

# **Cancer Patient Experience Survey**

2023 Results

## **Northern Cancer Alliance**

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

## Contents

Executive summary . . . . .	3
Introduction . . . . .	4
Methodology . . . . .	4
Understanding the results . . . . .	5
Further information . . . . .	7
Response rate . . . . .	8
Expected range charts . . . . .	10
Comparability tables . . . . .	14
Tumour group tables . . . . .	18
Age group tables . . . . .	23
Male/Female/Non-binary/Other tables . . . . .	27
Ethnicity tables . . . . .	32
IMD quintile tables . . . . .	36
Long-term condition status tables . . . . .	40
Year on year charts . . . . .	44
Trust expected range summary . . . . .	57
ICB expected range summary . . . . .	58

## Executive summary

### Questions above expected range

	Case mix adjusted scores			England score
	2023 score	Lower expected range	Upper expected range	
Q7. Patient felt the length of time waiting for diagnostic test results was about right	<b>81%</b>	75%	80%	<b>78%</b>
Q26. Care team reviewed the patient's care plan with them to ensure it was up to date	<b>100%</b>	98%	99%	<b>99%</b>
Q32. Patient's family, or someone close, was definitely able to talk to a member of the team looking after the patient in hospital	<b>73%</b>	67%	72%	<b>70%</b>
Q41_2. Beforehand patient completely had enough understandable information about chemotherapy	<b>88%</b>	84%	87%	<b>86%</b>
Q42_2. Patient completely had enough understandable information about their response to chemotherapy	<b>84%</b>	79%	83%	<b>81%</b>
Q42_4. Patient completely had enough understandable information about their response to hormone therapy	<b>81%</b>	73%	80%	<b>76%</b>
Q42_5. Patient completely had enough understandable information about their response to immunotherapy	<b>84%</b>	78%	83%	<b>81%</b>
Q44. Possible side effects from treatment were definitely explained in a way the patient could understand	<b>77%</b>	73%	76%	<b>74%</b>
Q48. Patient was definitely able to discuss options for managing the impact of any long-term side effects	<b>60%</b>	51%	58%	<b>55%</b>
Q50. During treatment, the patient definitely got enough care and support at home from community or voluntary services	<b>58%</b>	47%	58%	<b>52%</b>
Q53. After treatment, the patient definitely could get enough emotional support at home from community or voluntary services	<b>39%</b>	28%	37%	<b>32%</b>
Q56. The whole care team worked well together	<b>92%</b>	88%	92%	<b>90%</b>
Q57. Administration of care was very good or good	<b>90%</b>	85%	89%	<b>87%</b>

### Questions below expected range

Northern Cancer Alliance has no scores below expected range.

## Introduction

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

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

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

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

## Methodology

### Eligibility, fieldwork and survey methods

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

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

### Note on question comparability

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

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

### How alliance and ICB results are generated

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

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

### Case-mix adjustment

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

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

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

### Scoring methodology

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

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

### Statistical significance

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

### Suppression

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

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

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

### Additional suppression

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

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

## Understanding the results

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

### Expected range charts

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

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

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

### **Comparability tables**

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

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

### **Sub-group breakdowns**

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

### **Tumour group tables**

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

### **Age group tables**

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

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

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

### **Ethnicity tables**

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

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

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

### **IMD quintile tables**

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

### **Year on year charts**

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

### **Trust expected range summary**

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

### **ICB expected range summary**

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

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

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

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

## **Further information**

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

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

## Response rate

### Overall response rate

4,273 patients responded out of a total of 7,920 patients, resulting in a response rate of 54%.

	Sample size	Adjusted sample	Completed	Response rate
Overall response rate	8,552	7,920	4,273	54%
National	129,231	121,121	63,438	52%

### Respondents by survey type

	Number of respondents
Paper	3,537
Online	734
Phone	2
Translation service	0
<b>Total</b>	<b>4,273</b>

### Respondents by tumour group

	Number of respondents
Brain / CNS	11
Breast	895
Colorectal / LGT	512
Gynaecological	180
Haematological	778
Head and neck	103
Lung	362
Prostate	373
Sarcoma	18
Skin	74
Upper gastro	200
Urological	341
Other	426
<b>Total</b>	<b>4,273</b>

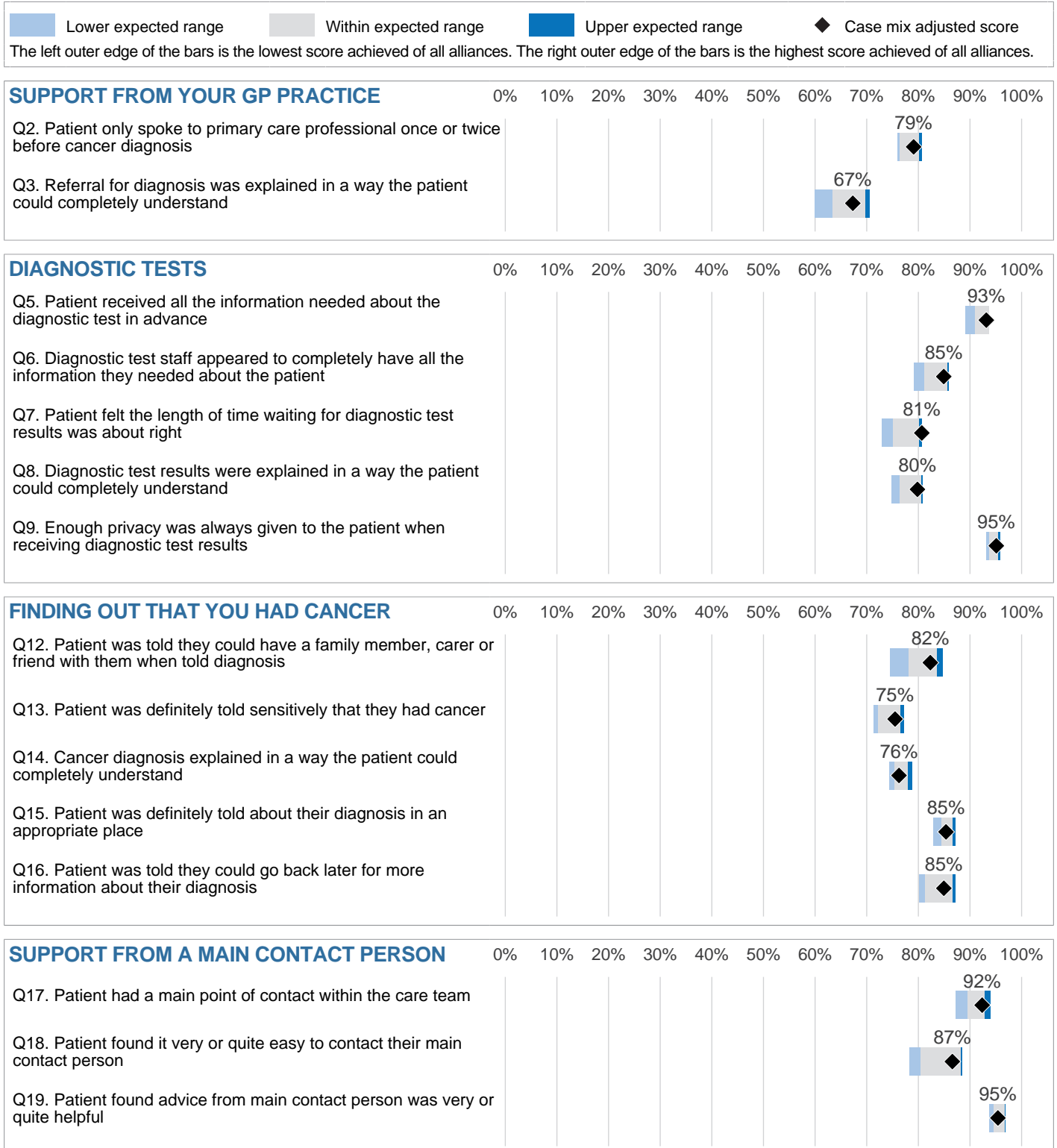


## Respondents by ethnicity

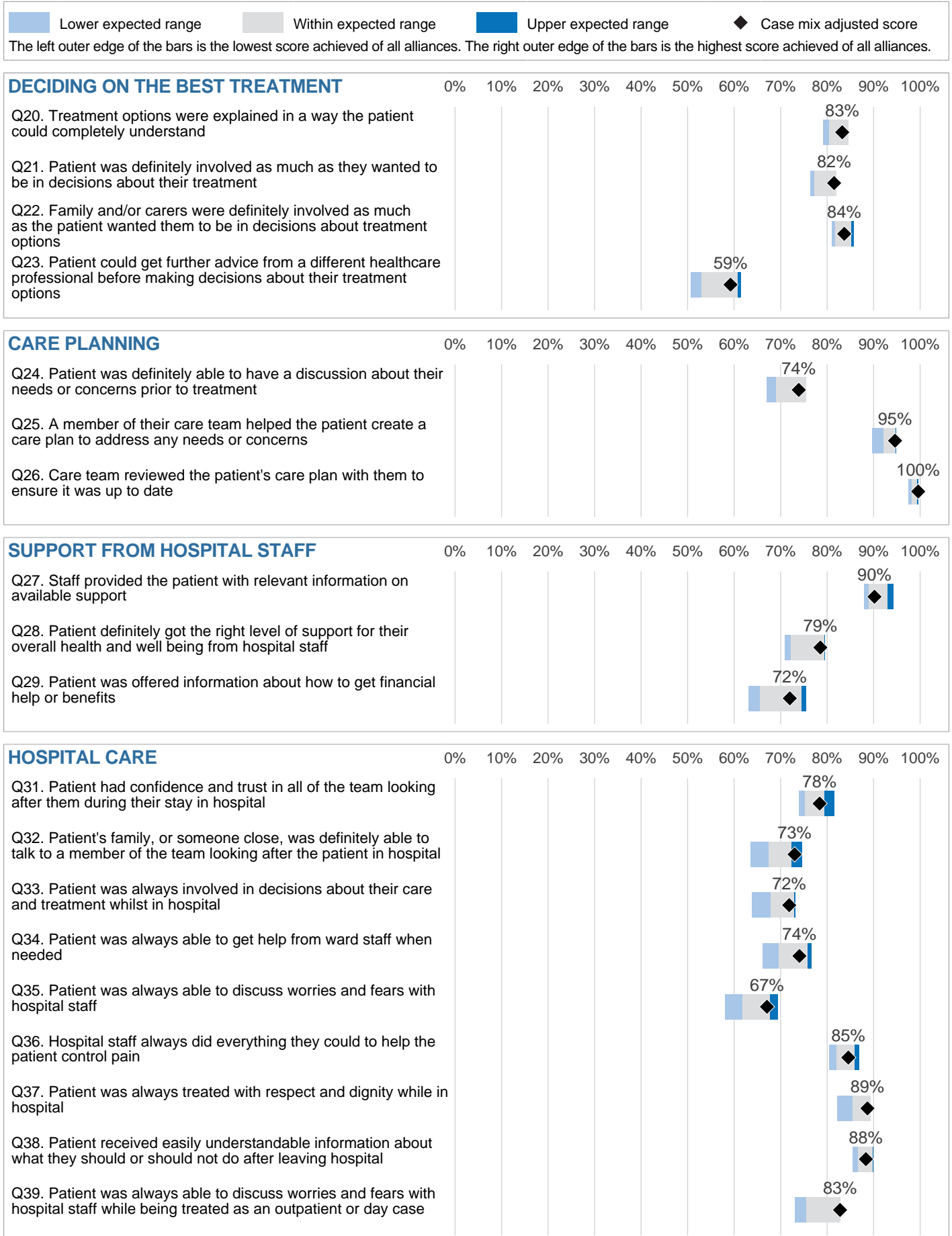
	Number of respondents
<b>White</b>	
English / Welsh / Scottish / Northern Irish / British	3,832
Irish	6
Gypsy or Irish Traveller	*
Roma	*
Any other White background	29
<b>Mixed / Multiple Ethnic Groups</b>	
White and Black Caribbean	8
White and Black African	*
White and Asian	*
Any other Mixed / multiple ethnic background	6
<b>Asian or Asian British</b>	
Indian	9
Pakistani	14
Bangladeshi	6
Chinese	7
Any other Asian background	9
<b>Black / African / Caribbean / Black British</b>	
African	*
Caribbean	*
Any other Black / African / Caribbean background	*
<b>Other Ethnicity</b>	
Arab	*
Any other ethnic group	6
<b>Not given</b>	
Not given	320
<b>Total</b>	<b>4,273</b>

