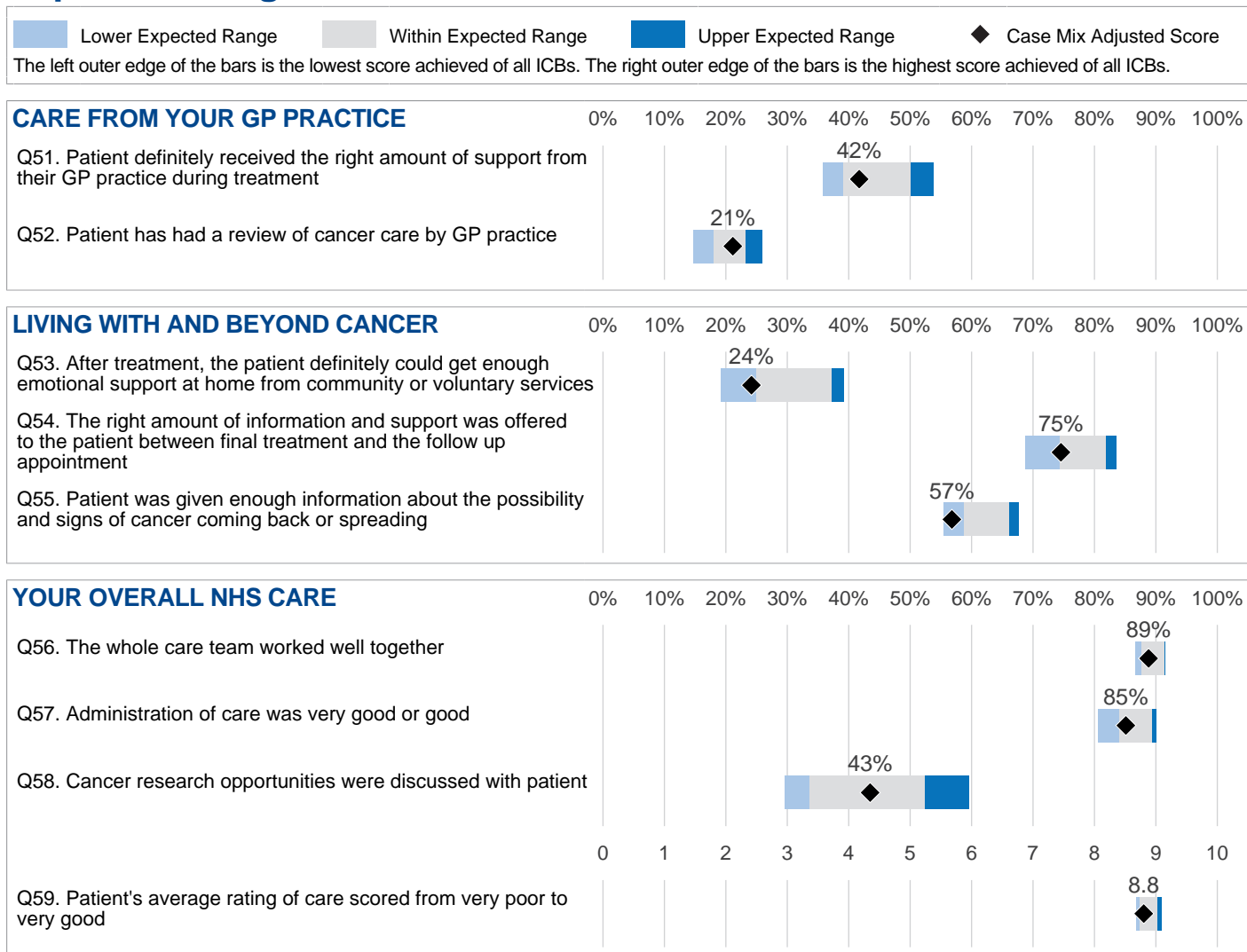
## Expected Range Charts



## Expected Range Charts



## Expected Range Charts



## Comparability tables

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

\*\* No score available for 2021.

▲ or ▼

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

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

SUPPORT FROM YOUR GP PRACTICE	Unadjusted Scores					Case Mix Adjusted Scores			England 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	700	80%	722	78%		77%	75%	81%	78%
Q3. Referral for diagnosis was explained in a way the patient could completely understand	942	64%	956	63%		62%	61%	69%	65%

DIAGNOSTIC TESTS	Unadjusted Scores					Case Mix Adjusted Scores			England 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	1120	93%	1141	91%		91%	90%	94%	92%
Q6. Diagnostic test staff appeared to completely have all the information they needed about the patient	1190	83%	1210	80%		80%	81%	86%	83%
Q7. Patient felt the length of time waiting for diagnostic test results was about right	1190	83%	1205	80%		80%	75%	81%	78%
Q8. Diagnostic test results were explained in a way the patient could completely understand	1196	76%	1211	76%		76%	76%	81%	78%
Q9. Enough privacy was always given to the patient when receiving diagnostic test results	1186	95%	1215	94%		94%	93%	96%	95%

FINDING OUT THAT YOU HAD CANCER	Unadjusted Scores					Case Mix Adjusted Scores			England 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	1356	68%	1387	71%		72%	72%	80%	76%
Q13. Patient was definitely told sensitively that they had cancer	1435	71%	1488	69%		70%	70%	77%	74%
Q14. Cancer diagnosis explained in a way the patient could completely understand	1433	75%	1495	73%		73%	74%	79%	76%
Q15. Patient was definitely told about their diagnosis in an appropriate place	1428	82%	1487	83%		84%	83%	87%	85%
Q16. Patient was told they could go back later for more information about their diagnosis	1253	81%	1320	82%		82%	81%	86%	84%

SUPPORT FROM A MAIN CONTACT PERSON	Unadjusted Scores					Case Mix Adjusted Scores			England 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	1385	92%	1440	92%		92%	89%	94%	92%
Q18. Patient found it very or quite easy to contact their main contact person	1196	82%	1243	80%		80%	80%	87%	84%
Q19. Patient found advice from main contact person was very or quite helpful	1221	95%	1271	95%		95%	94%	97%	95%

## Comparability tables

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

\*\* No score available for 2021.

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

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

<b>DECIDING ON THE BEST TREATMENT</b>	Unadjusted Scores					Case Mix Adjusted Scores			England 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	1328	<b>81%</b>	1383	<b>80%</b>		<b>80%</b>	80%	84%	<b>82%</b>
Q21. Patient was definitely involved as much as they wanted to be in decisions about their treatment	1420	<b>76%</b>	1477	<b>76%</b>		<b>76%</b>	77%	82%	<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	1176	<b>74%</b>	1229	<b>79%</b>	▲	<b>79%</b>	78%	82%	<b>80%</b>
Q23. Patient could get further advice or a second opinion before making decisions about their treatment options	639	<b>46%</b>	684	<b>46%</b>		<b>48%</b>	47%	56%	<b>52%</b>

<b>CARE PLANNING</b>	Unadjusted Scores					Case Mix Adjusted Scores			England 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	1275	<b>67%</b>	1350	<b>68%</b>		<b>68%</b>	68%	74%	<b>71%</b>
Q25. A member of their care team helped the patient create a care plan to address any needs or concerns	726	<b>90%</b>	741	<b>90%</b>		<b>90%</b>	91%	95%	<b>93%</b>
Q26. Care team reviewed the patient's care plan with them to ensure it was up to date	545	<b>98%</b>	539	<b>98%</b>		<b>99%</b>	98%	100%	<b>99%</b>

<b>SUPPORT FROM HOSPITAL STAFF</b>	Unadjusted Scores					Case Mix Adjusted Scores			England 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	1145	<b>87%</b>	1227	<b>89%</b>		<b>89%</b>	88%	92%	<b>90%</b>
Q28. Patient definitely got the right level of support for their overall health and well being from hospital staff	1427	<b>70%</b>	1489	<b>73%</b>		<b>74%</b>	72%	79%	<b>75%</b>
Q29. Patient was offered information about how to get financial help or benefits	681	<b>66%</b>	721	<b>66%</b>		<b>66%</b>	62%	73%	<b>67%</b>

<b>HOSPITAL CARE</b>	Unadjusted Scores					Case Mix Adjusted Scores			England 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	638	<b>76%</b>	656	<b>79%</b>		<b>80%</b>	75%	83%	<b>79%</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	496	<b>55%</b>	510	<b>61%</b>		<b>62%</b>	62%	70%	<b>66%</b>
Q33. Patient was always involved in decisions about their care and treatment whilst in hospital	631	<b>67%</b>	640	<b>67%</b>		<b>67%</b>	66%	73%	<b>69%</b>
Q34. Patient was always able to get help from ward staff when needed	625	<b>71%</b>	640	<b>70%</b>		<b>71%</b>	68%	78%	<b>73%</b>
Q35. Patient was always able to discuss worries and fears with hospital staff	610	<b>60%</b>	614	<b>61%</b>		<b>61%</b>	60%	68%	<b>64%</b>
Q36. Hospital staff always did everything they could to help the patient control pain	560	<b>85%</b>	558	<b>80%</b>		<b>81%</b>	81%	88%	<b>84%</b>
Q37. Patient was always treated with respect and dignity while in hospital	638	<b>87%</b>	651	<b>86%</b>		<b>86%</b>	85%	91%	<b>88%</b>
Q38. Patient received easily understandable information about what they should or should not do after leaving hospital	627	<b>86%</b>	639	<b>87%</b>		<b>87%</b>	85%	90%	<b>88%</b>
Q39. Patient was always able to discuss worries and fears with hospital staff while being treated as an outpatient or day case	1236	<b>73%</b>	1289	<b>74%</b>		<b>75%</b>	75%	82%	<b>78%</b>

## Comparability tables

<p>* Indicates where a score is not available due to suppression or a low base size.</p> <p>** No score available for 2021.</p>	<p>▲ or ▼ Change 2021-2022: Indicates where 2022 score is significantly higher or lower than 2021 score.</p>	<p><span style="display: inline-block; width: 15px; height: 10px; background-color: #d9e1f2; border: 1px solid #000;"></span> Adjusted Score below Lower Expected Range</p> <p><span style="display: inline-block; width: 15px; height: 10px; background-color: #e6e6e6; border: 1px solid #000;"></span> Adjusted Score between Upper and Lower Expected Ranges</p> <p><span style="display: inline-block; width: 15px; height: 10px; background-color: #0070c0; border: 1px solid #000;"></span> Adjusted Score above Upper Expected Range</p>
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YOUR TREATMENT	Unadjusted Scores					Case Mix Adjusted Scores			England 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	809	<b>88%</b>	821	<b>87%</b>		<b>88%</b>	87%	91%	<b>89%</b>
Q41_2. Beforehand patient completely had enough understandable information about chemotherapy	785	<b>83%</b>	780	<b>82%</b>		<b>83%</b>	83%	88%	<b>85%</b>
Q41_3. Beforehand patient completely had enough understandable information about radiotherapy	454	<b>84%</b>	456	<b>82%</b>		<b>83%</b>	86%	91%	<b>88%</b>
Q41_4. Beforehand patient completely had enough understandable information about hormone therapy	238	<b>71%</b>	267	<b>69%</b>		<b>70%</b>	74%	84%	<b>79%</b>
Q41_5. Beforehand patient completely had enough understandable information about immunotherapy	240	<b>87%</b>	238	<b>80%</b>		<b>81%</b>	79%	89%	<b>84%</b>
Q42_1. Patient completely had enough understandable information about progress with surgery	804	<b>81%</b>	813	<b>82%</b>		<b>82%</b>	82%	87%	<b>85%</b>
Q42_2. Patient completely had enough understandable information about progress with chemotherapy	779	<b>75%</b>	769	<b>74%</b>		<b>74%</b>	76%	81%	<b>79%</b>
Q42_3. Patient completely had enough understandable information about progress with radiotherapy	447	<b>76%</b>	439	<b>75%</b>		<b>75%</b>	77%	84%	<b>81%</b>
Q42_4. Patient completely had enough understandable information about progress with hormone therapy	236	<b>64%</b>	254	<b>63%</b>		<b>65%</b>	67%	78%	<b>72%</b>
Q42_5. Patient completely had enough understandable information about progress with immunotherapy	240	<b>76%</b>	230	<b>72%</b>		<b>73%</b>	74%	85%	<b>79%</b>
Q43. Patient felt the length of waiting time at clinic and day unit for cancer treatment was about right	1406	<b>71%</b>	1465	<b>70%</b>		<b>71%</b>	70%	86%	<b>78%</b>

IMMEDIATE AND LONG TERM SIDE EFFECTS	Unadjusted Scores					Case Mix Adjusted Scores			England 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	1387	<b>71%</b>	1418	<b>69%</b>		<b>69%</b>	71%	77%	<b>74%</b>
Q45. Patient was always offered practical advice on dealing with any immediate side effects from treatment	1317	<b>67%</b>	1380	<b>64%</b>		<b>64%</b>	66%	73%	<b>69%</b>
Q46. Patient was given information that they could access about support in dealing with immediate side effects from treatment	1041	<b>85%</b>	1090	<b>83%</b>		<b>83%</b>	83%	89%	<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	1299	<b>54%</b>	1318	<b>52%</b>		<b>53%</b>	56%	62%	<b>59%</b>
Q48. Patient was definitely able to discuss options for managing the impact of any long-term side effects	1108	<b>47%</b>	1132	<b>47%</b>		<b>49%</b>	49%	57%	<b>53%</b>

SUPPORT WHILE AT HOME	Unadjusted Scores					Case Mix Adjusted Scores			England 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	945	<b>50%</b>	983	<b>53%</b>		<b>54%</b>	54%	62%	<b>58%</b>
Q50. During treatment, the patient definitely got enough care and support at home from community or voluntary services	515	<b>43%</b>	566	<b>44%</b>		<b>45%</b>	44%	59%	<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			England 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	756	<b>38%</b>	808	<b>42%</b>		<b>42%</b>	39%	50%	<b>45%</b>
Q52. Patient has had a review of cancer care by GP practice	1387	<b>16%</b>	1420	<b>21%</b>	▲	<b>21%</b>	18%	23%	<b>21%</b>

<b>LIVING WITH AND BEYOND CANCER</b>	Unadjusted Scores					Case Mix Adjusted Scores			England 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	301	<b>26%</b>	324	<b>24%</b>		<b>24%</b>	25%	37%	<b>31%</b>
Q54. The right amount of information and support was offered to the patient between final treatment and the follow up appointment	695	<b>76%</b>	699	<b>74%</b>		<b>75%</b>	74%	82%	<b>78%</b>
Q55. Patient was given enough information about the possibility and signs of cancer coming back or spreading	1113	<b>57%</b>	1186	<b>56%</b>		<b>57%</b>	59%	66%	<b>62%</b>

<b>YOUR OVERALL NHS CARE</b>	Unadjusted Scores					Case Mix Adjusted Scores			England 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	1374	<b>88%</b>	1424	<b>88%</b>		<b>89%</b>	88%	91%	<b>90%</b>
Q57. Administration of care was very good or good	1426	<b>84%</b>	1479	<b>85%</b>		<b>85%</b>	84%	89%	<b>87%</b>
Q58. Cancer research opportunities were discussed with patient	882	<b>43%</b>	937	<b>43%</b>		<b>43%</b>	34%	52%	<b>43%</b>
Q59. Patient's average rating of care scored from very poor to very good	1396	<b>8.8</b>	1450	<b>8.8</b>		<b>8.8</b>	8.7	9.0	<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	*	95%	84%	67%	53%	83%	77%	78%	*	84%	72%	84%	70%	<b>78%</b>
Q3. Referral for diagnosis was explained in a way the patient could completely understand	*	75%	70%	59%	46%	58%	54%	70%	*	61%	49%	63%	64%	<b>63%</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	*	89%	94%	85%	87%	84%	93%	89%	*	100%	91%	96%	95%	<b>91%</b>
Q6. Diagnostic test staff appeared to completely have all the information they needed about the patient	*	82%	86%	74%	73%	80%	84%	81%	*	93%	75%	84%	80%	<b>80%</b>
Q7. Patient felt the length of time waiting for diagnostic test results was about right	*	78%	81%	80%	84%	77%	79%	78%	*	78%	80%	83%	78%	<b>80%</b>
Q8. Diagnostic test results were explained in a way the patient could completely understand	*	76%	84%	68%	71%	73%	76%	73%	*	88%	70%	81%	76%	<b>76%</b>
Q9. Enough privacy was always given to the patient when receiving diagnostic test results	*	97%	92%	88%	94%	93%	92%	94%	*	100%	94%	91%	92%	<b>94%</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	*	75%	85%	63%	62%	60%	75%	71%	*	77%	78%	63%	68%	<b>71%</b>
Q13. Patient was definitely told sensitively that they had cancer	60%	75%	79%	64%	65%	63%	64%	63%	*	75%	73%	62%	64%	<b>69%</b>
Q14. Cancer diagnosis explained in a way the patient could completely understand	70%	78%	87%	71%	63%	75%	67%	67%	*	83%	72%	73%	68%	<b>73%</b>
Q15. Patient was definitely told about their diagnosis in an appropriate place	90%	89%	89%	69%	78%	84%	79%	86%	*	85%	85%	84%	77%	<b>83%</b>
Q16. Patient was told they could go back later for more information about their diagnosis	80%	88%	81%	82%	76%	84%	69%	81%	*	91%	83%	80%	79%	<b>82%</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
Q17. Patient had a main point of contact within the care team	*	94%	97%	95%	91%	87%	94%	87%	*	89%	88%	87%	92%	<b>92%</b>
Q18. Patient found it very or quite easy to contact their main contact person	*	80%	85%	83%	79%	83%	92%	64%	*	82%	81%	82%	79%	<b>80%</b>
Q19. Patient found advice from main contact person was very or quite helpful	*	94%	96%	90%	94%	97%	97%	94%	*	95%	97%	99%	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
Q20. Treatment options were explained in a way the patient could completely understand	60%	80%	85%	80%	73%	84%	85%	76%	*	93%	80%	86%	81%	<b>80%</b>
Q21. Patient was definitely involved as much as they wanted to be in decisions about their treatment	60%	76%	76%	76%	71%	69%	80%	75%	*	87%	84%	81%	72%	<b>76%</b>
Q22. Family and/or carers were definitely involved as much as the patient wanted them to be in decisions about treatment options	70%	77%	85%	81%	77%	80%	87%	73%	*	80%	84%	83%	77%	<b>79%</b>
Q23. Patient could get further advice or a second opinion before making decisions about their treatment options	*	41%	54%	37%	43%	33%	41%	56%	*	47%	62%	48%	45%	<b>46%</b>

	Tumour Type													
	Brain / CNS	Breast	Colorectal / LGT	Gynaecological	Haematological	Head and Neck	Lung	Prostate	Sarcoma	Skin	Upper Gastro	Urological	Other	All Cancers
Q24. Patient was definitely able to have a discussion about their needs or concerns prior to treatment	50%	68%	76%	64%	60%	63%	67%	76%	*	80%	67%	69%	63%	<b>68%</b>
Q25. A member of their care team helped the patient create a care plan to address any needs or concerns	*	87%	93%	88%	86%	83%	100%	94%	*	94%	92%	94%	91%	<b>90%</b>
Q26. Care team reviewed the patient's care plan with them to ensure it was up to date	*	99%	99%	100%	97%	100%	100%	98%	*	100%	100%	100%	95%	<b>98%</b>