\* indicates the count is not shown due to suppression

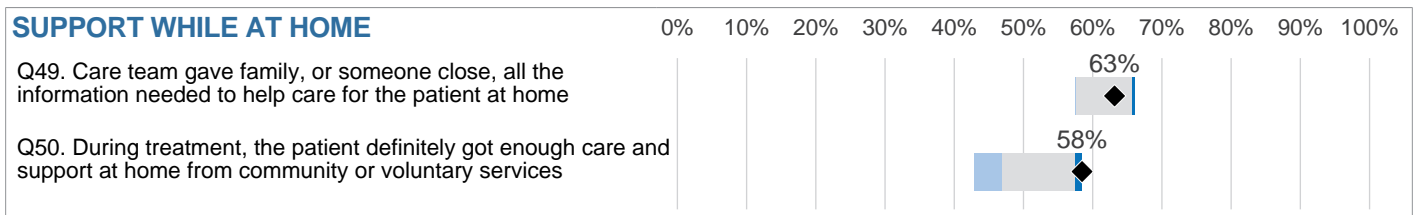
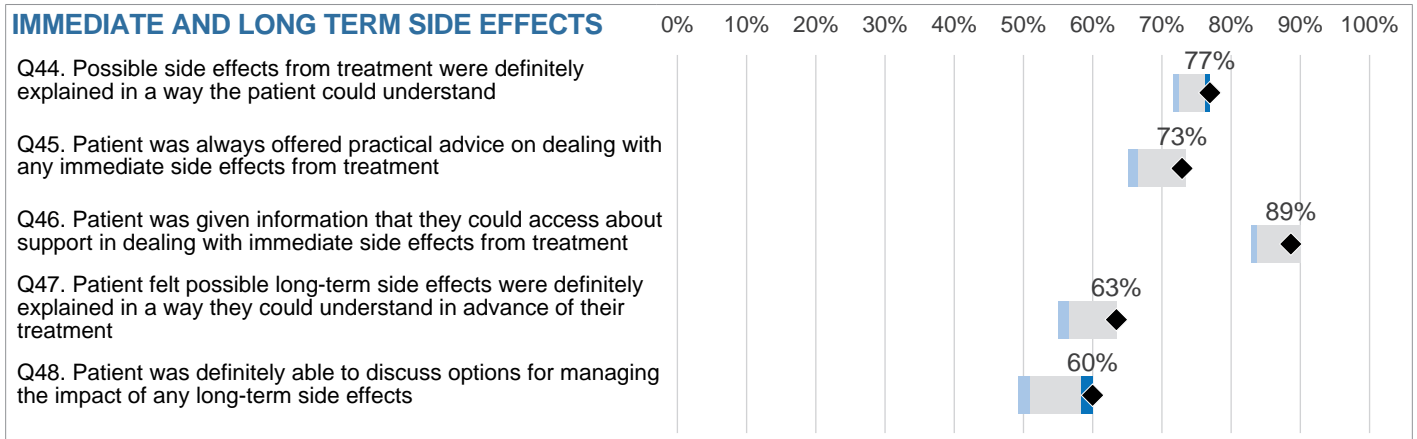
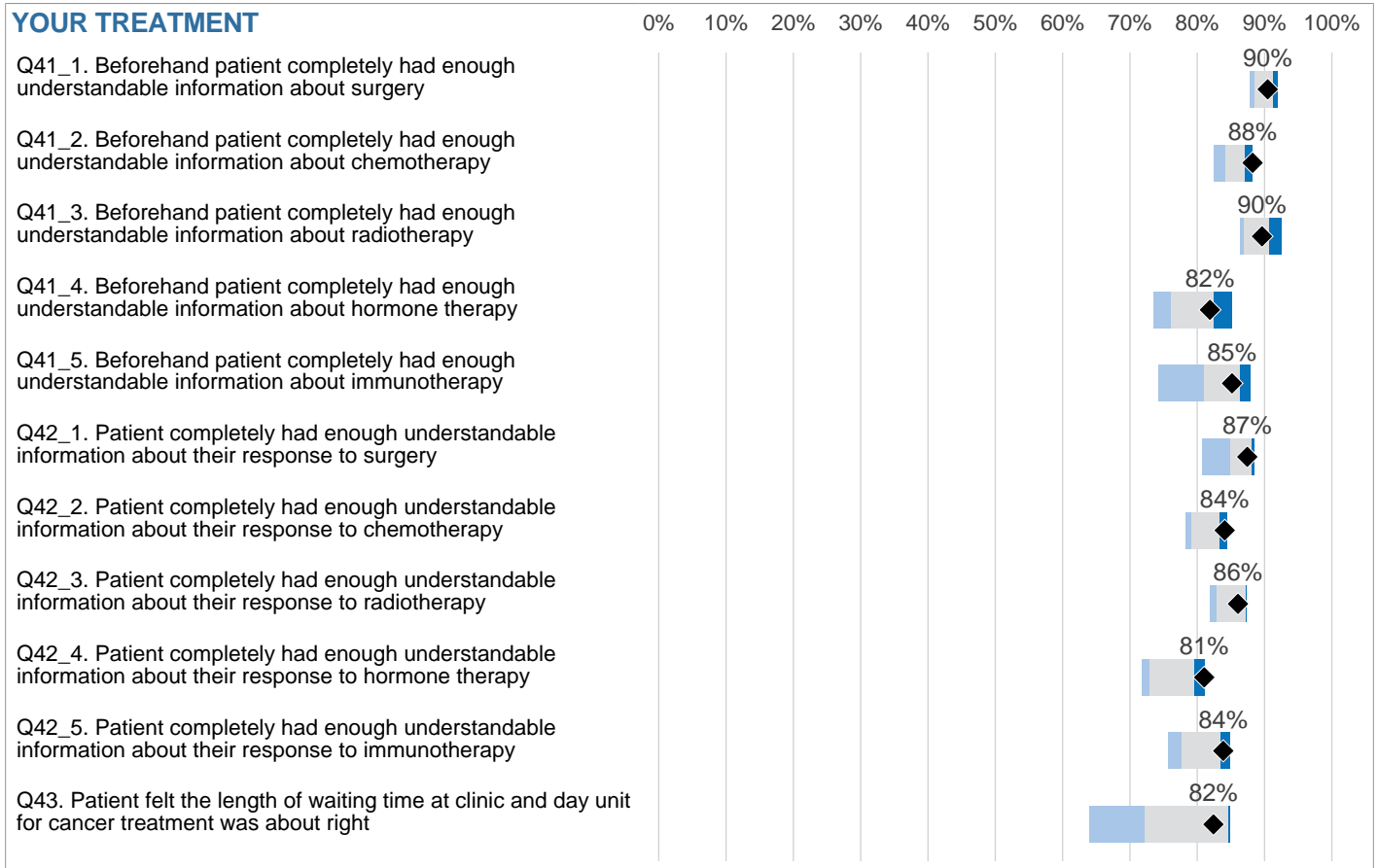
## Expected range charts



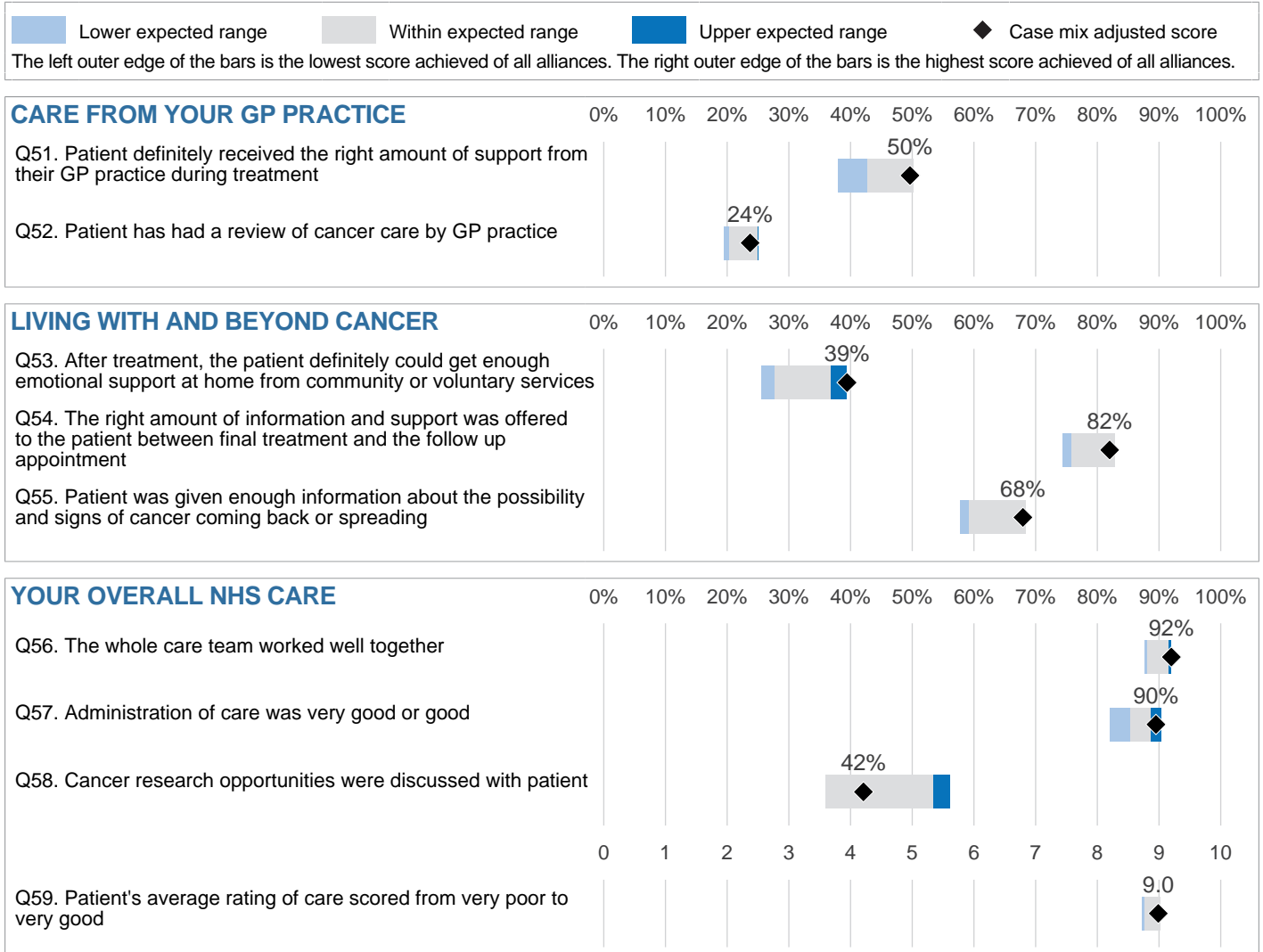
## Expected range charts



## Expected range charts






## Expected range charts



## Comparability tables

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

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

	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
	2022 n	2022 score	2023 n	2023 score	Change 2022-2023	Change overall	2023 score	Lower expected range	Upper expected range	
Q2. Patient only spoke to primary care professional once or twice before cancer diagnosis	2021	<b>78%</b>	1995	<b>78%</b>			<b>79%</b>	76%	80%	<b>78%</b>
Q3. Referral for diagnosis was explained in a way the patient could completely understand	2803	<b>64%</b>	2765	<b>66%</b>			<b>67%</b>	63%	70%	<b>67%</b>

DIAGNOSTIC TESTS	Unadjusted scores						Case mix adjusted scores			England score
	2022 n	2022 score	2023 n	2023 score	Change 2022-2023	Change overall	2023 score	Lower expected range	Upper expected range	
Q5. Patient received all the information needed about the diagnostic test in advance	3341	<b>93%</b>	3196	<b>93%</b>			<b>93%</b>	91%	94%	<b>92%</b>
Q6. Diagnostic test staff appeared to completely have all the information they needed about the patient	3535	<b>84%</b>	3342	<b>85%</b>			<b>85%</b>	81%	86%	<b>83%</b>
Q7. Patient felt the length of time waiting for diagnostic test results was about right	3539	<b>79%</b>	3335	<b>81%</b>		▼	<b>81%</b>	75%	80%	<b>78%</b>
Q8. Diagnostic test results were explained in a way the patient could completely understand	3549	<b>79%</b>	3353	<b>80%</b>			<b>80%</b>	76%	80%	<b>78%</b>
Q9. Enough privacy was always given to the patient when receiving diagnostic test results	3549	<b>95%</b>	3358	<b>95%</b>			<b>95%</b>	94%	96%	<b>95%</b>

FINDING OUT THAT YOU HAD CANCER	Unadjusted scores						Case mix adjusted scores			England score
	2022 n	2022 score	2023 n	2023 score	Change 2022-2023	Change overall	2023 score	Lower expected range	Upper expected range	
Q12. Patient was told they could have a family member, carer or friend with them when told diagnosis	4045	<b>80%</b>	3977	<b>82%</b>	▲	▲	<b>82%</b>	78%	84%	<b>81%</b>
Q13. Patient was definitely told sensitively that they had cancer	4275	<b>75%</b>	4209	<b>75%</b>			<b>75%</b>	72%	77%	<b>74%</b>
Q14. Cancer diagnosis explained in a way the patient could completely understand	4270	<b>77%</b>	4208	<b>76%</b>			<b>76%</b>	75%	78%	<b>77%</b>
Q15. Patient was definitely told about their diagnosis in an appropriate place	4264	<b>85%</b>	4201	<b>85%</b>			<b>85%</b>	84%	87%	<b>86%</b>
Q16. Patient was told they could go back later for more information about their diagnosis	3813	<b>84%</b>	3744	<b>85%</b>			<b>85%</b>	81%	87%	<b>84%</b>

SUPPORT FROM A MAIN CONTACT PERSON	Unadjusted scores						Case mix adjusted scores			England score
	2022 n	2022 score	2023 n	2023 score	Change 2022-2023	Change overall	2023 score	Lower expected range	Upper expected range	
Q17. Patient had a main point of contact within the care team	4119	<b>92%</b>	4024	<b>93%</b>			<b>92%</b>	90%	93%	<b>91%</b>
Q18. Patient found it very or quite easy to contact their main contact person	3407	<b>87%</b>	3388	<b>87%</b>			<b>87%</b>	80%	88%	<b>84%</b>
Q19. Patient found advice from main contact person was very or quite helpful	3583	<b>96%</b>	3527	<b>95%</b>			<b>95%</b>	95%	97%	<b>96%</b>




## Comparability tables

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

\*\* No score available for 2022.

▲ or ▼

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

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

DECIDING ON THE BEST TREATMENT	Unadjusted scores						Case mix adjusted scores			England score
	2022 n	2022 score	2023 n	2023 score	Change 2022-2023	Change overall	2023 score	Lower expected range	Upper expected range	
Q20. Treatment options were explained in a way the patient could completely understand	4028	84%	3965	83%			83%	80%	85%	82%
Q21. Patient was definitely involved as much as they wanted to be in decisions about their treatment	4224	82%	4167	81%			82%	77%	82%	80%
Q22. Family and/or carers were definitely involved as much as the patient wanted them to be in decisions about treatment options	3650	82%	3696	83%		▲	84%	82%	85%	83%
Q23. Patient could get further advice from a different healthcare professional before making decisions about their treatment options	**	**	2298	60%			59%	53%	61%	57%

CARE PLANNING	Unadjusted scores						Case mix adjusted scores			England score
	2022 n	2022 score	2023 n	2023 score	Change 2022-2023	Change overall	2023 score	Lower expected range	Upper expected range	
Q24. Patient was definitely able to have a discussion about their needs or concerns prior to treatment	3846	75%	3827	74%			74%	69%	75%	72%
Q25. A member of their care team helped the patient create a care plan to address any needs or concerns	2351	94%	2339	95%			95%	92%	95%	94%
Q26. Care team reviewed the patient's care plan with them to ensure it was up to date	1947	99%	1930	100%			100%	98%	99%	99%

SUPPORT FROM HOSPITAL STAFF	Unadjusted scores						Case mix adjusted scores			England score
	2022 n	2022 score	2023 n	2023 score	Change 2022-2023	Change overall	2023 score	Lower expected range	Upper expected range	
Q27. Staff provided the patient with relevant information on available support	3466	88%	3456	90%	▲		90%	89%	93%	91%
Q28. Patient definitely got the right level of support for their overall health and well being from hospital staff	4225	78%	4168	79%			79%	72%	79%	76%
Q29. Patient was offered information about how to get financial help or benefits	2502	68%	2528	72%	▲	▲	72%	66%	75%	70%

## Comparability tables

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

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

HOSPITAL CARE	Unadjusted scores						Case mix adjusted scores			England score
	2022 n	2022 score	2023 n	2023 score	Change 2022-2023	Change overall	2023 score	Lower expected range	Upper expected range	
Q31. Patient had confidence and trust in all of the team looking after them during their stay in hospital	1708	81%	1587	78%		▼	78%	75%	79%	77%
Q32. Patient's family, or someone close, was definitely able to talk to a member of the team looking after the patient in hospital	1430	71%	1337	73%		▲	73%	67%	72%	70%
Q33. Patient was always involved in decisions about their care and treatment whilst in hospital	1675	72%	1555	72%			72%	68%	73%	70%
Q34. Patient was always able to get help from ward staff when needed	1674	76%	1549	74%		▼	74%	70%	76%	73%
Q35. Patient was always able to discuss worries and fears with hospital staff	1632	69%	1516	68%			67%	62%	68%	65%
Q36. Hospital staff always did everything they could to help the patient control pain	1508	86%	1384	84%			85%	82%	86%	84%
Q37. Patient was always treated with respect and dignity while in hospital	1699	89%	1585	88%			89%	85%	89%	87%
Q38. Patient received easily understandable information about what they should or should not do after leaving hospital	1650	89%	1545	88%			88%	87%	90%	88%
Q39. Patient was always able to discuss worries and fears with hospital staff while being treated as an outpatient or day case	3742	81%	3761	83%			83%	75%	83%	79%

YOUR TREATMENT	Unadjusted scores						Case mix adjusted scores			England score
	2022 n	2022 score	2023 n	2023 score	Change 2022-2023	Change overall	2023 score	Lower expected range	Upper expected range	
Q41_1. Beforehand patient completely had enough understandable information about surgery	2166	91%	1865	90%			90%	88%	91%	90%
Q41_2. Beforehand patient completely had enough understandable information about chemotherapy	2286	89%	2291	88%			88%	84%	87%	86%
Q41_3. Beforehand patient completely had enough understandable information about radiotherapy	1251	89%	1070	90%			90%	87%	91%	89%
Q41_4. Beforehand patient completely had enough understandable information about hormone therapy	586	83%	642	82%			82%	76%	82%	79%
Q41_5. Beforehand patient completely had enough understandable information about immunotherapy	603	85%	731	85%			85%	81%	86%	84%
Q42_1. Patient completely had enough understandable information about their response to surgery	**	**	1837	87%			87%	85%	88%	86%
Q42_2. Patient completely had enough understandable information about their response to chemotherapy	**	**	2297	84%			84%	79%	83%	81%
Q42_3. Patient completely had enough understandable information about their response to radiotherapy	**	**	1062	86%			86%	83%	87%	85%
Q42_4. Patient completely had enough understandable information about their response to hormone therapy	**	**	636	82%			81%	73%	80%	76%
Q42_5. Patient completely had enough understandable information about their response to immunotherapy	**	**	727	84%			84%	78%	83%	81%
Q43. Patient felt the length of waiting time at clinic and day unit for cancer treatment was about right	4171	82%	4042	82%			82%	72%	85%	78%



## Comparability tables

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

\*\* No score available for 2022.