	Tumour Type													
	Brain / CNS	Breast	Colorectal / LGT	Gynaecological	Haematological	Head and Neck	Lung	Prostate	Sarcoma	Skin	Upper Gastro	Urological	Other	All Cancers
Q27. Staff provided the patient with relevant information on available support	*	90%	94%	91%	88%	93%	96%	89%	*	90%	95%	88%	81%	<b>89%</b>
Q28. Patient definitely got the right level of support for their overall health and well being from hospital staff	60%	70%	75%	69%	73%	73%	78%	71%	*	72%	77%	78%	70%	<b>73%</b>
Q29. Patient was offered information about how to get financial help or benefits	*	71%	71%	61%	68%	83%	85%	48%	*	40%	66%	54%	65%	<b>66%</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
<b>HOSPITAL CARE</b>														
Q31. Patient had confidence and trust in all of the team looking after them during their stay in hospital	*	78%	80%	75%	80%	88%	86%	79%	*	*	77%	84%	69%	<b>79%</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%	68%	61%	65%	70%	67%	55%	*	*	64%	68%	53%	<b>61%</b>
Q33. Patient was always involved in decisions about their care and treatment whilst in hospital	*	71%	71%	58%	64%	71%	64%	66%	*	*	67%	71%	66%	<b>67%</b>
Q34. Patient was always able to get help from ward staff when needed	*	70%	71%	58%	72%	77%	86%	79%	*	*	73%	71%	60%	<b>70%</b>
Q35. Patient was always able to discuss worries and fears with hospital staff	*	64%	57%	54%	54%	75%	62%	67%	*	*	59%	57%	61%	<b>61%</b>
Q36. Hospital staff always did everything they could to help the patient control pain	*	78%	83%	80%	87%	70%	89%	79%	*	*	80%	77%	72%	<b>80%</b>
Q37. Patient was always treated with respect and dignity while in hospital	*	82%	88%	85%	85%	90%	86%	88%	*	*	87%	85%	85%	<b>86%</b>
Q38. Patient received easily understandable information about what they should or should not do after leaving hospital	*	82%	89%	85%	85%	97%	79%	93%	*	*	83%	91%	85%	<b>87%</b>
Q39. Patient was always able to discuss worries and fears with hospital staff while being treated as an outpatient or day case	*	70%	83%	68%	78%	79%	66%	75%	*	85%	83%	78%	69%	<b>74%</b>
<b>YOUR TREATMENT</b>														
Q41_1. Beforehand patient completely had enough understandable information about surgery	*	88%	90%	89%	75%	76%	79%	88%	*	94%	90%	88%	88%	<b>87%</b>
Q41_2. Beforehand patient completely had enough understandable information about chemotherapy	*	83%	85%	84%	79%	78%	84%	80%	*	*	93%	83%	78%	<b>82%</b>
Q41_3. Beforehand patient completely had enough understandable information about radiotherapy	*	84%	74%	92%	77%	81%	70%	75%	*	*	91%	100%	79%	<b>82%</b>
Q41_4. Beforehand patient completely had enough understandable information about hormone therapy	*	69%	*	*	*	*	*	76%	*	*	*	*	56%	<b>69%</b>
Q41_5. Beforehand patient completely had enough understandable information about immunotherapy	*	70%	90%	83%	81%	*	70%	*	*	92%	*	93%	76%	<b>80%</b>
Q42_1. Patient completely had enough understandable information about progress with surgery	*	86%	82%	76%	71%	68%	83%	76%	*	91%	78%	83%	82%	<b>82%</b>
Q42_2. Patient completely had enough understandable information about progress with chemotherapy	*	78%	73%	79%	71%	78%	75%	65%	*	*	78%	76%	72%	<b>74%</b>
Q42_3. Patient completely had enough understandable information about progress with radiotherapy	*	78%	73%	92%	70%	78%	74%	63%	*	*	82%	*	69%	<b>75%</b>
Q42_4. Patient completely had enough understandable information about progress with hormone therapy	*	67%	*	*	*	*	*	56%	*	*	*	*	64%	<b>63%</b>
Q42_5. Patient completely had enough understandable information about progress with immunotherapy	*	68%	*	83%	76%	*	64%	*	*	96%	*	72%	59%	<b>72%</b>
Q43. Patient felt the length of waiting time at clinic and day unit for cancer treatment was about right	80%	66%	78%	62%	59%	90%	79%	79%	*	80%	76%	76%	64%	<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
Q44. Possible side effects from treatment were definitely explained in a way the patient could understand	*	69%	70%	68%	61%	65%	74%	61%	*	80%	77%	78%	72%	<b>69%</b>
Q45. Patient was always offered practical advice on dealing with any immediate side effects from treatment	80%	64%	70%	64%	61%	59%	68%	50%	*	81%	66%	70%	64%	<b>64%</b>
Q46. Patient was given information that they could access about support in dealing with immediate side effects from treatment	90%	86%	85%	88%	80%	85%	86%	77%	*	96%	89%	84%	77%	<b>83%</b>
Q47. Patient felt possible long-term side effects were definitely explained in a way they could understand in advance of their treatment	*	52%	57%	45%	42%	46%	53%	62%	*	65%	53%	57%	47%	<b>52%</b>
Q48. Patient was definitely able to discuss options for managing the impact of any long-term side effects	*	45%	51%	46%	42%	49%	46%	52%	*	57%	47%	59%	46%	<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
Q49. Care team gave family, or someone close, all the information needed to help care for the patient at home	*	48%	57%	49%	56%	63%	62%	53%	*	55%	50%	56%	50%	<b>53%</b>
Q50. During treatment, the patient definitely got enough care and support at home from community or voluntary services	*	44%	42%	53%	42%	50%	43%	41%	*	50%	39%	45%	43%	<b>44%</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	*	42%	51%	52%	32%	32%	43%	46%	*	59%	28%	40%	39%	<b>42%</b>
Q52. Patient has had a review of cancer care by GP practice	*	21%	27%	25%	20%	14%	18%	21%	*	22%	26%	14%	20%	<b>21%</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	*	24%	33%	30%	21%	14%	*	16%	*	*	29%	18%	33%	<b>24%</b>
Q54. The right amount of information and support was offered to the patient between final treatment and the follow up appointment	*	69%	78%	73%	81%	80%	86%	75%	*	79%	86%	75%	69%	<b>74%</b>
Q55. Patient was given enough information about the possibility and signs of cancer coming back or spreading	*	53%	57%	48%	60%	40%	50%	50%	*	80%	51%	74%	57%	<b>56%</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	*	90%	90%	88%	88%	85%	89%	86%	*	90%	86%	91%	88%	<b>88%</b>
Q57. Administration of care was very good or good	*	87%	83%	86%	86%	84%	91%	80%	*	83%	86%	86%	81%	<b>85%</b>
Q58. Cancer research opportunities were discussed with patient	*	33%	36%	59%	50%	29%	55%	37%	*	26%	61%	43%	53%	<b>43%</b>
Q59. Patient's average rating of care scored from very poor to very good	*	8.9	8.9	8.7	8.7	8.8	8.8	8.6	*	8.9	8.9	8.9	8.6	<b>8.8</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	*	*	90%	74%	74%	80%	79%	87%	<b>78%</b>
Q3. Referral for diagnosis was explained in a way the patient could completely understand	*	*	67%	76%	64%	62%	63%	47%	<b>63%</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	*	*	91%	87%	90%	91%	91%	90%	<b>91%</b>
Q6. Diagnostic test staff appeared to completely have all the information they needed about the patient	*	70%	74%	75%	81%	79%	84%	78%	<b>80%</b>
Q7. Patient felt the length of time waiting for diagnostic test results was about right	*	80%	74%	69%	77%	82%	84%	78%	<b>80%</b>
Q8. Diagnostic test results were explained in a way the patient could completely understand	*	90%	68%	69%	76%	79%	76%	65%	<b>76%</b>
Q9. Enough privacy was always given to the patient when receiving diagnostic test results	*	90%	85%	95%	93%	96%	92%	93%	<b>94%</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	*	58%	70%	66%	65%	74%	75%	71%	<b>71%</b>
Q13. Patient was definitely told sensitively that they had cancer	*	58%	73%	67%	64%	71%	71%	76%	<b>69%</b>
Q14. Cancer diagnosis explained in a way the patient could completely understand	*	58%	68%	68%	73%	77%	72%	74%	<b>73%</b>
Q15. Patient was definitely told about their diagnosis in an appropriate place	*	42%	82%	76%	81%	86%	87%	86%	<b>83%</b>
Q16. Patient was told they could go back later for more information about their diagnosis	*	92%	81%	84%	84%	83%	81%	70%	<b>82%</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	*	100%	93%	92%	91%	93%	93%	87%	<b>92%</b>
Q18. Patient found it very or quite easy to contact their main contact person	*	82%	76%	82%	78%	81%	80%	80%	<b>80%</b>
Q19. Patient found advice from main contact person was very or quite helpful	*	100%	95%	94%	93%	96%	96%	96%	<b>95%</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	*	*	71%	76%	80%	82%	82%	78%	<b>80%</b>
Q21. Patient was definitely involved as much as they wanted to be in decisions about their treatment	*	58%	68%	69%	75%	77%	78%	76%	<b>76%</b>
Q22. Family and/or carers were definitely involved as much as the patient wanted them to be in decisions about treatment options	*	*	65%	74%	75%	84%	80%	80%	<b>79%</b>
Q23. Patient could get further advice or a second opinion before making decisions about their treatment options	*	45%	38%	37%	46%	48%	48%	45%	<b>46%</b>

## Age group tables

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<b>CARE PLANNING</b>	Age								
	16 - 24	25 - 34	35 - 44	45 - 54	55 - 64	65 - 74	75 - 84	85+	All
Q24. Patient was definitely able to have a discussion about their needs or concerns prior to treatment	*	58%	63%	61%	69%	69%	68%	68%	<b>68%</b>
Q25. A member of their care team helped the patient create a care plan to address any needs or concerns	*	*	93%	84%	87%	92%	91%	100%	<b>90%</b>
Q26. Care team reviewed the patient's care plan with them to ensure it was up to date	*	*	94%	98%	97%	99%	98%	100%	<b>98%</b>

<b>SUPPORT FROM HOSPITAL STAFF</b>	Age								
	16 - 24	25 - 34	35 - 44	45 - 54	55 - 64	65 - 74	75 - 84	85+	All
Q27. Staff provided the patient with relevant information on available support	*	91%	89%	87%	89%	91%	89%	89%	<b>89%</b>
Q28. Patient definitely got the right level of support for their overall health and well being from hospital staff	*	75%	63%	60%	71%	76%	75%	69%	<b>73%</b>
Q29. Patient was offered information about how to get financial help or benefits	*	*	82%	77%	63%	69%	60%	42%	<b>66%</b>

<b>HOSPITAL CARE</b>	Age								
	16 - 24	25 - 34	35 - 44	45 - 54	55 - 64	65 - 74	75 - 84	85+	All
Q31. Patient had confidence and trust in all of the team looking after them during their stay in hospital	*	*	65%	74%	77%	83%	80%	86%	<b>79%</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	*	*	48%	57%	56%	66%	64%	64%	<b>61%</b>
Q33. Patient was always involved in decisions about their care and treatment whilst in hospital	*	*	61%	64%	68%	70%	62%	73%	<b>67%</b>
Q34. Patient was always able to get help from ward staff when needed	*	*	52%	65%	69%	76%	72%	69%	<b>70%</b>
Q35. Patient was always able to discuss worries and fears with hospital staff	*	*	43%	58%	63%	69%	54%	44%	<b>61%</b>
Q36. Hospital staff always did everything they could to help the patient control pain	*	*	73%	74%	79%	82%	87%	65%	<b>80%</b>
Q37. Patient was always treated with respect and dignity while in hospital	*	*	74%	83%	86%	88%	86%	86%	<b>86%</b>
Q38. Patient received easily understandable information about what they should or should not do after leaving hospital	*	*	73%	87%	89%	89%	84%	76%	<b>87%</b>
Q39. Patient was always able to discuss worries and fears with hospital staff while being treated as an outpatient or day case	*	64%	75%	70%	73%	77%	75%	69%	<b>74%</b>

## Age group tables

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

YOUR TREATMENT	Age								
	16 - 24	25 - 34	35 - 44	45 - 54	55 - 64	65 - 74	75 - 84	85+	All
Q41_1. Beforehand patient completely had enough understandable information about surgery	*	*	81%	82%	85%	90%	90%	80%	<b>87%</b>
Q41_2. Beforehand patient completely had enough understandable information about chemotherapy	*	*	83%	73%	81%	87%	80%	83%	<b>82%</b>
Q41_3. Beforehand patient completely had enough understandable information about radiotherapy	*	*	80%	79%	82%	84%	82%	80%	<b>82%</b>
Q41_4. Beforehand patient completely had enough understandable information about hormone therapy	*	*	60%	59%	71%	75%	72%	*	<b>69%</b>
Q41_5. Beforehand patient completely had enough understandable information about immunotherapy	*	*	*	80%	83%	81%	73%	*	<b>80%</b>
Q42_1. Patient completely had enough understandable information about progress with surgery	*	*	70%	77%	81%	85%	81%	74%	<b>82%</b>
Q42_2. Patient completely had enough understandable information about progress with chemotherapy	*	*	66%	72%	76%	78%	69%	68%	<b>74%</b>
Q42_3. Patient completely had enough understandable information about progress with radiotherapy	*	*	79%	79%	73%	78%	76%	53%	<b>75%</b>
Q42_4. Patient completely had enough understandable information about progress with hormone therapy	*	*	*	66%	68%	62%	60%	*	<b>63%</b>
Q42_5. Patient completely had enough understandable information about progress with immunotherapy	*	*	*	68%	75%	72%	65%	*	<b>72%</b>
Q43. Patient felt the length of waiting time at clinic and day unit for cancer treatment was about right	*	58%	60%	70%	69%	71%	74%	67%	<b>70%</b>

IMMEDIATE AND LONG TERM SIDE EFFECTS	Age								
	16 - 24	25 - 34	35 - 44	45 - 54	55 - 64	65 - 74	75 - 84	85+	All
Q44. Possible side effects from treatment were definitely explained in a way the patient could understand	*	83%	69%	72%	72%	71%	62%	67%	<b>69%</b>
Q45. Patient was always offered practical advice on dealing with any immediate side effects from treatment	*	73%	57%	62%	66%	66%	61%	62%	<b>64%</b>
Q46. Patient was given information that they could access about support in dealing with immediate side effects from treatment	*	*	79%	88%	83%	84%	81%	81%	<b>83%</b>
Q47. Patient felt possible long-term side effects were definitely explained in a way they could understand in advance of their treatment	*	73%	50%	51%	55%	53%	48%	43%	<b>52%</b>
Q48. Patient was definitely able to discuss options for managing the impact of any long-term side effects	*	*	41%	40%	51%	48%	48%	44%	<b>47%</b>

SUPPORT WHILE AT HOME	Age								
	16 - 24	25 - 34	35 - 44	45 - 54	55 - 64	65 - 74	75 - 84	85+	All
Q49. Care team gave family, or someone close, all the information needed to help care for the patient at home	*	*	42%	44%	51%	55%	54%	63%	<b>53%</b>
Q50. During treatment, the patient definitely got enough care and support at home from community or voluntary services	*	*	32%	38%	39%	46%	50%	48%	<b>44%</b>

CARE FROM YOUR GP PRACTICE	Age								
	16 - 24	25 - 34	35 - 44	45 - 54	55 - 64	65 - 74	75 - 84	85+	All
Q51. Patient definitely received the right amount of support from their GP practice during treatment	*	*	61%	36%	40%	44%	42%	40%	<b>42%</b>
Q52. Patient has had a review of cancer care by GP practice	*	*	33%	18%	22%	21%	20%	23%	<b>21%</b>

## Age group tables

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

<b>LIVING WITH AND BEYOND CANCER</b>	Age								
	16 - 24	25 - 34	35 - 44	45 - 54	55 - 64	65 - 74	75 - 84	85+	All
Q53. After treatment, the patient definitely could get enough emotional support at home from community or voluntary services	*	*	46%	23%	17%	21%	31%	25%	<b>24%</b>
Q54. The right amount of information and support was offered to the patient between final treatment and the follow up appointment	*	*	70%	74%	73%	76%	75%	79%	<b>74%</b>
Q55. Patient was given enough information about the possibility and signs of cancer coming back or spreading	*	45%	43%	48%	56%	61%	53%	63%	<b>56%</b>

<b>YOUR OVERALL NHS CARE</b>	Age								
	16 - 24	25 - 34	35 - 44	45 - 54	55 - 64	65 - 74	75 - 84	85+	All
Q56. The whole care team worked well together	*	100%	90%	85%	88%	89%	87%	94%	<b>88%</b>
Q57. Administration of care was very good or good	*	100%	77%	86%	83%	85%	86%	85%	<b>85%</b>
Q58. Cancer research opportunities were discussed with patient	*	*	35%	42%	44%	47%	42%	26%	<b>43%</b>
Q59. Patient's average rating of care scored from very poor to very good	*	8.8	8.5	8.5	8.7	8.9	8.8	8.5	<b>8.8</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	82%	70%	*	*	*	84%	<b>78%</b>
Q3. Referral for diagnosis was explained in a way the patient could completely understand	66%	60%	*	*	*	64%	<b>63%</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	89%	92%	*	*	*	95%	<b>91%</b>
Q6. Diagnostic test staff appeared to completely have all the information they needed about the patient	80%	81%	*	*	*	77%	<b>80%</b>
Q7. Patient felt the length of time waiting for diagnostic test results was about right	77%	83%	*	*	*	81%	<b>80%</b>
Q8. Diagnostic test results were explained in a way the patient could completely understand	74%	77%	*	*	*	80%	<b>76%</b>
Q9. Enough privacy was always given to the patient when receiving diagnostic test results	93%	94%	*	*	*	94%	<b>94%</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	71%	70%	*	*	*	74%	<b>71%</b>
Q13. Patient was definitely told sensitively that they had cancer	69%	68%	*	*	*	73%	<b>69%</b>
Q14. Cancer diagnosis explained in a way the patient could completely understand	73%	72%	*	*	*	75%	<b>73%</b>
Q15. Patient was definitely told about their diagnosis in an appropriate place	82%	85%	*	*	*	84%	<b>83%</b>
Q16. Patient was told they could go back later for more information about their diagnosis	83%	81%	*	*	*	84%	<b>82%</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	93%	90%	*	*	*	95%	<b>92%</b>
Q18. Patient found it very or quite easy to contact their main contact person	81%	78%	*	*	*	81%	<b>80%</b>
Q19. Patient found advice from main contact person was very or quite helpful	93%	96%	*	*	*	99%	<b>95%</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	80%	80%	*	*	*	83%	<b>80%</b>
Q21. Patient was definitely involved as much as they wanted to be in decisions about their treatment	75%	77%	*	*	*	75%	<b>76%</b>
Q22. Family and/or carers were definitely involved as much as the patient wanted them to be in decisions about treatment options	79%	80%	*	*	*	74%	<b>79%</b>
Q23. Patient could get further advice or a second opinion before making decisions about their treatment options	44%	49%	*	*	*	47%	<b>46%</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	66%	70%	*	*	*	63%	<b>68%</b>
Q25. A member of their care team helped the patient create a care plan to address any needs or concerns	88%	93%	*	*	*	92%	<b>90%</b>
Q26. Care team reviewed the patient's care plan with them to ensure it was up to date	98%	99%	*	*	*	100%	<b>98%</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	88%	91%	*	*	*	90%	<b>89%</b>
Q28. Patient definitely got the right level of support for their overall health and well being from hospital staff	69%	78%	*	*	*	69%	<b>73%</b>
Q29. Patient was offered information about how to get financial help or benefits	66%	66%	*	*	*	75%	<b>66%</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	75%	83%	*	*	*	79%	<b>79%</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%	65%	*	*	*	61%	<b>61%</b>
Q33. Patient was always involved in decisions about their care and treatment whilst in hospital	65%	69%	*	*	*	62%	<b>67%</b>
Q34. Patient was always able to get help from ward staff when needed	64%	79%	*	*	*	54%	<b>70%</b>
Q35. Patient was always able to discuss worries and fears with hospital staff	57%	66%	*	*	*	46%	<b>61%</b>
Q36. Hospital staff always did everything they could to help the patient control pain	79%	83%	*	*	*	75%	<b>80%</b>
Q37. Patient was always treated with respect and dignity while in hospital	82%	89%	*	*	*	89%	<b>86%</b>
Q38. Patient received easily understandable information about what they should or should not do after leaving hospital	86%	88%	*	*	*	77%	<b>87%</b>
Q39. Patient was always able to discuss worries and fears with hospital staff while being treated as an outpatient or day case	71%	79%	*	*	*	75%	<b>74%</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	87%	87%	*	*	*	94%	<b>87%</b>
Q41_2. Beforehand patient completely had enough understandable information about chemotherapy	80%	85%	*	*	*	82%	<b>82%</b>
Q41_3. Beforehand patient completely had enough understandable information about radiotherapy	85%	76%	*	*	*	86%	<b>82%</b>
Q41_4. Beforehand patient completely had enough understandable information about hormone therapy	66%	73%	*	*	*	83%	<b>69%</b>
Q41_5. Beforehand patient completely had enough understandable information about immunotherapy	76%	87%	*	*	*	73%	<b>80%</b>
Q42_1. Patient completely had enough understandable information about progress with surgery	82%	81%	*	*	*	81%	<b>82%</b>
Q42_2. Patient completely had enough understandable information about progress with chemotherapy	74%	72%	*	*	*	77%	<b>74%</b>
Q42_3. Patient completely had enough understandable information about progress with radiotherapy	78%	69%	*	*	*	78%	<b>75%</b>
Q42_4. Patient completely had enough understandable information about progress with hormone therapy	65%	59%	*	*	*	73%	<b>63%</b>
Q42_5. Patient completely had enough understandable information about progress with immunotherapy	69%	77%	*	*	*	60%	<b>72%</b>
Q43. Patient felt the length of waiting time at clinic and day unit for cancer treatment was about right	64%	78%	*	*	*	70%	<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						
	Female	Male	Non-binary	Prefer to self-describe	Prefer not to say	Not given	All
<b>IMMEDIATE AND LONG TERM SIDE EFFECTS</b>							
Q44. Possible side effects from treatment were definitely explained in a way the patient could understand	68%	71%	*	*	*	64%	<b>69%</b>
Q45. Patient was always offered practical advice on dealing with any immediate side effects from treatment	63%	65%	*	*	*	65%	<b>64%</b>
Q46. Patient was given information that they could access about support in dealing with immediate side effects from treatment	83%	83%	*	*	*	87%	<b>83%</b>
Q47. Patient felt possible long-term side effects were definitely explained in a way they could understand in advance of their treatment	47%	59%	*	*	*	38%	<b>52%</b>
Q48. Patient was definitely able to discuss options for managing the impact of any long-term side effects	45%	52%	*	*	*	41%	<b>47%</b>

	Male/Female/Non-binary/Other						
	Female	Male	Non-binary	Prefer to self-describe	Prefer not to say	Not given	All
<b>SUPPORT WHILE AT HOME</b>							
Q49. Care team gave family, or someone close, all the information needed to help care for the patient at home	47%	60%	*	*	*	51%	<b>53%</b>
Q50. During treatment, the patient definitely got enough care and support at home from community or voluntary services	44%	45%	*	*	*	44%	<b>44%</b>

	Male/Female/Non-binary/Other						
	Female	Male	Non-binary	Prefer to self-describe	Prefer not to say	Not given	All
<b>CARE FROM YOUR GP PRACTICE</b>							
Q51. Patient definitely received the right amount of support from their GP practice during treatment	41%	43%	*	*	*	42%	<b>42%</b>
Q52. Patient has had a review of cancer care by GP practice	22%	20%	*	*	*	22%	<b>21%</b>

	Male/Female/Non-binary/Other						
	Female	Male	Non-binary	Prefer to self-describe	Prefer not to say	Not given	All
<b>LIVING WITH AND BEYOND CANCER</b>							
Q53. After treatment, the patient definitely could get enough emotional support at home from community or voluntary services	24%	24%	*	*	*	33%	<b>24%</b>
Q54. The right amount of information and support was offered to the patient between final treatment and the follow up appointment	72%	80%	*	*	*	63%	<b>74%</b>
Q55. Patient was given enough information about the possibility and signs of cancer coming back or spreading	53%	59%	*	*	*	54%	<b>56%</b>