▲ or ▼

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

<span style="background-color: #4F81BD; width: 20px; height: 10px; display: inline-block;"></span>	Adjusted score below lower expected range
<span style="background-color: #D9D9D9; width: 20px; height: 10px; display: inline-block;"></span>	Adjusted score between upper and lower expected ranges
<span style="background-color: #0070C0; width: 20px; height: 10px; display: inline-block;"></span>	Adjusted score above upper expected range

IMMEDIATE AND LONG TERM SIDE EFFECTS	Unadjusted scores						Case mix adjusted scores			England score
	2022 n	2022 score	2023 n	2023 score	Change 2022-2023	Change overall	2023 score	Lower expected range	Upper expected range	
Q44. Possible side effects from treatment were definitely explained in a way the patient could understand	4064	<b>77%</b>	3973	<b>77%</b>			<b>77%</b>	73%	76%	<b>74%</b>
Q45. Patient was always offered practical advice on dealing with any immediate side effects from treatment	3907	<b>73%</b>	3805	<b>73%</b>			<b>73%</b>	67%	74%	<b>70%</b>
Q46. Patient was given information that they could access about support in dealing with immediate side effects from treatment	3126	<b>87%</b>	3067	<b>89%</b>		▲	<b>89%</b>	84%	90%	<b>87%</b>
Q47. Patient felt possible long-term side effects were definitely explained in a way they could understand in advance of their treatment	3811	<b>63%</b>	3791	<b>64%</b>			<b>63%</b>	57%	63%	<b>60%</b>
Q48. Patient was definitely able to discuss options for managing the impact of any long-term side effects	3357	<b>58%</b>	3332	<b>60%</b>			<b>60%</b>	51%	58%	<b>55%</b>

SUPPORT WHILE AT HOME	Unadjusted scores						Case mix adjusted scores			England score
	2022 n	2022 score	2023 n	2023 score	Change 2022-2023	Change overall	2023 score	Lower expected range	Upper expected range	
Q49. Care team gave family, or someone close, all the information needed to help care for the patient at home	2909	<b>61%</b>	2834	<b>63%</b>		▲	<b>63%</b>	58%	66%	<b>62%</b>
Q50. During treatment, the patient definitely got enough care and support at home from community or voluntary services	1793	<b>56%</b>	1763	<b>58%</b>			<b>58%</b>	47%	58%	<b>52%</b>

CARE FROM YOUR GP PRACTICE	Unadjusted scores						Case mix adjusted scores			England score
	2022 n	2022 score	2023 n	2023 score	Change 2022-2023	Change overall	2023 score	Lower expected range	Upper expected range	
Q51. Patient definitely received the right amount of support from their GP practice during treatment	2662	<b>49%</b>	2631	<b>49%</b>			<b>50%</b>	43%	50%	<b>46%</b>
Q52. Patient has had a review of cancer care by GP practice	4036	<b>25%</b>	3932	<b>23%</b>		▲	<b>24%</b>	20%	25%	<b>23%</b>

LIVING WITH AND BEYOND CANCER	Unadjusted scores						Case mix adjusted scores			England score
	2022 n	2022 score	2023 n	2023 score	Change 2022-2023	Change overall	2023 score	Lower expected range	Upper expected range	
Q53. After treatment, the patient definitely could get enough emotional support at home from community or voluntary services	1002	<b>37%</b>	905	<b>39%</b>			<b>39%</b>	28%	37%	<b>32%</b>
Q54. The right amount of information and support was offered to the patient between final treatment and the follow up appointment	1819	<b>80%</b>	1642	<b>82%</b>			<b>82%</b>	76%	83%	<b>79%</b>
Q55. Patient was given enough information about the possibility and signs of cancer coming back or spreading	3281	<b>68%</b>	3220	<b>68%</b>			<b>68%</b>	59%	68%	<b>64%</b>

YOUR OVERALL NHS CARE	Unadjusted scores						Case mix adjusted scores			England score
	2022 n	2022 score	2023 n	2023 score	Change 2022-2023	Change overall	2023 score	Lower expected range	Upper expected range	
Q56. The whole care team worked well together	4082	<b>90%</b>	3953	<b>92%</b>			<b>92%</b>	88%	92%	<b>90%</b>
Q57. Administration of care was very good or good	4198	<b>89%</b>	4129	<b>90%</b>			<b>90%</b>	85%	89%	<b>87%</b>
Q58. Cancer research opportunities were discussed with patient	2382	<b>40%</b>	2367	<b>43%</b>			<b>42%</b>	36%	53%	<b>45%</b>
Q59. Patient's average rating of care scored from very poor to very good	4088	<b>9.0</b>	4063	<b>9.0</b>			<b>9.0</b>	8.8	9.0	<b>8.9</b>

## Tumour group tables

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

	Tumour group													
	Brain / CNS	Breast	Colorectal / LGT	Gynaecological	Haematological	Head and neck	Lung	Prostate	Sarcoma	Skin	Upper gastro	Urological	Other	All cancers
Q2. Patient only spoke to primary care professional once or twice before cancer diagnosis	*	92%	78%	71%	65%	76%	69%	80%	*	90%	79%	78%	75%	<b>78%</b>
Q3. Referral for diagnosis was explained in a way the patient could completely understand	*	81%	71%	68%	57%	66%	50%	69%	*	75%	53%	59%	64%	<b>66%</b>

	Tumour group													
	Brain / CNS	Breast	Colorectal / LGT	Gynaecological	Haematological	Head and neck	Lung	Prostate	Sarcoma	Skin	Upper gastro	Urological	Other	All cancers
Q5. Patient received all the information needed about the diagnostic test in advance	*	93%	94%	84%	96%	92%	93%	94%	92%	95%	92%	92%	94%	<b>93%</b>
Q6. Diagnostic test staff appeared to completely have all the information they needed about the patient	*	86%	88%	76%	86%	83%	82%	90%	67%	83%	80%	85%	80%	<b>85%</b>
Q7. Patient felt the length of time waiting for diagnostic test results was about right	*	82%	84%	81%	87%	80%	81%	82%	80%	69%	72%	79%	76%	<b>81%</b>
Q8. Diagnostic test results were explained in a way the patient could completely understand	*	82%	82%	73%	84%	79%	81%	78%	73%	87%	75%	78%	73%	<b>80%</b>
Q9. Enough privacy was always given to the patient when receiving diagnostic test results	*	96%	95%	94%	94%	94%	95%	98%	87%	95%	97%	89%	95%	<b>95%</b>

	Tumour group													
	Brain / CNS	Breast	Colorectal / LGT	Gynaecological	Haematological	Head and neck	Lung	Prostate	Sarcoma	Skin	Upper gastro	Urological	Other	All cancers
Q12. Patient was told they could have a family member, carer or friend with them when told diagnosis	70%	87%	87%	83%	78%	87%	83%	82%	83%	75%	89%	75%	76%	<b>82%</b>
Q13. Patient was definitely told sensitively that they had cancer	60%	80%	77%	75%	75%	75%	76%	72%	78%	76%	79%	65%	75%	<b>75%</b>
Q14. Cancer diagnosis explained in a way the patient could completely understand	36%	78%	80%	78%	74%	83%	75%	77%	67%	82%	76%	74%	69%	<b>76%</b>
Q15. Patient was definitely told about their diagnosis in an appropriate place	82%	89%	86%	85%	84%	88%	85%	86%	89%	89%	85%	77%	79%	<b>85%</b>
Q16. Patient was told they could go back later for more information about their diagnosis	*	89%	85%	87%	85%	84%	82%	84%	94%	90%	82%	78%	81%	<b>85%</b>

## Tumour group tables

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

	Tumour group													
	Brain/ CNS	Breast	Colorectal/ LGT	Gynaecological	Haematological	Head and neck	Lung	Prostate	Sarcoma	Skin	Upper gastro	Urological	Other	All cancers
Q17. Patient had a main point of contact within the care team	100%	96%	91%	94%	93%	94%	96%	91%	82%	81%	93%	87%	89%	<b>93%</b>
Q18. Patient found it very or quite easy to contact their main contact person	82%	87%	89%	88%	88%	88%	85%	85%	71%	98%	81%	84%	87%	<b>87%</b>
Q19. Patient found advice from main contact person was very or quite helpful	100%	94%	96%	97%	98%	95%	94%	96%	100%	98%	94%	93%	96%	<b>95%</b>

	Tumour group													
	Brain/ CNS	Breast	Colorectal/ LGT	Gynaecological	Haematological	Head and neck	Lung	Prostate	Sarcoma	Skin	Upper gastro	Urological	Other	All cancers
Q20. Treatment options were explained in a way the patient could completely understand	90%	84%	83%	85%	86%	79%	83%	79%	71%	90%	85%	82%	80%	<b>83%</b>
Q21. Patient was definitely involved as much as they wanted to be in decisions about their treatment	73%	83%	82%	80%	84%	81%	83%	80%	78%	87%	81%	78%	76%	<b>81%</b>
Q22. Family and/or carers were definitely involved as much as the patient wanted them to be in decisions about treatment options	*	82%	87%	84%	86%	87%	86%	80%	94%	91%	83%	79%	78%	<b>83%</b>
Q23. Patient could get further advice from a different healthcare professional before making decisions about their treatment options	*	63%	59%	50%	59%	61%	63%	63%	57%	62%	64%	60%	53%	<b>60%</b>

	Tumour group													
	Brain/ CNS	Breast	Colorectal/ LGT	Gynaecological	Haematological	Head and neck	Lung	Prostate	Sarcoma	Skin	Upper gastro	Urological	Other	All cancers
Q24. Patient was definitely able to have a discussion about their needs or concerns prior to treatment	82%	75%	76%	75%	76%	76%	75%	68%	71%	87%	74%	68%	72%	<b>74%</b>
Q25. A member of their care team helped the patient create a care plan to address any needs or concerns	*	95%	93%	94%	96%	97%	97%	93%	100%	92%	97%	90%	93%	<b>95%</b>
Q26. Care team reviewed the patient's care plan with them to ensure it was up to date	*	100%	100%	99%	100%	100%	99%	99%	*	100%	100%	100%	99%	<b>100%</b>

	Tumour group													
	Brain/ CNS	Breast	Colorectal/ LGT	Gynaecological	Haematological	Head and neck	Lung	Prostate	Sarcoma	Skin	Upper gastro	Urological	Other	All cancers
Q27. Staff provided the patient with relevant information on available support	*	90%	91%	88%	93%	92%	90%	89%	81%	95%	93%	81%	86%	<b>90%</b>
Q28. Patient definitely got the right level of support for their overall health and well being from hospital staff	73%	77%	79%	76%	84%	79%	77%	78%	67%	87%	82%	76%	75%	<b>79%</b>
Q29. Patient was offered information about how to get financial help or benefits	*	73%	74%	78%	75%	75%	83%	58%	58%	88%	77%	56%	65%	<b>72%</b>

## Tumour group tables

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

	Tumour group													
	Brain/ CNS	Breast	Colorectal/ LGT	Gynaecological	Haematological	Head and neck	Lung	Prostate	Sarcoma	Skin	Upper gastro	Urological	Other	All cancers
Q31. Patient had confidence and trust in all of the team looking after them during their stay in hospital	*	74%	77%	88%	79%	84%	82%	82%	*	*	73%	81%	74%	<b>78%</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	*	68%	72%	74%	77%	73%	77%	76%	*	*	77%	69%	74%	<b>73%</b>
Q33. Patient was always involved in decisions about their care and treatment whilst in hospital	*	70%	71%	79%	71%	77%	62%	73%	*	*	72%	75%	74%	<b>72%</b>
Q34. Patient was always able to get help from ward staff when needed	*	72%	71%	75%	77%	76%	79%	77%	*	*	76%	75%	73%	<b>74%</b>
Q35. Patient was always able to discuss worries and fears with hospital staff	*	62%	68%	67%	69%	70%	70%	65%	*	*	71%	71%	66%	<b>68%</b>
Q36. Hospital staff always did everything they could to help the patient control pain	*	85%	83%	86%	83%	84%	88%	86%	*	*	87%	82%	84%	<b>84%</b>
Q37. Patient was always treated with respect and dignity while in hospital	*	85%	86%	95%	91%	86%	92%	90%	*	*	89%	88%	88%	<b>88%</b>
Q38. Patient received easily understandable information about what they should or should not do after leaving hospital	*	88%	89%	89%	90%	87%	89%	84%	*	*	89%	89%	85%	<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	70%	81%	83%	81%	88%	81%	86%	83%	69%	94%	85%	78%	79%	<b>83%</b>

	Tumour group													
	Brain/ CNS	Breast	Colorectal/ LGT	Gynaecological	Haematological	Head and neck	Lung	Prostate	Sarcoma	Skin	Upper gastro	Urological	Other	All cancers
Q41_1. Beforehand patient completely had enough understandable information about surgery	*	92%	91%	91%	87%	92%	89%	88%	83%	96%	89%	87%	88%	<b>90%</b>
Q41_2. Beforehand patient completely had enough understandable information about chemotherapy	*	88%	88%	86%	91%	85%	90%	90%	82%	*	90%	83%	86%	<b>88%</b>
Q41_3. Beforehand patient completely had enough understandable information about radiotherapy	*	91%	82%	91%	89%	86%	92%	87%	*	*	96%	90%	90%	<b>90%</b>
Q41_4. Beforehand patient completely had enough understandable information about hormone therapy	*	80%	*	*	*	*	*	86%	*	*	*	*	86%	<b>82%</b>
Q41_5. Beforehand patient completely had enough understandable information about immunotherapy	*	75%	78%	*	90%	100%	88%	100%	*	73%	83%	90%	81%	<b>85%</b>
Q42_1. Patient completely had enough understandable information about their response to surgery	*	90%	89%	85%	84%	89%	82%	87%	80%	91%	90%	83%	85%	<b>87%</b>
Q42_2. Patient completely had enough understandable information about their response to chemotherapy	*	85%	82%	78%	87%	85%	88%	82%	82%	*	86%	79%	81%	<b>84%</b>
Q42_3. Patient completely had enough understandable information about their response to radiotherapy	*	89%	75%	86%	84%	83%	83%	87%	*	*	92%	83%	87%	<b>86%</b>
Q42_4. Patient completely had enough understandable information about their response to hormone therapy	*	77%	*	*	64%	*	*	88%	*	*	*	*	85%	<b>82%</b>
Q42_5. Patient completely had enough understandable information about their response to immunotherapy	*	77%	74%	*	89%	86%	85%	91%	*	93%	82%	88%	80%	<b>84%</b>
Q43. Patient felt the length of waiting time at clinic and day unit for cancer treatment was about right	82%	80%	85%	77%	84%	88%	85%	92%	56%	93%	79%	83%	72%	<b>82%</b>

## Tumour group tables

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

	Tumour group													
	Brain / CNS	Breast	Colorectal / LGT	Gynaecological	Haematological	Head and neck	Lung	Prostate	Sarcoma	Skin	Upper gastro	Urological	Other	All cancers
Q44. Possible side effects from treatment were definitely explained in a way the patient could understand	64%	77%	79%	72%	78%	77%	80%	75%	81%	80%	78%	78%	74%	<b>77%</b>
Q45. Patient was always offered practical advice on dealing with any immediate side effects from treatment	64%	75%	75%	73%	75%	77%	75%	67%	83%	84%	79%	68%	69%	<b>73%</b>
Q46. Patient was given information that they could access about support in dealing with immediate side effects from treatment	82%	88%	91%	92%	91%	89%	88%	86%	100%	100%	88%	83%	86%	<b>89%</b>
Q47. Patient felt possible long-term side effects were definitely explained in a way they could understand in advance of their treatment	73%	64%	64%	59%	63%	66%	66%	69%	50%	71%	66%	60%	57%	<b>64%</b>
Q48. Patient was definitely able to discuss options for managing the impact of any long-term side effects	50%	60%	58%	55%	65%	65%	62%	62%	54%	82%	58%	56%	55%	<b>60%</b>