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

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

YOUR OVERALL NHS CARE	Male/Female/Non-binary/Other						
	Female	Male	Non-binary	Prefer to self-describe	Prefer not to say	Not given	All
Q56. The whole care team worked well together	88%	89%	*	*	*	89%	<b>88%</b>
Q57. Administration of care was very good or good	85%	84%	*	*	*	85%	<b>85%</b>
Q58. Cancer research opportunities were discussed with patient	44%	43%	*	*	*	42%	<b>43%</b>
Q59. Patient's average rating of care scored from very poor to very good	8.8	8.8	*	*	*	8.9	<b>8.8</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	78%	*	88%	56%	*	81%	<b>78%</b>
Q3. Referral for diagnosis was explained in a way the patient could completely understand	64%	*	45%	60%	*	63%	<b>63%</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	91%	*	87%	91%	*	94%	<b>91%</b>
Q6. Diagnostic test staff appeared to completely have all the information they needed about the patient	81%	*	74%	76%	*	76%	<b>80%</b>
Q7. Patient felt the length of time waiting for diagnostic test results was about right	80%	*	74%	92%	*	79%	<b>80%</b>
Q8. Diagnostic test results were explained in a way the patient could completely understand	76%	*	65%	69%	*	80%	<b>76%</b>
Q9. Enough privacy was always given to the patient when receiving diagnostic test results	93%	*	100%	92%	*	98%	<b>94%</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	71%	60%	76%	73%	*	72%	<b>71%</b>
Q13. Patient was definitely told sensitively that they had cancer	69%	73%	77%	71%	*	70%	<b>69%</b>
Q14. Cancer diagnosis explained in a way the patient could completely understand	74%	91%	63%	64%	*	72%	<b>73%</b>
Q15. Patient was definitely told about their diagnosis in an appropriate place	83%	100%	88%	86%	*	80%	<b>83%</b>
Q16. Patient was told they could go back later for more information about their diagnosis	82%	*	87%	89%	*	83%	<b>82%</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	92%	80%	94%	96%	*	91%	<b>92%</b>
Q18. Patient found it very or quite easy to contact their main contact person	81%	*	58%	74%	*	78%	<b>80%</b>
Q19. Patient found advice from main contact person was very or quite helpful	94%	*	97%	96%	*	99%	<b>95%</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	81%	55%	80%	78%	*	76%	<b>80%</b>
Q21. Patient was definitely involved as much as they wanted to be in decisions about their treatment	76%	64%	74%	65%	*	72%	<b>76%</b>
Q22. Family and/or carers were definitely involved as much as the patient wanted them to be in decisions about treatment options	80%	80%	81%	68%	*	75%	<b>79%</b>
Q23. Patient could get further advice or a second opinion before making decisions about their treatment options	46%	*	59%	68%	*	42%	<b>46%</b>

## Ethnicity tables

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

CARE PLANNING	Ethnicity						
	White	Mixed	Asian	Black	Other	Not given	All
Q24. Patient was definitely able to have a discussion about their needs or concerns prior to treatment	68%	40%	68%	84%	*	65%	<b>68%</b>
Q25. A member of their care team helped the patient create a care plan to address any needs or concerns	89%	*	95%	94%	*	91%	<b>90%</b>
Q26. Care team reviewed the patient's care plan with them to ensure it was up to date	98%	*	94%	100%	*	100%	<b>98%</b>

SUPPORT FROM HOSPITAL STAFF	Ethnicity						
	White	Mixed	Asian	Black	Other	Not given	All
Q27. Staff provided the patient with relevant information on available support	89%	82%	88%	100%	*	94%	<b>89%</b>
Q28. Patient definitely got the right level of support for their overall health and well being from hospital staff	73%	73%	72%	93%	*	66%	<b>73%</b>
Q29. Patient was offered information about how to get financial help or benefits	66%	*	46%	82%	*	69%	<b>66%</b>

HOSPITAL CARE	Ethnicity						
	White	Mixed	Asian	Black	Other	Not given	All
Q31. Patient had confidence and trust in all of the team looking after them during their stay in hospital	79%	*	75%	79%	*	82%	<b>79%</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	62%	*	60%	55%	*	58%	<b>61%</b>
Q33. Patient was always involved in decisions about their care and treatment whilst in hospital	67%	*	69%	71%	*	63%	<b>67%</b>
Q34. Patient was always able to get help from ward staff when needed	71%	*	69%	85%	*	58%	<b>70%</b>
Q35. Patient was always able to discuss worries and fears with hospital staff	61%	*	80%	57%	*	46%	<b>61%</b>
Q36. Hospital staff always did everything they could to help the patient control pain	82%	*	60%	85%	*	72%	<b>80%</b>
Q37. Patient was always treated with respect and dignity while in hospital	86%	*	88%	86%	*	89%	<b>86%</b>
Q38. Patient received easily understandable information about what they should or should not do after leaving hospital	87%	*	75%	100%	*	80%	<b>87%</b>
Q39. Patient was always able to discuss worries and fears with hospital staff while being treated as an outpatient or day case	75%	*	68%	75%	*	73%	<b>74%</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	87%	*	95%	92%	*	88%	<b>87%</b>
Q41_2. Beforehand patient completely had enough understandable information about chemotherapy	82%	*	90%	95%	*	84%	<b>82%</b>
Q41_3. Beforehand patient completely had enough understandable information about radiotherapy	82%	*	86%	90%	*	87%	<b>82%</b>
Q41_4. Beforehand patient completely had enough understandable information about hormone therapy	68%	*	*	*	*	75%	<b>69%</b>
Q41_5. Beforehand patient completely had enough understandable information about immunotherapy	81%	*	*	*	*	77%	<b>80%</b>
Q42_1. Patient completely had enough understandable information about progress with surgery	82%	*	83%	86%	*	75%	<b>82%</b>
Q42_2. Patient completely had enough understandable information about progress with chemotherapy	74%	*	78%	75%	*	73%	<b>74%</b>
Q42_3. Patient completely had enough understandable information about progress with radiotherapy	75%	*	85%	80%	*	79%	<b>75%</b>
Q42_4. Patient completely had enough understandable information about progress with hormone therapy	63%	*	*	*	*	73%	<b>63%</b>
Q42_5. Patient completely had enough understandable information about progress with immunotherapy	72%	*	*	*	*	67%	<b>72%</b>
Q43. Patient felt the length of waiting time at clinic and day unit for cancer treatment was about right	71%	73%	61%	71%	*	61%	<b>70%</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	69%	55%	64%	86%	*	64%	<b>69%</b>
Q45. Patient was always offered practical advice on dealing with any immediate side effects from treatment	64%	55%	53%	68%	*	60%	<b>64%</b>
Q46. Patient was given information that they could access about support in dealing with immediate side effects from treatment	84%	*	75%	83%	*	81%	<b>83%</b>
Q47. Patient felt possible long-term side effects were definitely explained in a way they could understand in advance of their treatment	53%	*	50%	50%	*	42%	<b>52%</b>
Q48. Patient was definitely able to discuss options for managing the impact of any long-term side effects	48%	*	45%	53%	*	41%	<b>47%</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	53%	*	52%	56%	*	48%	<b>53%</b>
Q50. During treatment, the patient definitely got enough care and support at home from community or voluntary services	44%	*	50%	50%	*	44%	<b>44%</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	42%	*	50%	29%	*	43%	<b>42%</b>
Q52. Patient has had a review of cancer care by GP practice	21%	*	31%	17%	*	22%	<b>21%</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	24%	*	30%	*	*	26%	<b>24%</b>
Q54. The right amount of information and support was offered to the patient between final treatment and the follow up appointment	75%	*	73%	100%	*	66%	<b>74%</b>
Q55. Patient was given enough information about the possibility and signs of cancer coming back or spreading	56%	*	50%	58%	*	52%	<b>56%</b>

<b>YOUR OVERALL NHS CARE</b>	Ethnicity						
	White	Mixed	Asian	Black	Other	Not given	All
Q56. The whole care team worked well together	89%	91%	86%	83%	*	89%	<b>88%</b>
Q57. Administration of care was very good or good	84%	82%	88%	92%	*	86%	<b>85%</b>
Q58. Cancer research opportunities were discussed with patient	43%	*	31%	68%	*	48%	<b>43%</b>
Q59. Patient's average rating of care scored from very poor to very good	8.8	9.2	8.9	8.8	*	8.7	<b>8.8</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	83%	71%	75%	76%	81%	*	<b>78%</b>
Q3. Referral for diagnosis was explained in a way the patient could completely understand	75%	58%	60%	61%	67%	*	<b>63%</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	88%	91%	89%	92%	90%	*	<b>91%</b>
Q6. Diagnostic test staff appeared to completely have all the information they needed about the patient	84%	80%	78%	82%	80%	*	<b>80%</b>
Q7. Patient felt the length of time waiting for diagnostic test results was about right	79%	83%	79%	81%	78%	*	<b>80%</b>
Q8. Diagnostic test results were explained in a way the patient could completely understand	80%	72%	73%	75%	78%	*	<b>76%</b>
Q9. Enough privacy was always given to the patient when receiving diagnostic test results	100%	93%	92%	93%	95%	*	<b>94%</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	62%	80%	72%	70%	69%	*	<b>71%</b>
Q13. Patient was definitely told sensitively that they had cancer	71%	76%	69%	65%	68%	*	<b>69%</b>
Q14. Cancer diagnosis explained in a way the patient could completely understand	86%	73%	73%	71%	74%	*	<b>73%</b>
Q15. Patient was definitely told about their diagnosis in an appropriate place	91%	90%	84%	82%	82%	*	<b>83%</b>
Q16. Patient was told they could go back later for more information about their diagnosis	75%	84%	82%	80%	83%	*	<b>82%</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	86%	94%	90%	92%	93%	*	<b>92%</b>
Q18. Patient found it very or quite easy to contact their main contact person	71%	80%	82%	78%	80%	*	<b>80%</b>
Q19. Patient found advice from main contact person was very or quite helpful	100%	96%	96%	95%	94%	*	<b>95%</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
Q20. Treatment options were explained in a way the patient could completely understand	85%	79%	79%	83%	80%	*	<b>80%</b>
Q21. Patient was definitely involved as much as they wanted to be in decisions about their treatment	86%	76%	78%	74%	74%	*	<b>76%</b>
Q22. Family and/or carers were definitely involved as much as the patient wanted them to be in decisions about treatment options	59%	81%	78%	79%	80%	*	<b>79%</b>
Q23. Patient could get further advice or a second opinion before making decisions about their treatment options	38%	46%	44%	47%	48%	*	<b>46%</b>

	IMD Quintile						
	1 (most deprived)	2	3	4	5 (least deprived)	Non-England	All
Q24. Patient was definitely able to have a discussion about their needs or concerns prior to treatment	79%	74%	65%	69%	66%	*	<b>68%</b>
Q25. A member of their care team helped the patient create a care plan to address any needs or concerns	*	97%	87%	93%	88%	*	<b>90%</b>
Q26. Care team reviewed the patient's care plan with them to ensure it was up to date	*	100%	98%	99%	98%	*	<b>98%</b>

	IMD Quintile						
	1 (most deprived)	2	3	4	5 (least deprived)	Non-England	All
Q27. Staff provided the patient with relevant information on available support	94%	92%	87%	87%	91%	*	<b>89%</b>
Q28. Patient definitely got the right level of support for their overall health and well being from hospital staff	73%	79%	72%	75%	70%	*	<b>73%</b>
Q29. Patient was offered information about how to get financial help or benefits	79%	58%	74%	65%	65%	*	<b>66%</b>

	IMD Quintile						
	1 (most deprived)	2	3	4	5 (least deprived)	Non-England	All
Q31. Patient had confidence and trust in all of the team looking after them during their stay in hospital	*	79%	73%	80%	81%	*	<b>79%</b>
Q32. Patient's family, or someone close, was definitely able to talk to a member of the team looking after the patient in hospital	*	61%	63%	63%	60%	*	<b>61%</b>
Q33. Patient was always involved in decisions about their care and treatment whilst in hospital	*	63%	63%	75%	67%	*	<b>67%</b>
Q34. Patient was always able to get help from ward staff when needed	*	73%	66%	75%	70%	*	<b>70%</b>
Q35. Patient was always able to discuss worries and fears with hospital staff	*	66%	56%	65%	60%	*	<b>61%</b>
Q36. Hospital staff always did everything they could to help the patient control pain	*	79%	73%	88%	81%	*	<b>80%</b>
Q37. Patient was always treated with respect and dignity while in hospital	*	81%	82%	92%	86%	*	<b>86%</b>
Q38. Patient received easily understandable information about what they should or should not do after leaving hospital	*	85%	87%	88%	86%	*	<b>87%</b>
Q39. Patient was always able to discuss worries and fears with hospital staff while being treated as an outpatient or day case	69%	76%	68%	77%	75%	*	<b>74%</b>

## IMD quintile tables

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

YOUR TREATMENT	IMD Quintile						
	1 (most deprived)	2	3	4	5 (least deprived)	Non-England	All
Q41_1. Beforehand patient completely had enough understandable information about surgery	80%	88%	84%	92%	87%	*	<b>87%</b>
Q41_2. Beforehand patient completely had enough understandable information about chemotherapy	*	87%	82%	80%	82%	*	<b>82%</b>
Q41_3. Beforehand patient completely had enough understandable information about radiotherapy	*	86%	83%	89%	78%	*	<b>82%</b>
Q41_4. Beforehand patient completely had enough understandable information about hormone therapy	*	63%	76%	67%	67%	*	<b>69%</b>
Q41_5. Beforehand patient completely had enough understandable information about immunotherapy	*	96%	86%	79%	75%	*	<b>80%</b>
Q42_1. Patient completely had enough understandable information about progress with surgery	*	83%	79%	86%	80%	*	<b>82%</b>
Q42_2. Patient completely had enough understandable information about progress with chemotherapy	*	79%	71%	76%	73%	*	<b>74%</b>
Q42_3. Patient completely had enough understandable information about progress with radiotherapy	*	84%	71%	76%	74%	*	<b>75%</b>
Q42_4. Patient completely had enough understandable information about progress with hormone therapy	*	64%	65%	58%	65%	*	<b>63%</b>
Q42_5. Patient completely had enough understandable information about progress with immunotherapy	*	96%	83%	69%	66%	*	<b>72%</b>
Q43. Patient felt the length of waiting time at clinic and day unit for cancer treatment was about right	82%	70%	70%	74%	68%	*	<b>70%</b>

IMMEDIATE AND LONG TERM SIDE EFFECTS	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	62%	74%	70%	67%	69%	*	<b>69%</b>
Q45. Patient was always offered practical advice on dealing with any immediate side effects from treatment	68%	70%	62%	66%	62%	*	<b>64%</b>
Q46. Patient was given information that they could access about support in dealing with immediate side effects from treatment	81%	83%	79%	85%	85%	*	<b>83%</b>
Q47. Patient felt possible long-term side effects were definitely explained in a way they could understand in advance of their treatment	55%	58%	51%	51%	51%	*	<b>52%</b>
Q48. Patient was definitely able to discuss options for managing the impact of any long-term side effects	56%	51%	47%	45%	48%	*	<b>47%</b>

SUPPORT WHILE AT HOME	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	53%	61%	50%	52%	53%	*	<b>53%</b>
Q50. During treatment, the patient definitely got enough care and support at home from community or voluntary services	45%	47%	42%	46%	43%	*	<b>44%</b>

CARE FROM YOUR GP PRACTICE	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	59%	41%	37%	43%	43%	*	<b>42%</b>
Q52. Patient has had a review of cancer care by GP practice	24%	20%	20%	20%	22%	*	<b>21%</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	*	11%	32%	32%	21%	*	<b>24%</b>
Q54. The right amount of information and support was offered to the patient between final treatment and the follow up appointment	73%	75%	74%	74%	75%	*	<b>74%</b>
Q55. Patient was given enough information about the possibility and signs of cancer coming back or spreading	53%	57%	58%	54%	55%	*	<b>56%</b>

	IMD Quintile						
	1 (most deprived)	2	3	4	5 (least deprived)	Non-England	All
Q56. The whole care team worked well together	95%	88%	89%	86%	89%	*	<b>88%</b>
Q57. Administration of care was very good or good	91%	82%	85%	87%	84%	*	<b>85%</b>
Q58. Cancer research opportunities were discussed with patient	36%	56%	46%	43%	40%	*	<b>43%</b>
Q59. Patient's average rating of care scored from very poor to very good	8.9	8.8	8.7	8.8	8.8	*	<b>8.8</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	78%	75%	84%	78%
Q3. Referral for diagnosis was explained in a way the patient could completely understand	60%	68%	65%	63%

	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%	92%	93%	91%
Q6. Diagnostic test staff appeared to completely have all the information they needed about the patient	78%	85%	75%	80%
Q7. Patient felt the length of time waiting for diagnostic test results was about right	79%	81%	81%	80%
Q8. Diagnostic test results were explained in a way the patient could completely understand	74%	78%	80%	76%
Q9. Enough privacy was always given to the patient when receiving diagnostic test results	92%	96%	98%	94%

	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	70%	72%	75%	71%
Q13. Patient was definitely told sensitively that they had cancer	66%	72%	75%	69%
Q14. Cancer diagnosis explained in a way the patient could completely understand	71%	75%	77%	73%
Q15. Patient was definitely told about their diagnosis in an appropriate place	83%	83%	86%	83%
Q16. Patient was told they could go back later for more information about their diagnosis	80%	85%	84%	82%

	Long term condition status			
	Yes	No	Not given	All
<b>SUPPORT FROM A MAIN CONTACT PERSON</b>				
Q17. Patient had a main point of contact within the care team	91%	93%	93%	92%
Q18. Patient found it very or quite easy to contact their main contact person	79%	80%	82%	80%
Q19. Patient found advice from main contact person was very or quite helpful	95%	94%	99%	95%

	Long term condition status			
	Yes	No	Not given	All
<b>DECIDING ON THE BEST TREATMENT</b>				
Q20. Treatment options were explained in a way the patient could completely understand	79%	82%	85%	80%
Q21. Patient was definitely involved as much as they wanted to be in decisions about their treatment	72%	81%	75%	76%
Q22. Family and/or carers were definitely involved as much as the patient wanted them to be in decisions about treatment options	79%	79%	78%	79%
Q23. Patient could get further advice or a second opinion before making decisions about their treatment options	46%	45%	57%	46%

## Long term condition status tables

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

<b>CARE PLANNING</b>	Long term condition status			
	Yes	No	Not given	All
Q24. Patient was definitely able to have a discussion about their needs or concerns prior to treatment	66%	70%	71%	<b>68%</b>
Q25. A member of their care team helped the patient create a care plan to address any needs or concerns	89%	90%	96%	<b>90%</b>
Q26. Care team reviewed the patient's care plan with them to ensure it was up to date	99%	96%	100%	<b>98%</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	89%	90%	91%	<b>89%</b>
Q28. Patient definitely got the right level of support for their overall health and well being from hospital staff	71%	76%	70%	<b>73%</b>
Q29. Patient was offered information about how to get financial help or benefits	65%	69%	68%	<b>66%</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	77%	82%	81%	<b>79%</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%	66%	65%	<b>61%</b>
Q33. Patient was always involved in decisions about their care and treatment whilst in hospital	64%	71%	68%	<b>67%</b>
Q34. Patient was always able to get help from ward staff when needed	70%	71%	68%	<b>70%</b>
Q35. Patient was always able to discuss worries and fears with hospital staff	59%	65%	53%	<b>61%</b>
Q36. Hospital staff always did everything they could to help the patient control pain	79%	82%	84%	<b>80%</b>
Q37. Patient was always treated with respect and dignity while in hospital	85%	86%	95%	<b>86%</b>
Q38. Patient received easily understandable information about what they should or should not do after leaving hospital	86%	89%	84%	<b>87%</b>
Q39. Patient was always able to discuss worries and fears with hospital staff while being treated as an outpatient or day case	72%	77%	80%	<b>74%</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	87%	87%	94%	<b>87%</b>
Q41_2. Beforehand patient completely had enough understandable information about chemotherapy	82%	82%	86%	<b>82%</b>
Q41_3. Beforehand patient completely had enough understandable information about radiotherapy	82%	82%	83%	<b>82%</b>
Q41_4. Beforehand patient completely had enough understandable information about hormone therapy	69%	66%	92%	<b>69%</b>
Q41_5. Beforehand patient completely had enough understandable information about immunotherapy	78%	84%	82%	<b>80%</b>
Q42_1. Patient completely had enough understandable information about progress with surgery	79%	85%	84%	<b>82%</b>
Q42_2. Patient completely had enough understandable information about progress with chemotherapy	71%	77%	74%	<b>74%</b>
Q42_3. Patient completely had enough understandable information about progress with radiotherapy	74%	79%	63%	<b>75%</b>
Q42_4. Patient completely had enough understandable information about progress with hormone therapy	62%	64%	80%	<b>63%</b>
Q42_5. Patient completely had enough understandable information about progress with immunotherapy	68%	79%	*	<b>72%</b>
Q43. Patient felt the length of waiting time at clinic and day unit for cancer treatment was about right	70%	71%	68%	<b>70%</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	65%	76%	68%	<b>69%</b>
Q45. Patient was always offered practical advice on dealing with any immediate side effects from treatment	60%	69%	69%	<b>64%</b>
Q46. Patient was given information that they could access about support in dealing with immediate side effects from treatment	82%	85%	89%	<b>83%</b>
Q47. Patient felt possible long-term side effects were definitely explained in a way they could understand in advance of their treatment	50%	55%	49%	<b>52%</b>
Q48. Patient was definitely able to discuss options for managing the impact of any long-term side effects	46%	50%	46%	<b>47%</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	53%	52%	55%	<b>53%</b>
Q50. During treatment, the patient definitely got enough care and support at home from community or voluntary services	42%	47%	55%	<b>44%</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	42%	42%	45%	<b>42%</b>
Q52. Patient has had a review of cancer care by GP practice	21%	20%	23%	<b>21%</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	21%	31%	30%	<b>24%</b>
Q54. The right amount of information and support was offered to the patient between final treatment and the follow up appointment	72%	79%	68%	<b>74%</b>
Q55. Patient was given enough information about the possibility and signs of cancer coming back or spreading	54%	58%	56%	<b>56%</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	87%	90%	91%	<b>88%</b>
Q57. Administration of care was very good or good	84%	85%	89%	<b>85%</b>
Q58. Cancer research opportunities were discussed with patient	41%	48%	40%	<b>43%</b>
Q59. Patient's average rating of care scored from very poor to very good	8.7	8.9	9.0	<b>8.8</b>