	Tumour group													
	Brain / CNS	Breast	Colorectal / LGT	Gynaecological	Haematological	Head and neck	Lung	Prostate	Sarcoma	Skin	Upper gastro	Urological	Other	All cancers
Q49. Care team gave family, or someone close, all the information needed to help care for the patient at home	60%	58%	68%	60%	68%	72%	67%	55%	69%	70%	65%	63%	62%	<b>63%</b>
Q50. During treatment, the patient definitely got enough care and support at home from community or voluntary services	*	59%	57%	53%	59%	61%	63%	48%	*	62%	61%	58%	59%	<b>58%</b>

	Tumour group													
	Brain / CNS	Breast	Colorectal / LGT	Gynaecological	Haematological	Head and neck	Lung	Prostate	Sarcoma	Skin	Upper gastro	Urological	Other	All cancers
Q51. Patient definitely received the right amount of support from their GP practice during treatment	*	54%	48%	46%	49%	48%	41%	54%	25%	68%	44%	47%	44%	<b>49%</b>
Q52. Patient has had a review of cancer care by GP practice	27%	24%	26%	31%	19%	21%	26%	26%	19%	19%	26%	22%	21%	<b>23%</b>

## Tumour group tables

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

	Tumour group													
	Brain / CNS	Breast	Colorectal / LGT	Gynaecological	Haematological	Head and neck	Lung	Prostate	Sarcoma	Skin	Upper gastro	Urological	Other	All cancers
Q53. After treatment, the patient definitely could get enough emotional support at home from community or voluntary services	*	35%	40%	50%	36%	46%	42%	35%	*	*	50%	35%	33%	<b>39%</b>
Q54. The right amount of information and support was offered to the patient between final treatment and the follow up appointment	*	81%	82%	84%	87%	86%	74%	70%	*	88%	87%	78%	84%	<b>82%</b>
Q55. Patient was given enough information about the possibility and signs of cancer coming back or spreading	60%	59%	69%	61%	81%	64%	65%	74%	69%	85%	68%	73%	63%	<b>68%</b>

	Tumour group													
	Brain / CNS	Breast	Colorectal / LGT	Gynaecological	Haematological	Head and neck	Lung	Prostate	Sarcoma	Skin	Upper gastro	Urological	Other	All cancers
Q56. The whole care team worked well together	91%	92%	91%	90%	93%	95%	90%	92%	94%	99%	95%	93%	88%	<b>92%</b>
Q57. Administration of care was very good or good	91%	90%	90%	88%	94%	87%	87%	89%	78%	88%	88%	87%	87%	<b>90%</b>
Q58. Cancer research opportunities were discussed with patient	*	31%	41%	43%	47%	48%	51%	49%	54%	42%	52%	43%	41%	<b>43%</b>
Q59. Patient's average rating of care scored from very poor to very good	8.8	9.1	8.9	8.9	9.2	9.0	9.0	8.9	8.4	9.4	8.9	8.9	8.8	<b>9.0</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	*	84%	76%	80%	74%	78%	81%	78%	<b>78%</b>
Q3. Referral for diagnosis was explained in a way the patient could completely understand	*	67%	72%	80%	66%	64%	63%	64%	<b>66%</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	*	87%	86%	94%	94%	93%	93%	95%	<b>93%</b>
Q6. Diagnostic test staff appeared to completely have all the information they needed about the patient	*	67%	86%	84%	84%	86%	83%	87%	<b>85%</b>
Q7. Patient felt the length of time waiting for diagnostic test results was about right	*	67%	70%	71%	79%	82%	85%	90%	<b>81%</b>
Q8. Diagnostic test results were explained in a way the patient could completely understand	*	67%	69%	77%	78%	82%	80%	81%	<b>80%</b>
Q9. Enough privacy was always given to the patient when receiving diagnostic test results	*	87%	92%	93%	94%	95%	96%	98%	<b>95%</b>

<b>FINDING OUT THAT YOU HAD CANCER</b>	Age								
	16 - 24	25 - 34	35 - 44	45 - 54	55 - 64	65 - 74	75 - 84	85+	All
Q12. Patient was told they could have a family member, carer or friend with them when told diagnosis	67%	83%	85%	80%	80%	82%	84%	86%	<b>82%</b>
Q13. Patient was definitely told sensitively that they had cancer	58%	71%	67%	73%	72%	75%	79%	84%	<b>75%</b>
Q14. Cancer diagnosis explained in a way the patient could completely understand	58%	77%	73%	76%	74%	75%	78%	77%	<b>76%</b>
Q15. Patient was definitely told about their diagnosis in an appropriate place	75%	77%	75%	86%	81%	84%	89%	90%	<b>85%</b>
Q16. Patient was told they could go back later for more information about their diagnosis	92%	90%	87%	87%	87%	85%	82%	79%	<b>85%</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	91%	88%	87%	94%	93%	92%	92%	93%	<b>93%</b>
Q18. Patient found it very or quite easy to contact their main contact person	*	89%	87%	84%	87%	87%	87%	88%	<b>87%</b>
Q19. Patient found advice from main contact person was very or quite helpful	*	93%	96%	94%	95%	95%	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	82%	84%	79%	83%	83%	84%	84%	80%	<b>83%</b>
Q21. Patient was definitely involved as much as they wanted to be in decisions about their treatment	75%	80%	65%	75%	81%	83%	84%	80%	<b>81%</b>
Q22. Family and/or carers were definitely involved as much as the patient wanted them to be in decisions about treatment options	82%	79%	70%	79%	83%	84%	86%	83%	<b>83%</b>
Q23. Patient could get further advice from a different healthcare professional before making decisions about their treatment options	*	56%	58%	62%	60%	61%	57%	58%	<b>60%</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	73%	70%	67%	75%	73%	75%	75%	68%	<b>74%</b>
Q25. A member of their care team helped the patient create a care plan to address any needs or concerns	*	95%	98%	94%	94%	94%	96%	95%	<b>95%</b>
Q26. Care team reviewed the patient's care plan with them to ensure it was up to date	*	100%	98%	99%	100%	99%	100%	100%	<b>100%</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	100%	94%	90%	88%	90%	90%	90%	87%	<b>90%</b>
Q28. Patient definitely got the right level of support for their overall health and well being from hospital staff	83%	69%	68%	75%	75%	80%	82%	84%	<b>79%</b>
Q29. Patient was offered information about how to get financial help or benefits	91%	81%	67%	74%	75%	72%	68%	69%	<b>72%</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	80%	81%	71%	67%	74%	82%	83%	85%	<b>78%</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	90%	80%	65%	62%	73%	75%	76%	75%	<b>73%</b>
Q33. Patient was always involved in decisions about their care and treatment whilst in hospital	*	81%	73%	64%	69%	73%	74%	79%	<b>72%</b>
Q34. Patient was always able to get help from ward staff when needed	80%	69%	68%	63%	71%	77%	78%	80%	<b>74%</b>
Q35. Patient was always able to discuss worries and fears with hospital staff	90%	67%	56%	56%	66%	72%	70%	62%	<b>68%</b>
Q36. Hospital staff always did everything they could to help the patient control pain	80%	71%	83%	73%	84%	87%	86%	98%	<b>84%</b>
Q37. Patient was always treated with respect and dignity while in hospital	90%	88%	88%	79%	87%	91%	91%	92%	<b>88%</b>
Q38. Patient received easily understandable information about what they should or should not do after leaving hospital	*	81%	93%	83%	88%	89%	89%	85%	<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	90%	84%	69%	77%	82%	84%	85%	85%	<b>83%</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	*	89%	85%	89%	90%	92%	91%	89%	<b>90%</b>
Q41_2. Beforehand patient completely had enough understandable information about chemotherapy	90%	88%	84%	87%	88%	89%	90%	88%	<b>88%</b>
Q41_3. Beforehand patient completely had enough understandable information about radiotherapy	*	91%	83%	88%	92%	92%	87%	81%	<b>90%</b>
Q41_4. Beforehand patient completely had enough understandable information about hormone therapy	*	*	54%	77%	82%	88%	84%	85%	<b>82%</b>
Q41_5. Beforehand patient completely had enough understandable information about immunotherapy	*	*	67%	84%	80%	89%	89%	76%	<b>85%</b>
Q42_1. Patient completely had enough understandable information about their response to surgery	*	80%	80%	85%	87%	88%	89%	90%	<b>87%</b>
Q42_2. Patient completely had enough understandable information about their response to chemotherapy	91%	79%	71%	86%	83%	86%	85%	83%	<b>84%</b>
Q42_3. Patient completely had enough understandable information about their response to radiotherapy	*	82%	74%	86%	88%	88%	83%	81%	<b>86%</b>
Q42_4. Patient completely had enough understandable information about their response to hormone therapy	*	*	56%	79%	80%	86%	83%	79%	<b>82%</b>
Q42_5. Patient completely had enough understandable information about their response to immunotherapy	*	*	76%	81%	78%	88%	86%	80%	<b>84%</b>
Q43. Patient felt the length of waiting time at clinic and day unit for cancer treatment was about right	55%	59%	71%	79%	80%	83%	86%	84%	<b>82%</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	92%	69%	79%	76%	79%	77%	76%	74%	<b>77%</b>
Q45. Patient was always offered practical advice on dealing with any immediate side effects from treatment	83%	72%	63%	73%	74%	75%	73%	65%	<b>73%</b>
Q46. Patient was given information that they could access about support in dealing with immediate side effects from treatment	*	90%	82%	89%	90%	89%	88%	82%	<b>89%</b>
Q47. Patient felt possible long-term side effects were definitely explained in a way they could understand in advance of their treatment	75%	63%	62%	63%	63%	65%	62%	64%	<b>64%</b>
Q48. Patient was definitely able to discuss options for managing the impact of any long-term side effects	67%	52%	54%	58%	60%	62%	59%	58%	<b>60%</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	80%	63%	54%	59%	63%	64%	65%	68%	<b>63%</b>
Q50. During treatment, the patient definitely got enough care and support at home from community or voluntary services	*	77%	54%	61%	56%	58%	60%	59%	<b>58%</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	*	56%	55%	56%	50%	45%	49%	57%	<b>49%</b>
Q52. Patient has had a review of cancer care by GP practice	33%	29%	27%	33%	24%	22%	22%	22%	<b>23%</b>

## Age group tables

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

	Age								
	16 - 24	25 - 34	35 - 44	45 - 54	55 - 64	65 - 74	75 - 84	85+	All
Q53. After treatment, the patient definitely could get enough emotional support at home from community or voluntary services	*	55%	38%	38%	38%	37%	39%	56%	<b>39%</b>
Q54. The right amount of information and support was offered to the patient between final treatment and the follow up appointment	*	88%	75%	73%	83%	83%	83%	85%	<b>82%</b>
Q55. Patient was given enough information about the possibility and signs of cancer coming back or spreading	73%	45%	55%	61%	65%	72%	71%	72%	<b>68%</b>

	Age								
	16 - 24	25 - 34	35 - 44	45 - 54	55 - 64	65 - 74	75 - 84	85+	All
Q56. The whole care team worked well together	92%	91%	90%	89%	91%	92%	93%	93%	<b>92%</b>
Q57. Administration of care was very good or good	83%	85%	84%	88%	88%	90%	92%	88%	<b>90%</b>
Q58. Cancer research opportunities were discussed with patient	*	35%	41%	36%	44%	45%	43%	35%	<b>43%</b>
Q59. Patient's average rating of care scored from very poor to very good	8.7	9.0	8.6	8.8	9.0	9.1	9.0	8.9	<b>9.0</b>

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

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

SUPPORT FROM YOUR GP PRACTICE							
	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	80%	77%	*	*	*	73%	78%
Q3. Referral for diagnosis was explained in a way the patient could completely understand	70%	61%	*	*	*	65%	66%

DIAGNOSTIC TESTS							
	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	93%	93%	*	*	*	92%	93%
Q6. Diagnostic test staff appeared to completely have all the information they needed about the patient	83%	87%	*	*	*	82%	85%
Q7. Patient felt the length of time waiting for diagnostic test results was about right	80%	82%	*	*	*	83%	81%
Q8. Diagnostic test results were explained in a way the patient could completely understand	79%	81%	*	*	*	82%	80%
Q9. Enough privacy was always given to the patient when receiving diagnostic test results	94%	96%	*	*	*	96%	95%

FINDING OUT THAT YOU HAD CANCER							
	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	82%	82%	*	*	*	87%	82%
Q13. Patient was definitely told sensitively that they had cancer	76%	74%	*	*	*	82%	75%
Q14. Cancer diagnosis explained in a way the patient could completely understand	74%	77%	*	*	*	78%	76%
Q15. Patient was definitely told about their diagnosis in an appropriate place	84%	86%	*	*	*	89%	85%
Q16. Patient was told they could go back later for more information about their diagnosis	85%	84%	*	*	*	85%	85%

SUPPORT FROM A MAIN CONTACT PERSON							
	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%	92%	*	*	*	93%	93%
Q18. Patient found it very or quite easy to contact their main contact person	87%	86%	*	*	*	86%	87%
Q19. Patient found advice from main contact person was very or quite helpful	95%	96%	*	*	*	96%	95%

## 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	83%	83%	*	*	*	85%	<b>83%</b>
Q21. Patient was definitely involved as much as they wanted to be in decisions about their treatment	81%	82%	*	*	*	83%	<b>81%</b>
Q22. Family and/or carers were definitely involved as much as the patient wanted them to be in decisions about treatment options	83%	84%	*	*	*	85%	<b>83%</b>
Q23. Patient could get further advice from a different healthcare professional before making decisions about their treatment options	61%	59%	*	*	*	56%	<b>60%</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	74%	74%	*	*	*	75%	<b>74%</b>
Q25. A member of their care team helped the patient create a care plan to address any needs or concerns	94%	95%	*	*	*	95%	<b>95%</b>
Q26. Care team reviewed the patient's care plan with them to ensure it was up to date	99%	100%	*	*	*	100%	<b>100%</b>

	Male/Female/Non-binary/Other						
	Female	Male	Non-binary	Prefer to self-describe	Prefer not to say	Not given	All
Q27. Staff provided the patient with relevant information on available support	89%	91%	*	*	*	90%	<b>90%</b>
Q28. Patient definitely got the right level of support for their overall health and well being from hospital staff	77%	81%	*	*	*	77%	<b>79%</b>
Q29. Patient was offered information about how to get financial help or benefits	74%	71%	*	*	*	73%	<b>72%</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	77%	80%	*	*	*	79%	<b>78%</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	74%	74%	*	*	*	62%	<b>73%</b>
Q33. Patient was always involved in decisions about their care and treatment whilst in hospital	72%	72%	*	*	*	72%	<b>72%</b>
Q34. Patient was always able to get help from ward staff when needed	71%	77%	*	*	*	80%	<b>74%</b>
Q35. Patient was always able to discuss worries and fears with hospital staff	65%	71%	*	*	*	65%	<b>68%</b>
Q36. Hospital staff always did everything they could to help the patient control pain	84%	85%	*	*	*	83%	<b>84%</b>
Q37. Patient was always treated with respect and dignity while in hospital	87%	90%	*	*	*	86%	<b>88%</b>
Q38. Patient received easily understandable information about what they should or should not do after leaving hospital	88%	89%	*	*	*	83%	<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	82%	85%	*	*	*	83%	<b>83%</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	91%	89%	*	*	*	90%	<b>90%</b>
Q41_2. Beforehand patient completely had enough understandable information about chemotherapy	88%	90%	*	*	*	87%	<b>88%</b>
Q41_3. Beforehand patient completely had enough understandable information about radiotherapy	91%	89%	*	*	*	84%	<b>90%</b>
Q41_4. Beforehand patient completely had enough understandable information about hormone therapy	80%	87%	*	*	*	77%	<b>82%</b>
Q41_5. Beforehand patient completely had enough understandable information about immunotherapy	83%	89%	*	*	*	84%	<b>85%</b>
Q42_1. Patient completely had enough understandable information about their response to surgery	88%	86%	*	*	*	88%	<b>87%</b>
Q42_2. Patient completely had enough understandable information about their response to chemotherapy	84%	85%	*	*	*	86%	<b>84%</b>
Q42_3. Patient completely had enough understandable information about their response to radiotherapy	87%	86%	*	*	*	85%	<b>86%</b>
Q42_4. Patient completely had enough understandable information about their response to hormone therapy	77%	88%	*	*	*	81%	<b>82%</b>
Q42_5. Patient completely had enough understandable information about their response to immunotherapy	83%	86%	*	*	*	82%	<b>84%</b>
Q43. Patient felt the length of waiting time at clinic and day unit for cancer treatment was about right	80%	86%	*	*	*	81%	<b>82%</b>