## Year on Year Charts

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

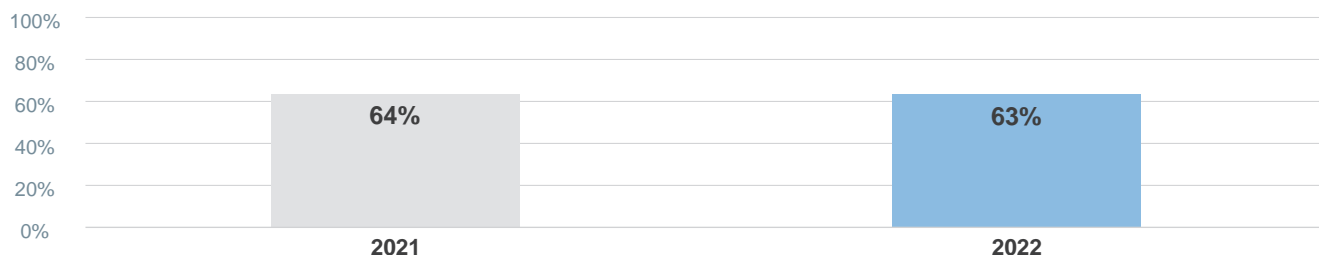
The scores are unadjusted and based on England scores only.

### SUPPORT FROM YOUR GP PRACTICE

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

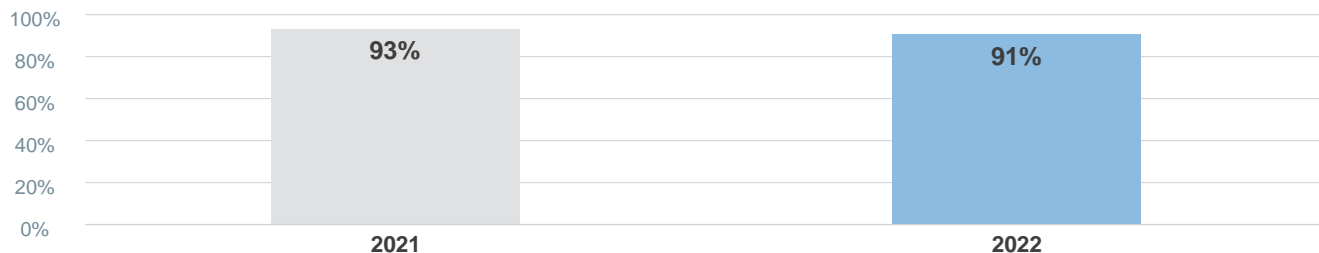


Q3. Referral for diagnosis was explained in a way the patient could completely understand

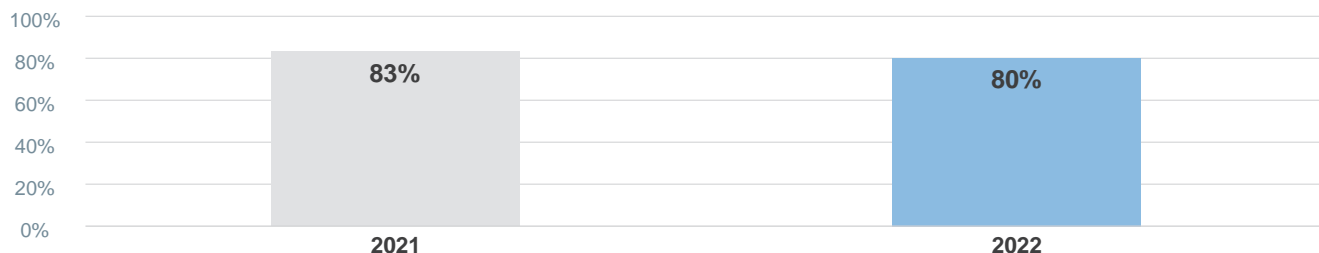


### DIAGNOSTIC TESTS

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

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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



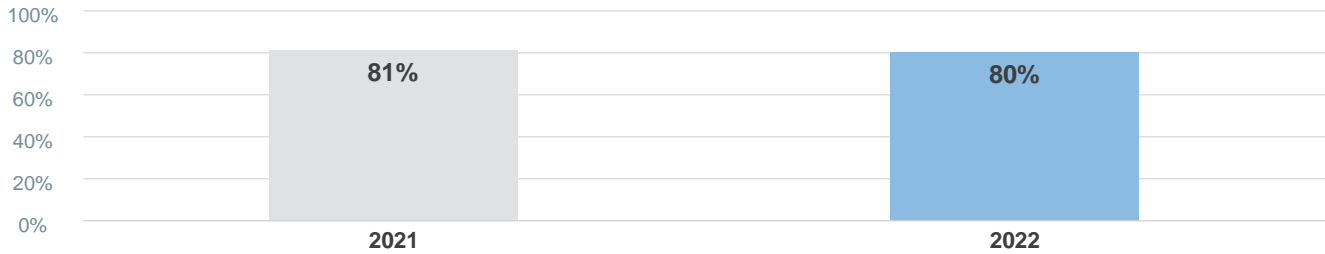
## Year on Year Charts

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### DECIDING ON THE BEST TREATMENT

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



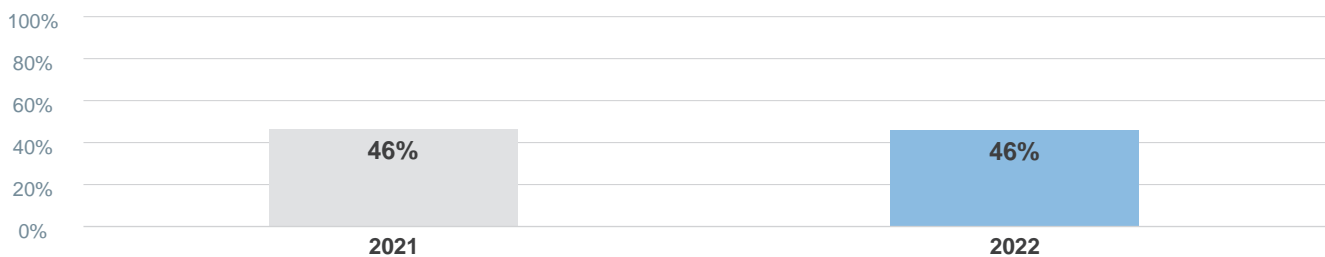
Q21. Patient was definitely involved as much as they wanted to be in decisions about their treatment



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

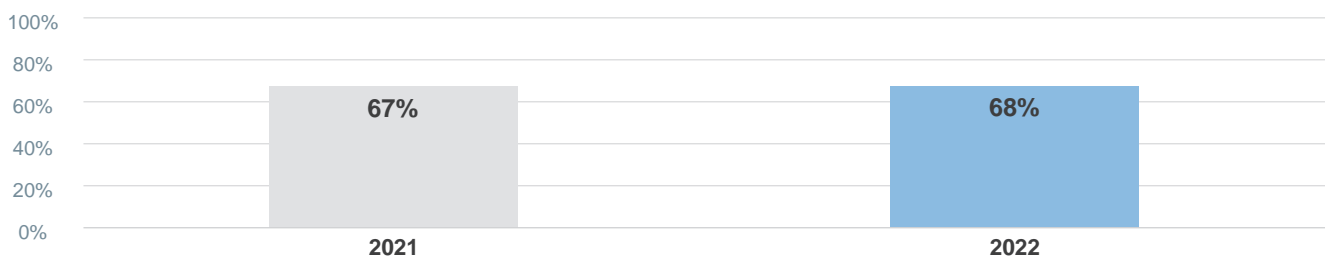


Q23. Patient could get further advice or a second opinion before making decisions about their treatment options



### CARE PLANNING

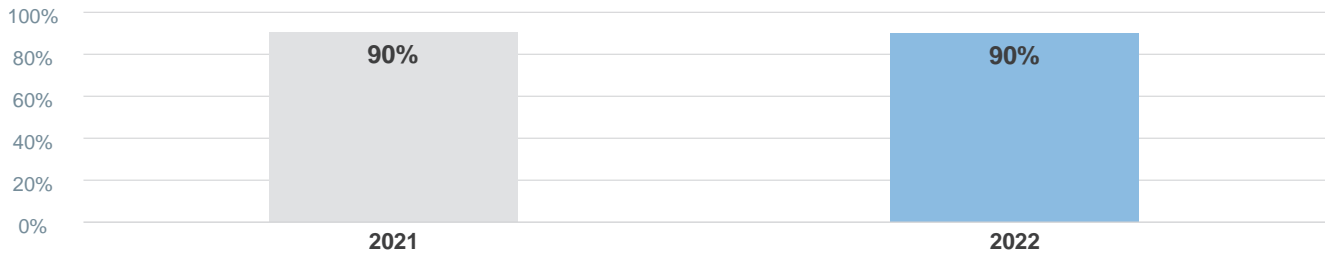
Q24. Patient was definitely able to have a discussion about their needs or concerns prior to treatment



## Year on Year Charts

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



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



### SUPPORT FROM HOSPITAL STAFF

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



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



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



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### 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

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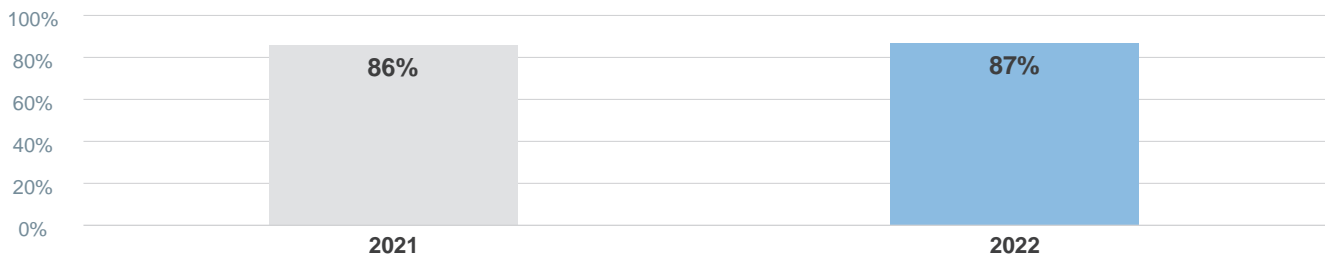
Q36. Hospital staff always did everything they could to help the patient control pain