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

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

<b>IMMEDIATE AND LONG TERM SIDE EFFECTS</b>							
	Male/Female/Non-binary/Other						
	Female	Male	Non-binary	Prefer to self-describe	Prefer not to say	Not given	All
Q44. Possible side effects from treatment were definitely explained in a way the patient could understand	76%	78%	*	*	*	73%	<b>77%</b>
Q45. Patient was always offered practical advice on dealing with any immediate side effects from treatment	73%	74%	*	*	*	72%	<b>73%</b>
Q46. Patient was given information that they could access about support in dealing with immediate side effects from treatment	89%	88%	*	*	*	87%	<b>89%</b>
Q47. Patient felt possible long-term side effects were definitely explained in a way they could understand in advance of their treatment	62%	66%	*	*	*	59%	<b>64%</b>
Q48. Patient was definitely able to discuss options for managing the impact of any long-term side effects	58%	64%	*	*	*	59%	<b>60%</b>

<b>SUPPORT WHILE AT HOME</b>							
	Male/Female/Non-binary/Other						
	Female	Male	Non-binary	Prefer to self-describe	Prefer not to say	Not given	All
Q49. Care team gave family, or someone close, all the information needed to help care for the patient at home	59%	69%	*	*	*	62%	<b>63%</b>
Q50. During treatment, the patient definitely got enough care and support at home from community or voluntary services	57%	60%	*	*	*	59%	<b>58%</b>

<b>CARE FROM YOUR GP PRACTICE</b>							
	Male/Female/Non-binary/Other						
	Female	Male	Non-binary	Prefer to self-describe	Prefer not to say	Not given	All
Q51. Patient definitely received the right amount of support from their GP practice during treatment	48%	49%	*	*	*	54%	<b>49%</b>
Q52. Patient has had a review of cancer care by GP practice	24%	23%	*	*	*	25%	<b>23%</b>

<b>LIVING WITH AND BEYOND CANCER</b>							
	Male/Female/Non-binary/Other						
	Female	Male	Non-binary	Prefer to self-describe	Prefer not to say	Not given	All
Q53. After treatment, the patient definitely could get enough emotional support at home from community or voluntary services	36%	43%	*	*	*	39%	<b>39%</b>
Q54. The right amount of information and support was offered to the patient between final treatment and the follow up appointment	81%	83%	*	*	*	78%	<b>82%</b>
Q55. Patient was given enough information about the possibility and signs of cancer coming back or spreading	64%	74%	*	*	*	63%	<b>68%</b>

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

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

YOUR OVERALL NHS CARE	Male/Female/Non-binary/Other						All
	Female	Male	Non-binary	Prefer to self-describe	Prefer not to say	Not given	
Q56. The whole care team worked well together	91%	93%	*	*	*	94%	<b>92%</b>
Q57. Administration of care was very good or good	88%	91%	*	*	*	93%	<b>90%</b>
Q58. Cancer research opportunities were discussed with patient	38%	50%	*	*	*	37%	<b>43%</b>
Q59. Patient's average rating of care scored from very poor to very good	9.0	9.0	*	*	*	9.0	<b>9.0</b>

## Ethnicity tables

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

SUPPORT FROM YOUR GP PRACTICE	Ethnicity						
	White	Mixed	Asian	Black	Other	Not given	All
Q2. Patient only spoke to primary care professional once or twice before cancer diagnosis	79%	79%	65%	*	*	77%	<b>78%</b>
Q3. Referral for diagnosis was explained in a way the patient could completely understand	66%	61%	62%	*	*	64%	<b>66%</b>

DIAGNOSTIC TESTS	Ethnicity						
	White	Mixed	Asian	Black	Other	Not given	All
Q5. Patient received all the information needed about the diagnostic test in advance	93%	94%	94%	90%	*	93%	<b>93%</b>
Q6. Diagnostic test staff appeared to completely have all the information they needed about the patient	85%	82%	92%	70%	*	80%	<b>85%</b>
Q7. Patient felt the length of time waiting for diagnostic test results was about right	81%	76%	85%	90%	*	82%	<b>81%</b>
Q8. Diagnostic test results were explained in a way the patient could completely understand	80%	89%	78%	90%	*	77%	<b>80%</b>
Q9. Enough privacy was always given to the patient when receiving diagnostic test results	95%	89%	98%	100%	*	95%	<b>95%</b>

FINDING OUT THAT YOU HAD CANCER	Ethnicity						
	White	Mixed	Asian	Black	Other	Not given	All
Q12. Patient was told they could have a family member, carer or friend with them when told diagnosis	82%	90%	88%	*	*	83%	<b>82%</b>
Q13. Patient was definitely told sensitively that they had cancer	75%	76%	84%	80%	*	78%	<b>75%</b>
Q14. Cancer diagnosis explained in a way the patient could completely understand	76%	68%	82%	90%	*	77%	<b>76%</b>
Q15. Patient was definitely told about their diagnosis in an appropriate place	85%	86%	93%	90%	*	88%	<b>85%</b>
Q16. Patient was told they could go back later for more information about their diagnosis	85%	89%	86%	80%	*	83%	<b>85%</b>

SUPPORT FROM A MAIN CONTACT PERSON	Ethnicity						
	White	Mixed	Asian	Black	Other	Not given	All
Q17. Patient had a main point of contact within the care team	93%	90%	95%	80%	*	92%	<b>93%</b>
Q18. Patient found it very or quite easy to contact their main contact person	87%	76%	86%	*	*	86%	<b>87%</b>
Q19. Patient found advice from main contact person was very or quite helpful	95%	89%	100%	*	*	95%	<b>95%</b>

DECIDING ON THE BEST TREATMENT	Ethnicity						
	White	Mixed	Asian	Black	Other	Not given	All
Q20. Treatment options were explained in a way the patient could completely understand	83%	86%	98%	90%	*	83%	<b>83%</b>
Q21. Patient was definitely involved as much as they wanted to be in decisions about their treatment	81%	81%	95%	100%	*	80%	<b>81%</b>
Q22. Family and/or carers were definitely involved as much as the patient wanted them to be in decisions about treatment options	83%	81%	86%	*	*	84%	<b>83%</b>
Q23. Patient could get further advice from a different healthcare professional before making decisions about their treatment options	60%	56%	75%	*	*	56%	<b>60%</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	74%	70%	88%	*	*	72%	74%
Q25. A member of their care team helped the patient create a care plan to address any needs or concerns	94%	100%	97%	*	*	96%	95%
Q26. Care team reviewed the patient's care plan with them to ensure it was up to date	100%	100%	100%	*	*	100%	100%

SUPPORT FROM HOSPITAL STAFF	Ethnicity						
	White	Mixed	Asian	Black	Other	Not given	All
Q27. Staff provided the patient with relevant information on available support	90%	95%	92%	100%	*	89%	90%
Q28. Patient definitely got the right level of support for their overall health and well being from hospital staff	79%	81%	84%	60%	*	76%	79%
Q29. Patient was offered information about how to get financial help or benefits	72%	84%	74%	*	*	69%	72%

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	78%	50%	93%	*	*	81%	78%
Q32. Patient's family, or someone close, was definitely able to talk to a member of the team looking after the patient in hospital	74%	*	78%	*	*	66%	73%
Q33. Patient was always involved in decisions about their care and treatment whilst in hospital	72%	60%	83%	*	*	74%	72%
Q34. Patient was always able to get help from ward staff when needed	74%	70%	79%	*	*	77%	74%
Q35. Patient was always able to discuss worries and fears with hospital staff	68%	50%	79%	*	*	61%	68%
Q36. Hospital staff always did everything they could to help the patient control pain	84%	60%	89%	*	*	83%	84%
Q37. Patient was always treated with respect and dignity while in hospital	88%	90%	93%	*	*	87%	88%
Q38. Patient received easily understandable information about what they should or should not do after leaving hospital	89%	*	90%	*	*	81%	88%
Q39. Patient was always able to discuss worries and fears with hospital staff while being treated as an outpatient or day case	83%	82%	80%	*	*	80%	83%

## Ethnicity tables

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

YOUR TREATMENT	Ethnicity						
	White	Mixed	Asian	Black	Other	Not given	All
Q41_1. Beforehand patient completely had enough understandable information about surgery	90%	92%	93%	*	*	88%	<b>90%</b>
Q41_2. Beforehand patient completely had enough understandable information about chemotherapy	89%	93%	97%	*	*	83%	<b>88%</b>
Q41_3. Beforehand patient completely had enough understandable information about radiotherapy	90%	*	100%	*	*	82%	<b>90%</b>
Q41_4. Beforehand patient completely had enough understandable information about hormone therapy	82%	*	*	*	*	81%	<b>82%</b>
Q41_5. Beforehand patient completely had enough understandable information about immunotherapy	85%	*	*	*	*	83%	<b>85%</b>
Q42_1. Patient completely had enough understandable information about their response to surgery	88%	82%	89%	*	*	86%	<b>87%</b>
Q42_2. Patient completely had enough understandable information about their response to chemotherapy	85%	87%	90%	*	*	80%	<b>84%</b>
Q42_3. Patient completely had enough understandable information about their response to radiotherapy	86%	*	100%	*	*	83%	<b>86%</b>
Q42_4. Patient completely had enough understandable information about their response to hormone therapy	81%	*	*	*	*	84%	<b>82%</b>
Q42_5. Patient completely had enough understandable information about their response to immunotherapy	84%	*	*	*	*	82%	<b>84%</b>
Q43. Patient felt the length of waiting time at clinic and day unit for cancer treatment was about right	83%	89%	75%	*	*	80%	<b>82%</b>

IMMEDIATE AND LONG TERM SIDE EFFECTS	Ethnicity						
	White	Mixed	Asian	Black	Other	Not given	All
Q44. Possible side effects from treatment were definitely explained in a way the patient could understand	77%	85%	82%	70%	*	72%	<b>77%</b>
Q45. Patient was always offered practical advice on dealing with any immediate side effects from treatment	74%	70%	88%	*	*	69%	<b>73%</b>
Q46. Patient was given information that they could access about support in dealing with immediate side effects from treatment	89%	89%	91%	*	*	88%	<b>89%</b>
Q47. Patient felt possible long-term side effects were definitely explained in a way they could understand in advance of their treatment	64%	62%	74%	70%	*	59%	<b>64%</b>
Q48. Patient was definitely able to discuss options for managing the impact of any long-term side effects	60%	62%	78%	*	*	58%	<b>60%</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	63%	63%	69%	*	*	64%	<b>63%</b>
Q50. During treatment, the patient definitely got enough care and support at home from community or voluntary services	58%	75%	71%	*	*	58%	<b>58%</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	48%	59%	61%	*	*	51%	<b>49%</b>
Q52. Patient has had a review of cancer care by GP practice	23%	30%	38%	40%	*	23%	<b>23%</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	38%	*	47%	*	*	41%	<b>39%</b>
Q54. The right amount of information and support was offered to the patient between final treatment and the follow up appointment	82%	83%	71%	*	*	76%	<b>82%</b>
Q55. Patient was given enough information about the possibility and signs of cancer coming back or spreading	69%	62%	76%	*	*	63%	<b>68%</b>

<b>YOUR OVERALL NHS CARE</b>	Ethnicity						
	White	Mixed	Asian	Black	Other	Not given	All
Q56. The whole care team worked well together	92%	90%	98%	90%	*	94%	<b>92%</b>
Q57. Administration of care was very good or good	89%	81%	93%	100%	*	91%	<b>90%</b>
Q58. Cancer research opportunities were discussed with patient	43%	44%	58%	*	*	38%	<b>43%</b>
Q59. Patient's average rating of care scored from very poor to very good	9.0	8.6	9.0	8.6	*	9.0	<b>9.0</b>

## IMD quintile tables

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

	IMD quintile					
	1 (most deprived)	2	3	4	5 (least deprived)	All
<b>SUPPORT FROM YOUR GP PRACTICE</b>						
Q2. Patient only spoke to primary care professional once or twice before cancer diagnosis	74%	74%	77%	84%	82%	<b>78%</b>
Q3. Referral for diagnosis was explained in a way the patient could completely understand	60%	64%	68%	70%	69%	<b>66%</b>

	IMD quintile					
	1 (most deprived)	2	3	4	5 (least deprived)	All
<b>DIAGNOSTIC TESTS</b>						
Q5. Patient received all the information needed about the diagnostic test in advance	94%	91%	94%	92%	94%	<b>93%</b>
Q6. Diagnostic test staff appeared to completely have all the information they needed about the patient	83%	83%	86%	85%	87%	<b>85%</b>
Q7. Patient felt the length of time waiting for diagnostic test results was about right	82%	81%	82%	81%	80%	<b>81%</b>
Q8. Diagnostic test results were explained in a way the patient could completely understand	78%	78%	83%	81%	80%	<b>80%</b>
Q9. Enough privacy was always given to the patient when receiving diagnostic test results	94%	95%	95%	95%	95%	<b>95%</b>

	IMD quintile					
	1 (most deprived)	2	3	4	5 (least deprived)	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	83%	83%	81%	82%	83%	<b>82%</b>
Q13. Patient was definitely told sensitively that they had cancer	77%	75%	76%	73%	76%	<b>75%</b>
Q14. Cancer diagnosis explained in a way the patient could completely understand	74%	73%	76%	77%	78%	<b>76%</b>
Q15. Patient was definitely told about their diagnosis in an appropriate place	86%	83%	85%	85%	86%	<b>85%</b>
Q16. Patient was told they could go back later for more information about their diagnosis	85%	83%	84%	84%	88%	<b>85%</b>

	IMD quintile					
	1 (most deprived)	2	3	4	5 (least deprived)	All
<b>SUPPORT FROM A MAIN CONTACT PERSON</b>						
Q17. Patient had a main point of contact within the care team	94%	93%	93%	91%	92%	<b>93%</b>
Q18. Patient found it very or quite easy to contact their main contact person	87%	86%	86%	88%	87%	<b>87%</b>
Q19. Patient found advice from main contact person was very or quite helpful	96%	95%	95%	96%	96%	<b>95%</b>

## IMD quintile tables

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

DECIDING ON THE BEST TREATMENT	IMD quintile					
	1 (most deprived)	2	3	4	5 (least deprived)	All
Q20. Treatment options were explained in a way the patient could completely understand	82%	84%	84%	84%	83%	<b>83%</b>
Q21. Patient was definitely involved as much as they wanted to be in decisions about their treatment	80%	80%	81%	83%	82%	<b>81%</b>
Q22. Family and/or carers were definitely involved as much as the patient wanted them to be in decisions about treatment options	82%	81%	83%	85%	87%	<b>83%</b>
Q23. Patient could get further advice from a different healthcare professional before making decisions about their treatment options	62%	58%	57%	61%	59%	<b>60%</b>