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

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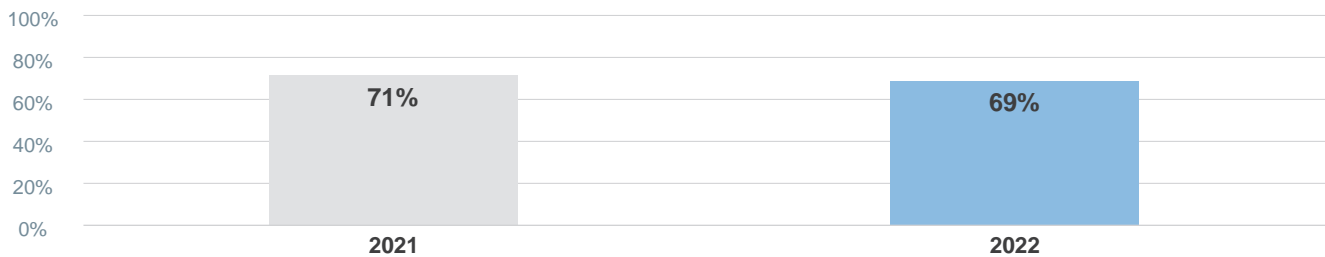
**Q41\_2. Beforehand patient completely had enough understandable information about chemotherapy**



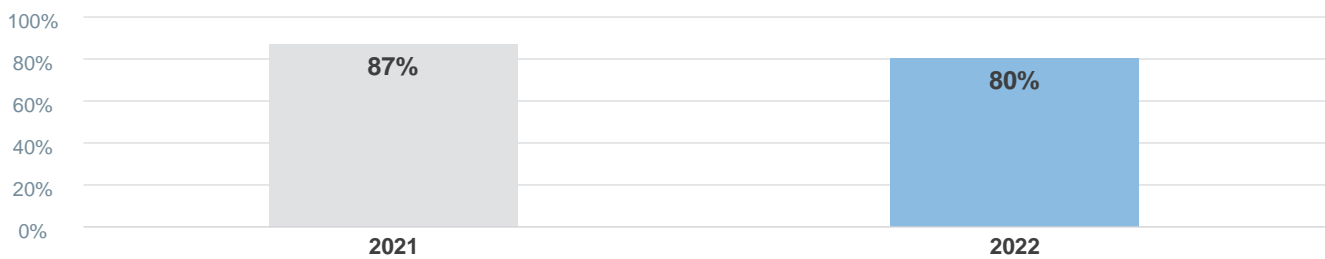
**Q41\_3. Beforehand patient completely had enough understandable information about radiotherapy**



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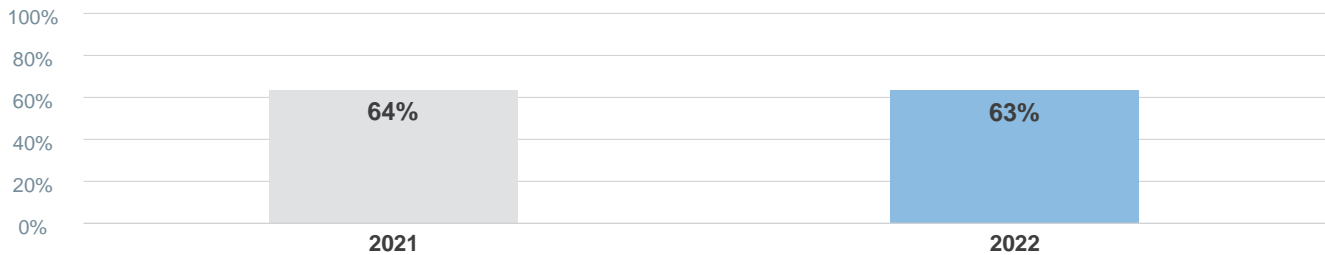
**Q42\_2. Patient completely had enough understandable information about progress with chemotherapy**



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



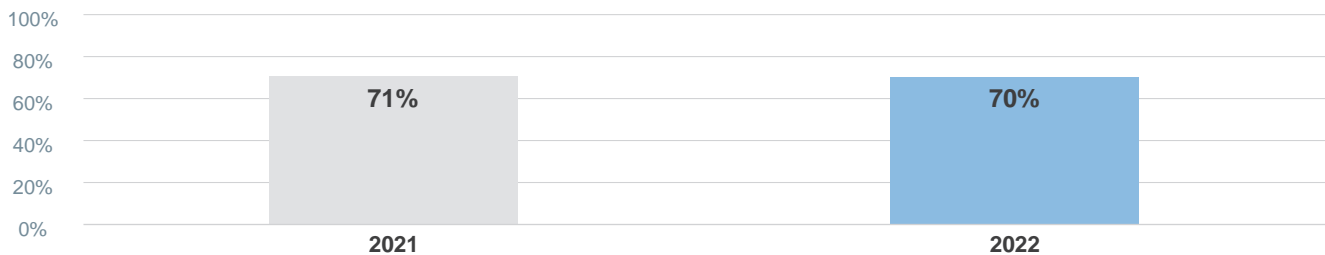
**Q42\_4. Patient completely had enough understandable information about progress with hormone therapy**



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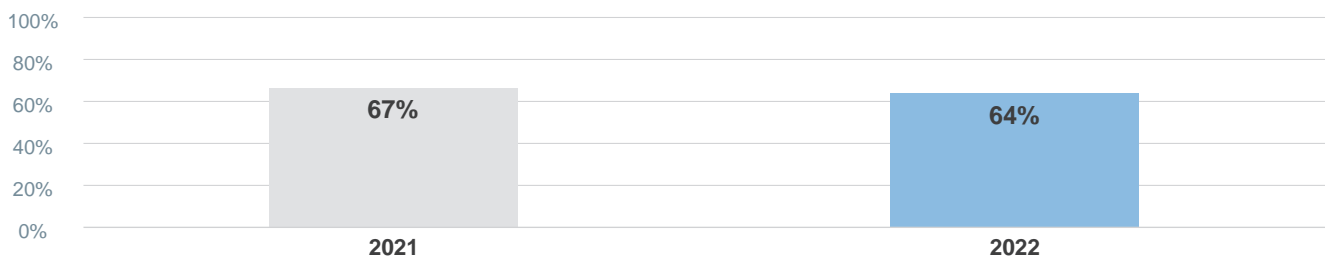
The scores are unadjusted and based on England scores only.

### IMMEDIATE AND LONG TERM SIDE EFFECTS

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



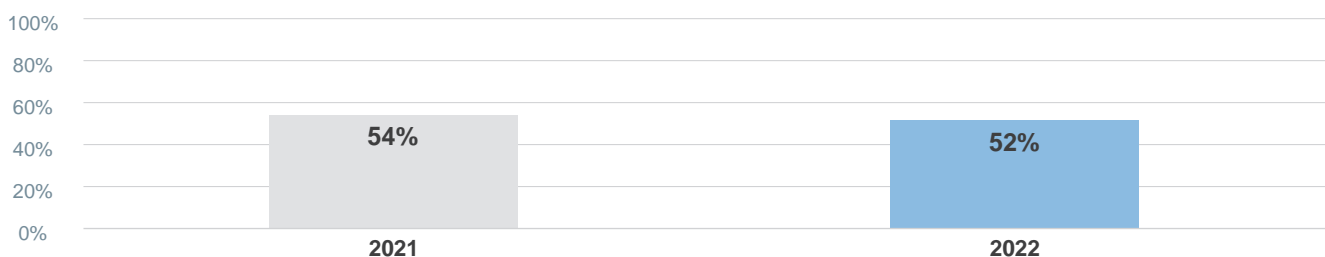
Q45. Patient was always offered practical advice on dealing with any immediate side effects from treatment



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



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



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



## Year on Year Charts

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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

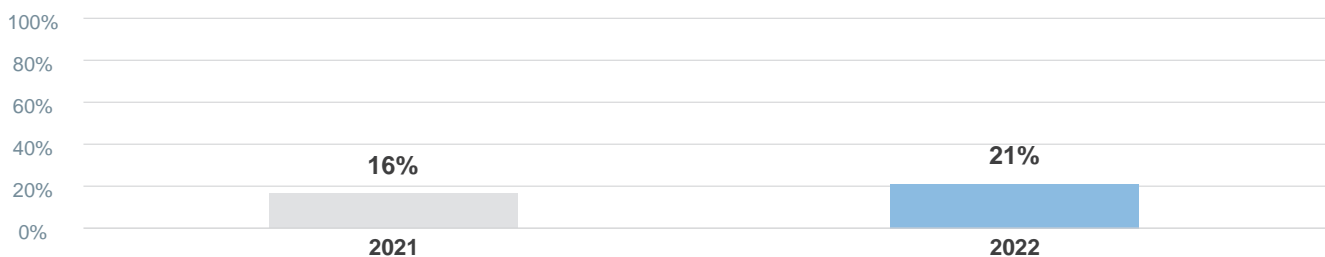


### CARE FROM YOUR GP PRACTICE

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



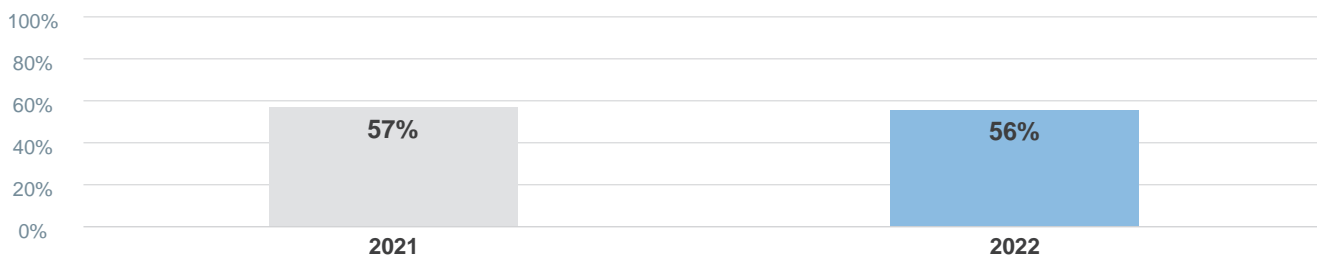
## Year on Year Charts

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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**

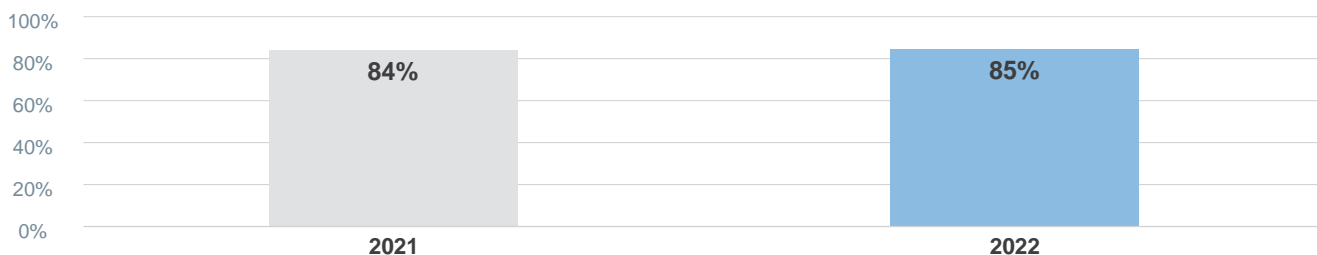


### YOUR OVERALL NHS CARE

**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

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