CARE PLANNING	IMD quintile					
	1 (most deprived)	2	3	4	5 (least deprived)	All
Q24. Patient was definitely able to have a discussion about their needs or concerns prior to treatment	76%	73%	74%	71%	76%	<b>74%</b>
Q25. A member of their care team helped the patient create a care plan to address any needs or concerns	94%	95%	94%	94%	96%	<b>95%</b>
Q26. Care team reviewed the patient's care plan with them to ensure it was up to date	99%	100%	100%	100%	99%	<b>100%</b>

SUPPORT FROM HOSPITAL STAFF	IMD quintile					
	1 (most deprived)	2	3	4	5 (least deprived)	All
Q27. Staff provided the patient with relevant information on available support	89%	89%	91%	91%	91%	<b>90%</b>
Q28. Patient definitely got the right level of support for their overall health and well being from hospital staff	79%	78%	80%	78%	79%	<b>79%</b>
Q29. Patient was offered information about how to get financial help or benefits	73%	72%	70%	69%	76%	<b>72%</b>

HOSPITAL CARE	IMD quintile					
	1 (most deprived)	2	3	4	5 (least deprived)	All
Q31. Patient had confidence and trust in all of the team looking after them during their stay in hospital	81%	76%	79%	76%	80%	<b>78%</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	74%	73%	74%	71%	74%	<b>73%</b>
Q33. Patient was always involved in decisions about their care and treatment whilst in hospital	74%	72%	71%	69%	73%	<b>72%</b>
Q34. Patient was always able to get help from ward staff when needed	78%	72%	75%	72%	73%	<b>74%</b>
Q35. Patient was always able to discuss worries and fears with hospital staff	71%	64%	71%	63%	69%	<b>68%</b>
Q36. Hospital staff always did everything they could to help the patient control pain	86%	82%	81%	85%	87%	<b>84%</b>
Q37. Patient was always treated with respect and dignity while in hospital	89%	87%	89%	89%	87%	<b>88%</b>
Q38. Patient received easily understandable information about what they should or should not do after leaving hospital	88%	86%	90%	87%	91%	<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	81%	83%	83%	84%	85%	<b>83%</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)	All
Q41_1. Beforehand patient completely had enough understandable information about surgery	91%	90%	91%	88%	91%	90%
Q41_2. Beforehand patient completely had enough understandable information about chemotherapy	89%	90%	88%	89%	87%	88%
Q41_3. Beforehand patient completely had enough understandable information about radiotherapy	90%	92%	87%	92%	87%	90%
Q41_4. Beforehand patient completely had enough understandable information about hormone therapy	84%	86%	83%	77%	82%	82%
Q41_5. Beforehand patient completely had enough understandable information about immunotherapy	87%	84%	88%	82%	85%	85%
Q42_1. Patient completely had enough understandable information about their response to surgery	87%	87%	88%	87%	87%	87%
Q42_2. Patient completely had enough understandable information about their response to chemotherapy	84%	86%	85%	84%	84%	84%
Q42_3. Patient completely had enough understandable information about their response to radiotherapy	87%	89%	83%	89%	83%	86%
Q42_4. Patient completely had enough understandable information about their response to hormone therapy	83%	85%	79%	79%	82%	82%
Q42_5. Patient completely had enough understandable information about their response to immunotherapy	86%	82%	87%	76%	89%	84%
Q43. Patient felt the length of waiting time at clinic and day unit for cancer treatment was about right	81%	83%	82%	82%	83%	82%

IMMEDIATE AND LONG TERM SIDE EFFECTS	IMD quintile					
	1 (most deprived)	2	3	4	5 (least deprived)	All
Q44. Possible side effects from treatment were definitely explained in a way the patient could understand	78%	76%	76%	77%	77%	77%
Q45. Patient was always offered practical advice on dealing with any immediate side effects from treatment	75%	72%	72%	75%	72%	73%
Q46. Patient was given information that they could access about support in dealing with immediate side effects from treatment	87%	89%	89%	87%	92%	89%
Q47. Patient felt possible long-term side effects were definitely explained in a way they could understand in advance of their treatment	65%	63%	63%	64%	62%	64%
Q48. Patient was definitely able to discuss options for managing the impact of any long-term side effects	62%	60%	58%	59%	61%	60%

SUPPORT WHILE AT HOME	IMD quintile					
	1 (most deprived)	2	3	4	5 (least deprived)	All
Q49. Care team gave family, or someone close, all the information needed to help care for the patient at home	64%	60%	66%	64%	64%	63%
Q50. During treatment, the patient definitely got enough care and support at home from community or voluntary services	56%	58%	57%	61%	62%	58%

CARE FROM YOUR GP PRACTICE	IMD quintile					
	1 (most deprived)	2	3	4	5 (least deprived)	All
Q51. Patient definitely received the right amount of support from their GP practice during treatment	47%	46%	50%	51%	50%	49%
Q52. Patient has had a review of cancer care by GP practice	25%	24%	22%	25%	21%	23%

## IMD quintile tables

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

<b>LIVING WITH AND BEYOND CANCER</b>	IMD quintile					
	1 (most deprived)	2	3	4	5 (least deprived)	All
Q53. After treatment, the patient definitely could get enough emotional support at home from community or voluntary services	35%	41%	40%	34%	48%	<b>39%</b>
Q54. The right amount of information and support was offered to the patient between final treatment and the follow up appointment	82%	79%	82%	81%	83%	<b>82%</b>
Q55. Patient was given enough information about the possibility and signs of cancer coming back or spreading	69%	67%	69%	70%	68%	<b>68%</b>

<b>YOUR OVERALL NHS CARE</b>	IMD quintile					
	1 (most deprived)	2	3	4	5 (least deprived)	All
Q56. The whole care team worked well together	92%	91%	93%	92%	92%	<b>92%</b>
Q57. Administration of care was very good or good	89%	88%	92%	90%	89%	<b>90%</b>
Q58. Cancer research opportunities were discussed with patient	49%	39%	44%	41%	40%	<b>43%</b>
Q59. Patient's average rating of care scored from very poor to very good	9.0	8.9	9.0	9.0	9.0	<b>9.0</b>

## Long-term condition status tables

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

	Long-term condition status			
	Yes	No	Not given	All
<b>SUPPORT FROM YOUR GP PRACTICE</b>				
Q2. Patient only spoke to primary care professional once or twice before cancer diagnosis	76%	82%	79%	<b>78%</b>
Q3. Referral for diagnosis was explained in a way the patient could completely understand	64%	70%	66%	<b>66%</b>

	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	93%	94%	93%	<b>93%</b>
Q6. Diagnostic test staff appeared to completely have all the information they needed about the patient	84%	87%	83%	<b>85%</b>
Q7. Patient felt the length of time waiting for diagnostic test results was about right	82%	79%	81%	<b>81%</b>
Q8. Diagnostic test results were explained in a way the patient could completely understand	79%	82%	79%	<b>80%</b>
Q9. Enough privacy was always given to the patient when receiving diagnostic test results	95%	95%	95%	<b>95%</b>

	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	81%	84%	86%	<b>82%</b>
Q13. Patient was definitely told sensitively that they had cancer	75%	75%	81%	<b>75%</b>
Q14. Cancer diagnosis explained in a way the patient could completely understand	75%	76%	77%	<b>76%</b>
Q15. Patient was definitely told about their diagnosis in an appropriate place	84%	85%	88%	<b>85%</b>
Q16. Patient was told they could go back later for more information about their diagnosis	84%	87%	85%	<b>85%</b>

	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	92%	93%	93%	<b>93%</b>
Q18. Patient found it very or quite easy to contact their main contact person	87%	88%	84%	<b>87%</b>
Q19. Patient found advice from main contact person was very or quite helpful	95%	96%	96%	<b>95%</b>

	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	82%	85%	83%	<b>83%</b>
Q21. Patient was definitely involved as much as they wanted to be in decisions about their treatment	81%	82%	80%	<b>81%</b>
Q22. Family and/or carers were definitely involved as much as the patient wanted them to be in decisions about treatment options	83%	85%	85%	<b>83%</b>
Q23. Patient could get further advice from a different healthcare professional before making decisions about their treatment options	59%	63%	58%	<b>60%</b>



## Long-term condition status tables

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

<b>CARE PLANNING</b>	Long-term condition status			
	Yes	No	Not given	All
Q24. Patient was definitely able to have a discussion about their needs or concerns prior to treatment	73%	76%	76%	<b>74%</b>
Q25. A member of their care team helped the patient create a care plan to address any needs or concerns	94%	95%	97%	<b>95%</b>
Q26. Care team reviewed the patient's care plan with them to ensure it was up to date	99%	100%	100%	<b>100%</b>

<b>SUPPORT FROM HOSPITAL STAFF</b>	Long-term condition status			
	Yes	No	Not given	All
Q27. Staff provided the patient with relevant information on available support	89%	93%	91%	<b>90%</b>
Q28. Patient definitely got the right level of support for their overall health and well being from hospital staff	79%	80%	76%	<b>79%</b>
Q29. Patient was offered information about how to get financial help or benefits	71%	75%	73%	<b>72%</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	78%	78%	85%	<b>78%</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	73%	74%	74%	<b>73%</b>
Q33. Patient was always involved in decisions about their care and treatment whilst in hospital	69%	76%	78%	<b>72%</b>
Q34. Patient was always able to get help from ward staff when needed	73%	75%	82%	<b>74%</b>
Q35. Patient was always able to discuss worries and fears with hospital staff	67%	69%	69%	<b>68%</b>
Q36. Hospital staff always did everything they could to help the patient control pain	84%	86%	82%	<b>84%</b>
Q37. Patient was always treated with respect and dignity while in hospital	87%	92%	90%	<b>88%</b>
Q38. Patient received easily understandable information about what they should or should not do after leaving hospital	88%	89%	89%	<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	82%	85%	83%	<b>83%</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	90%	91%	91%	<b>90%</b>
Q41_2. Beforehand patient completely had enough understandable information about chemotherapy	89%	88%	88%	<b>88%</b>
Q41_3. Beforehand patient completely had enough understandable information about radiotherapy	91%	90%	83%	<b>90%</b>
Q41_4. Beforehand patient completely had enough understandable information about hormone therapy	84%	81%	75%	<b>82%</b>
Q41_5. Beforehand patient completely had enough understandable information about immunotherapy	86%	85%	82%	<b>85%</b>
Q42_1. Patient completely had enough understandable information about their response to surgery	87%	89%	88%	<b>87%</b>
Q42_2. Patient completely had enough understandable information about their response to chemotherapy	84%	85%	86%	<b>84%</b>
Q42_3. Patient completely had enough understandable information about their response to radiotherapy	86%	87%	83%	<b>86%</b>
Q42_4. Patient completely had enough understandable information about their response to hormone therapy	82%	82%	79%	<b>82%</b>
Q42_5. Patient completely had enough understandable information about their response to immunotherapy	84%	86%	81%	<b>84%</b>
Q43. Patient felt the length of waiting time at clinic and day unit for cancer treatment was about right	83%	82%	80%	<b>82%</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	76%	81%	72%	<b>77%</b>
Q45. Patient was always offered practical advice on dealing with any immediate side effects from treatment	72%	76%	72%	<b>73%</b>
Q46. Patient was given information that they could access about support in dealing with immediate side effects from treatment	87%	91%	89%	<b>89%</b>
Q47. Patient felt possible long-term side effects were definitely explained in a way they could understand in advance of their treatment	62%	67%	61%	<b>64%</b>
Q48. Patient was definitely able to discuss options for managing the impact of any long-term side effects	59%	64%	60%	<b>60%</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	63%	66%	61%	<b>63%</b>
Q50. During treatment, the patient definitely got enough care and support at home from community or voluntary services	56%	65%	59%	<b>58%</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	47%	53%	47%	<b>49%</b>
Q52. Patient has had a review of cancer care by GP practice	24%	23%	23%	<b>23%</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	All
Q53. After treatment, the patient definitely could get enough emotional support at home from community or voluntary services	37%	44%	43%	<b>39%</b>
Q54. The right amount of information and support was offered to the patient between final treatment and the follow up appointment	79%	87%	79%	<b>82%</b>
Q55. Patient was given enough information about the possibility and signs of cancer coming back or spreading	68%	70%	65%	<b>68%</b>

<b>YOUR OVERALL NHS CARE</b>	Long-term condition status			
	Yes	No	Not given	All
Q56. The whole care team worked well together	91%	93%	95%	<b>92%</b>
Q57. Administration of care was very good or good	89%	91%	92%	<b>90%</b>
Q58. Cancer research opportunities were discussed with patient	42%	44%	41%	<b>43%</b>
Q59. Patient's average rating of care scored from very poor to very good	9.0	9.1	9.0	<b>9.0</b>

## Year on year charts

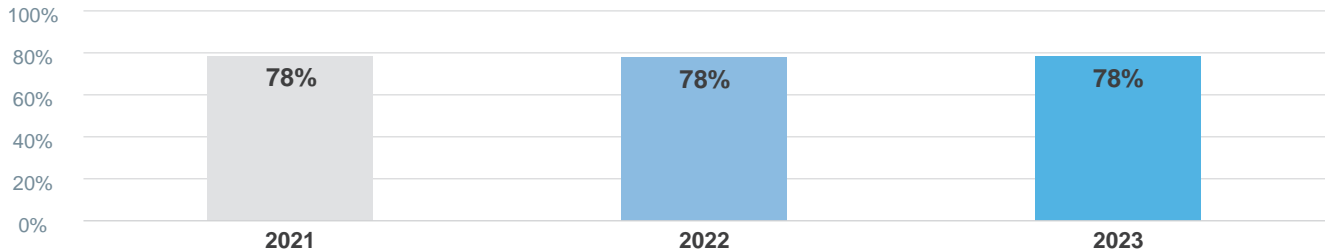
\* Indicates where a score is not available due to suppression or a low base size.

\*\* No score available for these years.

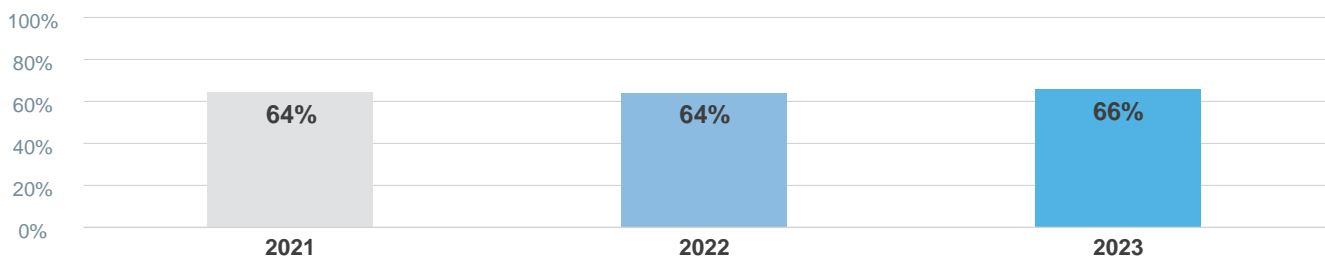
The scores are unadjusted and based on England scores only.

### SUPPORT FROM YOUR GP PRACTICE

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

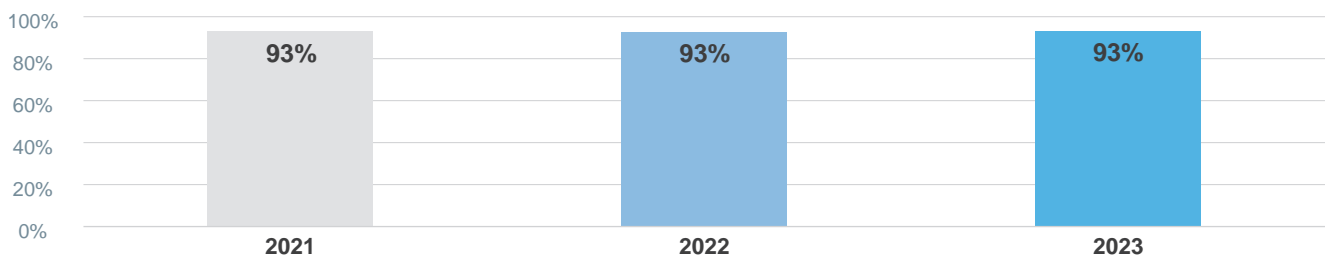


Q3. Referral for diagnosis was explained in a way the patient could completely understand

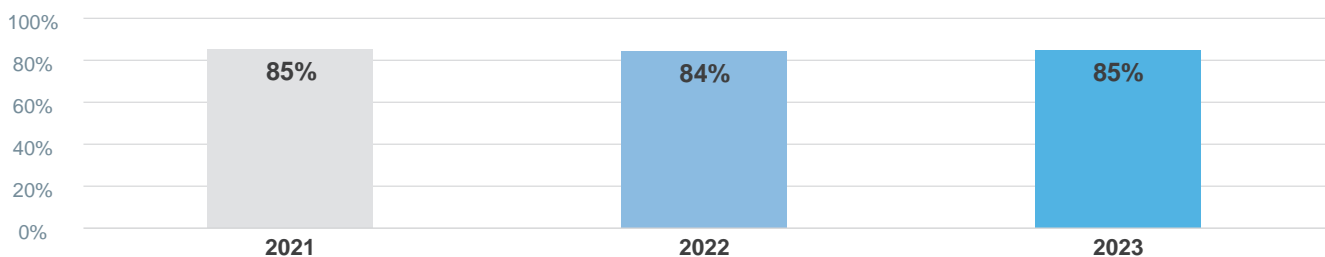


### DIAGNOSTIC TESTS

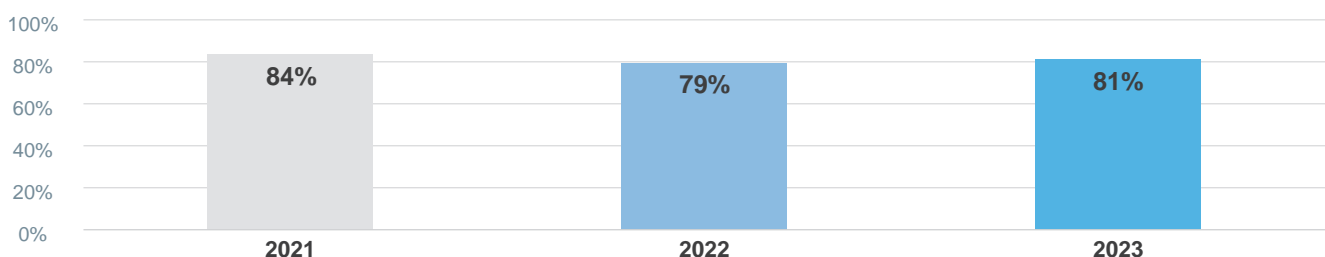
Q5. Patient received all the information needed about the diagnostic test in advance



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



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



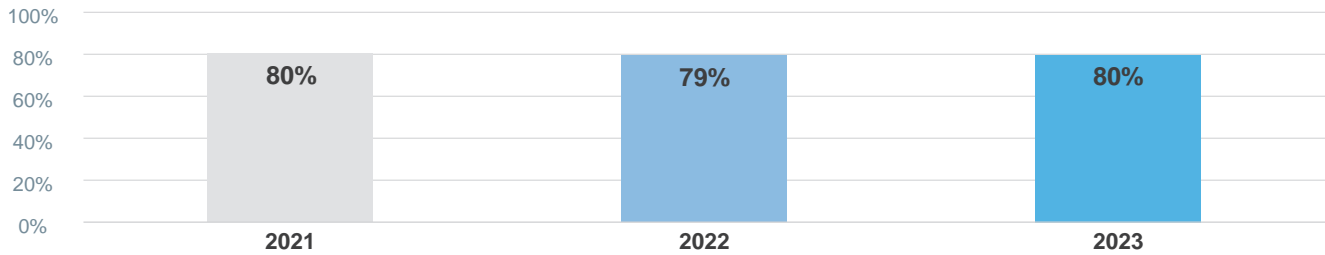
## Year on year charts

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

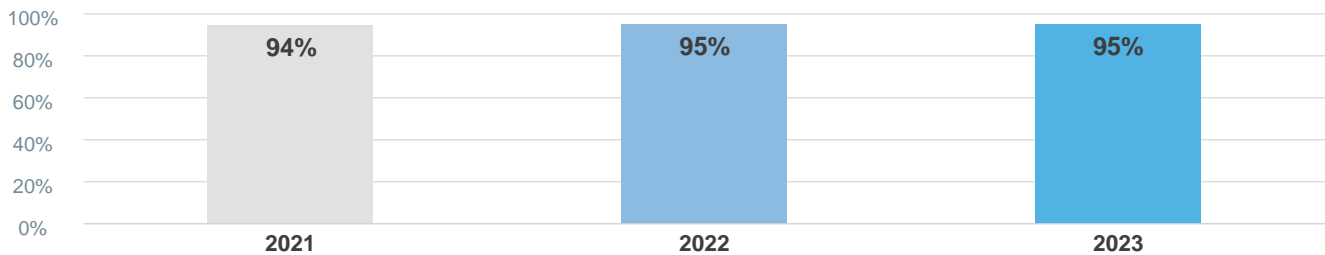
\*\* No score available for these years.

The scores are unadjusted and based on England scores only.

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

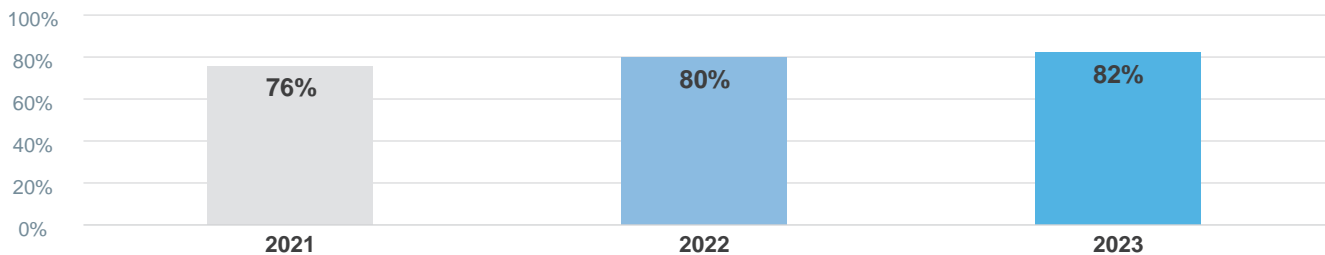


Q9. Enough privacy was always given to the patient when receiving diagnostic test results

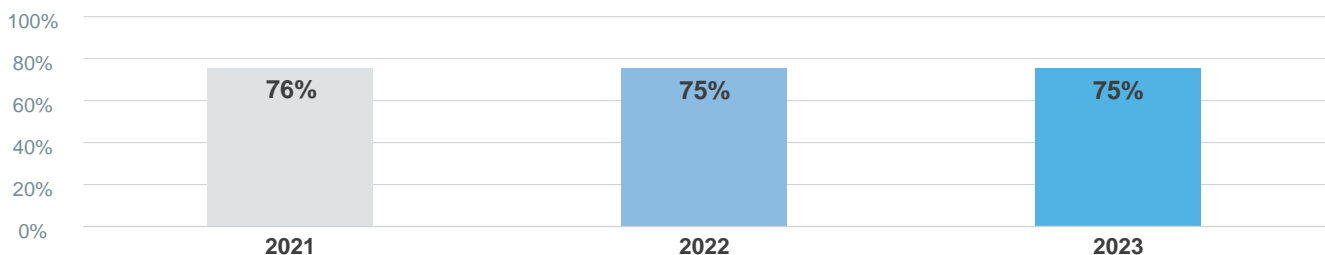


### FINDING OUT THAT YOU HAD CANCER

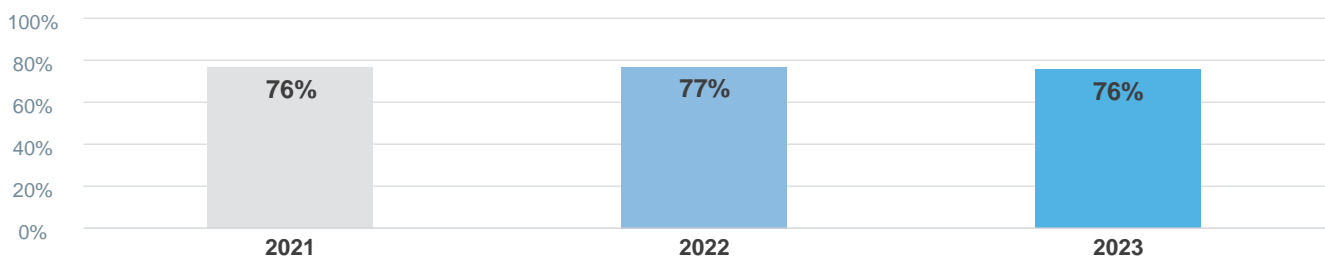
Q12. Patient was told they could have a family member, carer or friend with them when told diagnosis



Q13. Patient was definitely told sensitively that they had cancer



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



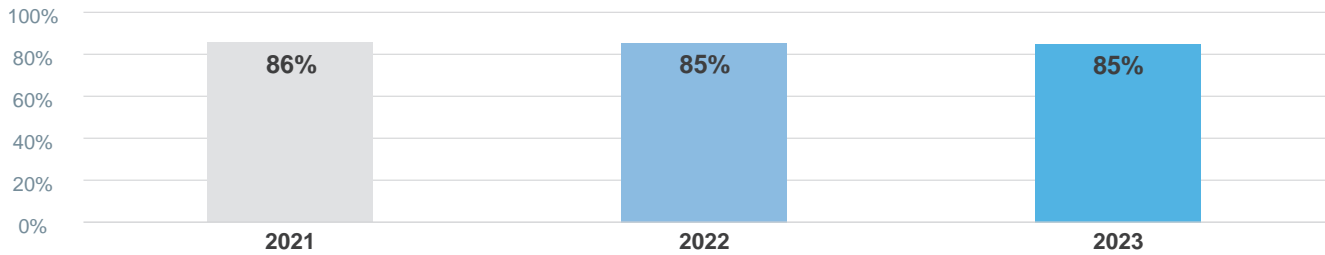
## Year on year charts

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

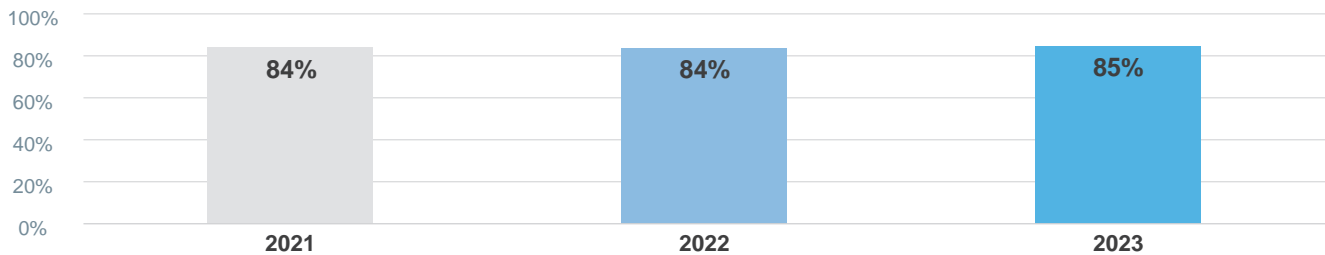
\*\* No score available for these years.

The scores are unadjusted and based on England scores only.

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

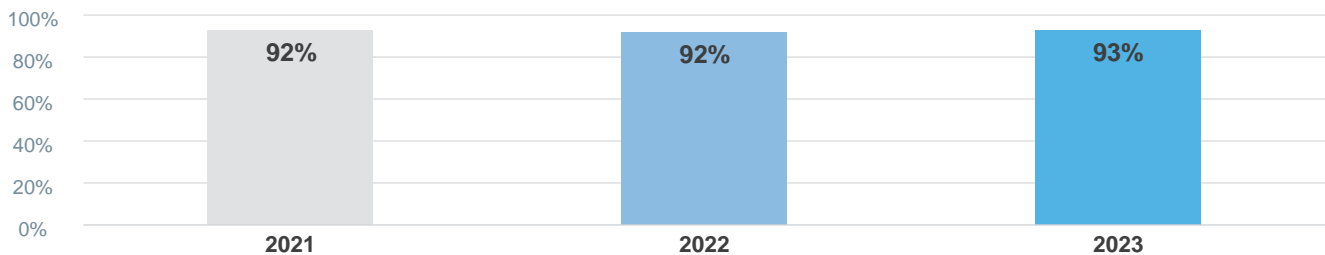


Q16. Patient was told they could go back later for more information about their diagnosis

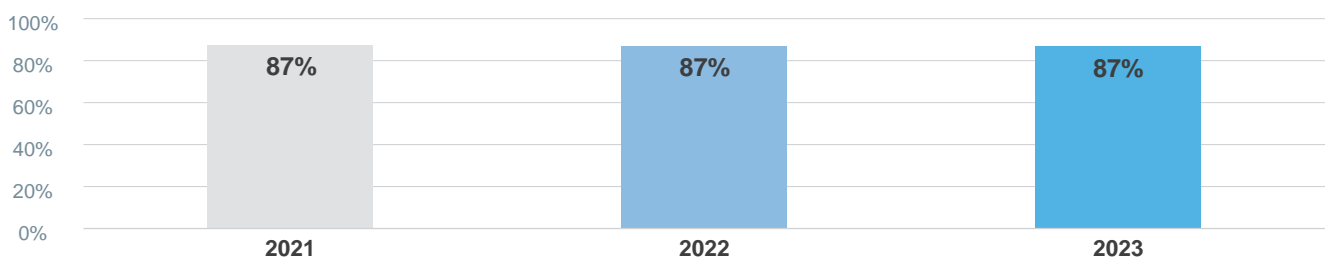


### SUPPORT FROM A MAIN CONTACT PERSON

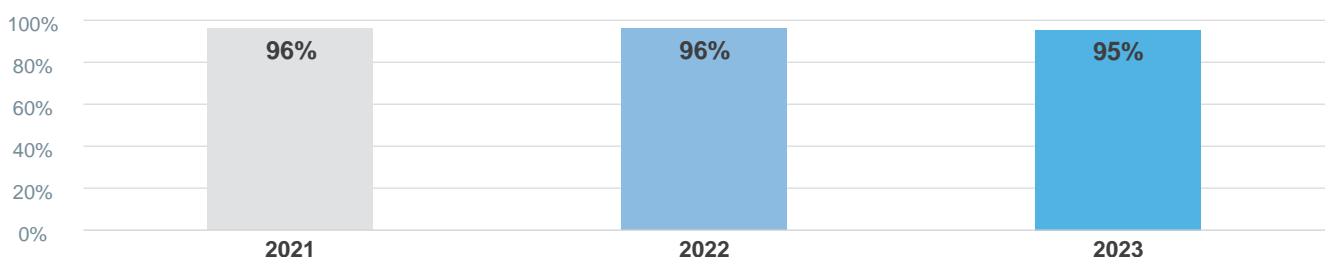
Q17. Patient had a main point of contact within the care team



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



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



## Year on year charts

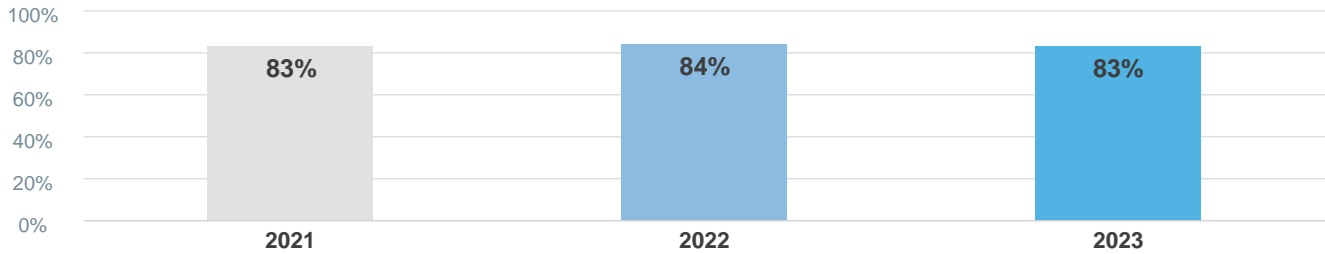
\* Indicates where a score is not available due to suppression or a low base size.

\*\* No score available for these years.

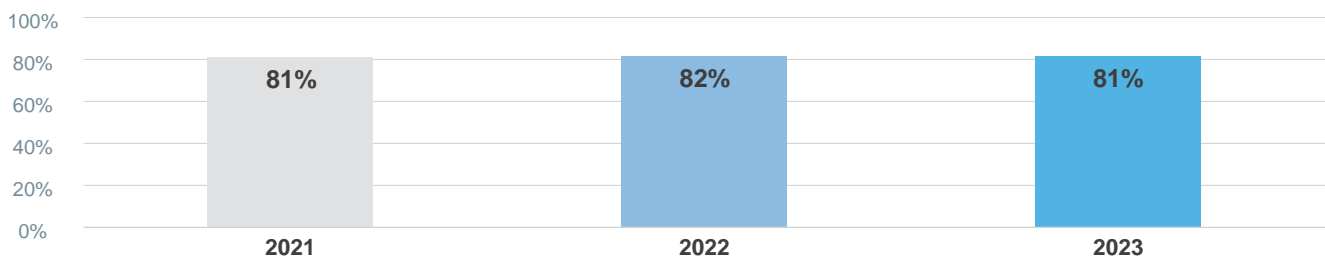
The scores are unadjusted and based on England scores only.

### DECIDING ON THE BEST TREATMENT

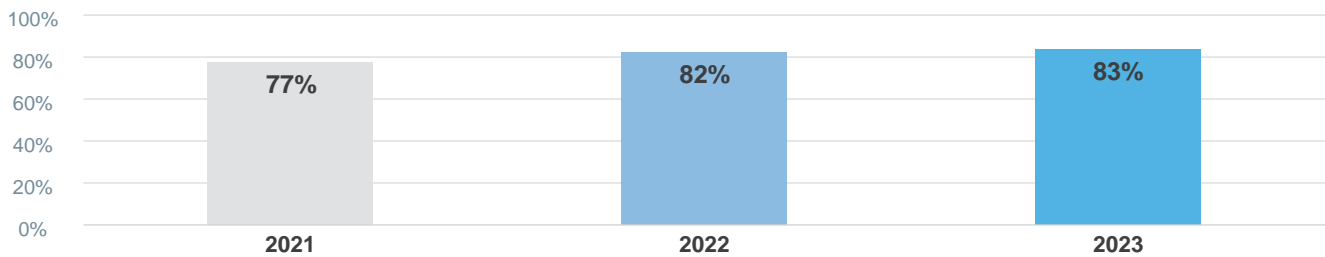
Q20. Treatment options were explained in a way the patient could completely understand



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



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

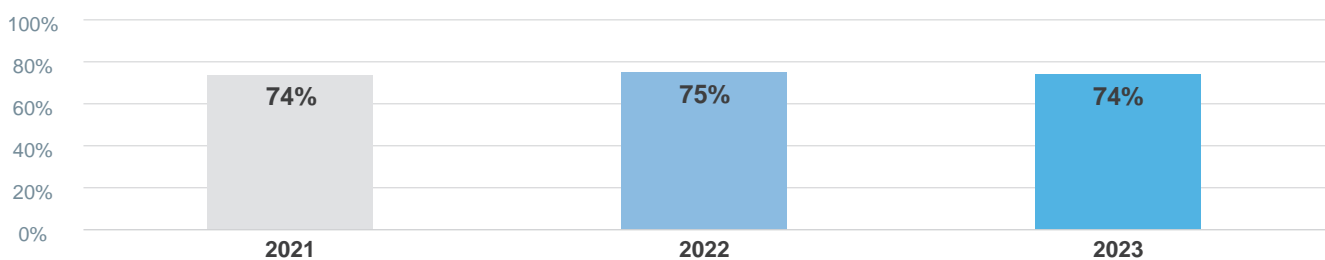


Q23. Patient could get further advice from a different healthcare professional before making decisions about their treatment options



### CARE PLANNING

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



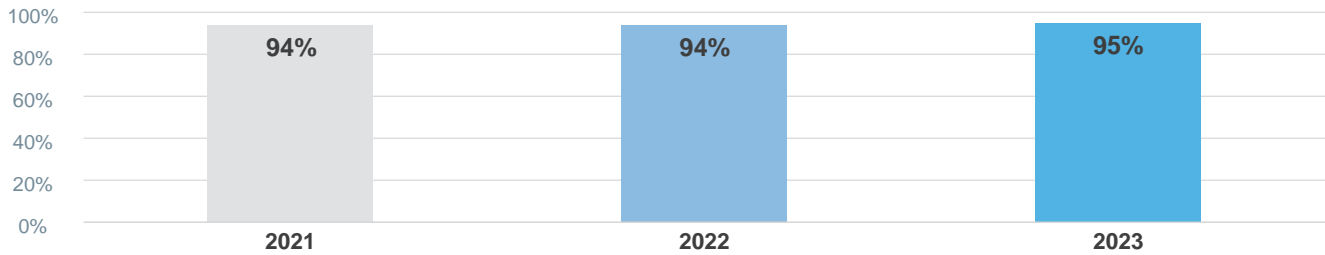
## Year on year charts

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

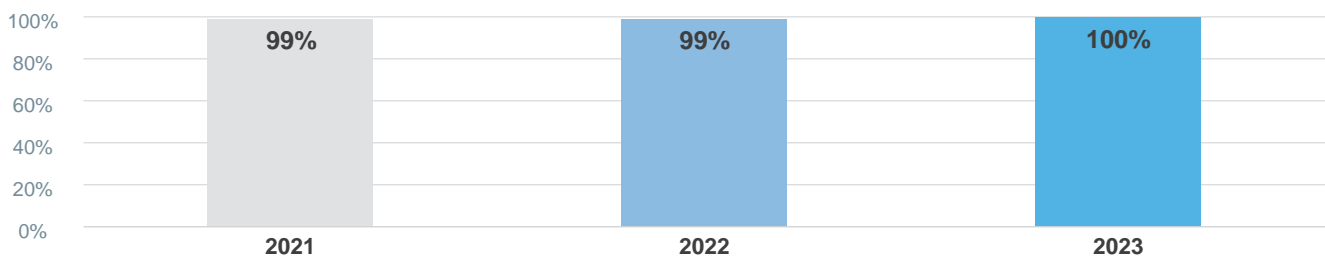
\*\* No score available for these years.

The scores are unadjusted and based on England scores only.

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

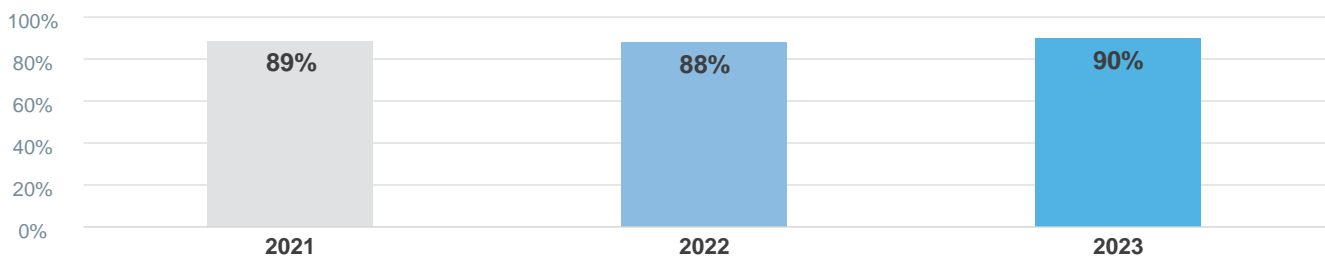


Q26. Care team reviewed the patient's care plan with them to ensure it was up to date

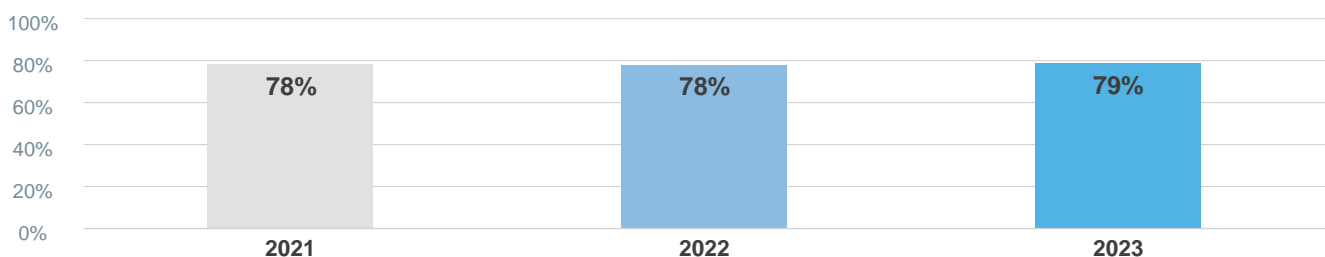


### SUPPORT FROM HOSPITAL STAFF

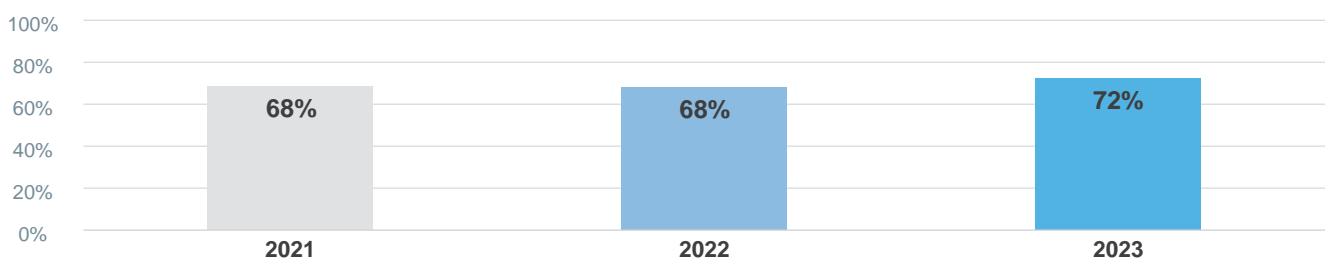
Q27. Staff provided the patient with relevant information on available support



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



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





## Year on year charts

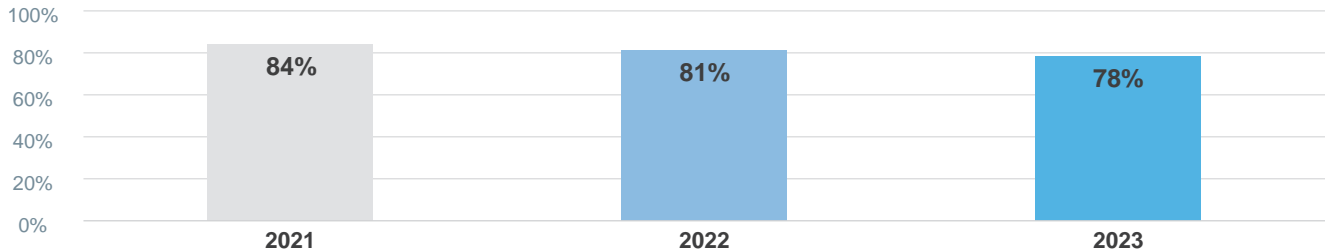
\* Indicates where a score is not available due to suppression or a low base size.

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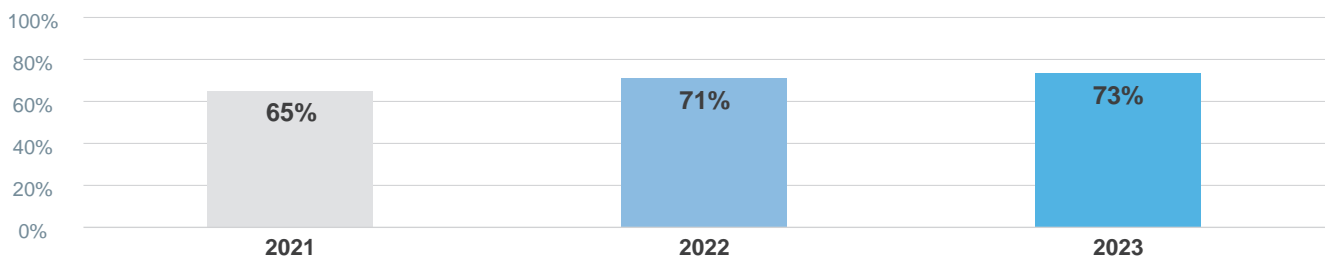
The scores are unadjusted and based on England scores only.

### HOSPITAL CARE

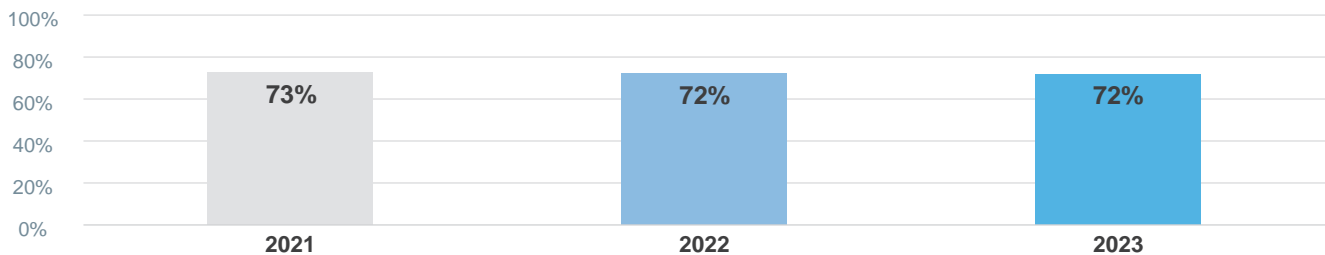
Q31. Patient had confidence and trust in all of the team looking after them during their stay in hospital



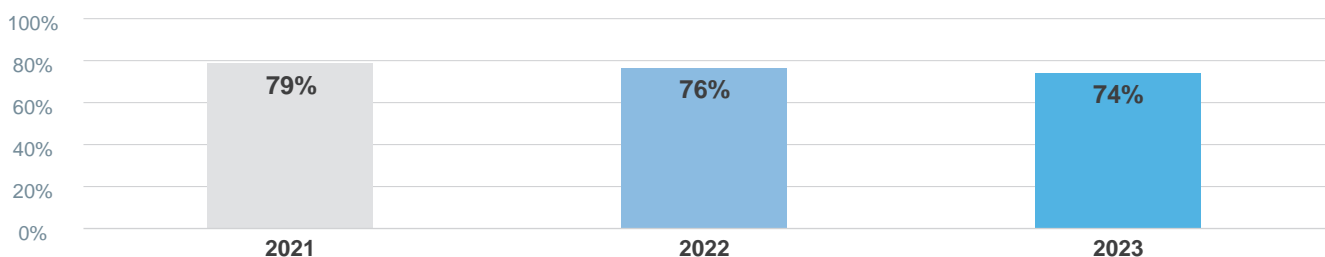
Q32. Patient's family, or someone close, was definitely able to talk to a member of the team looking after the patient in hospital



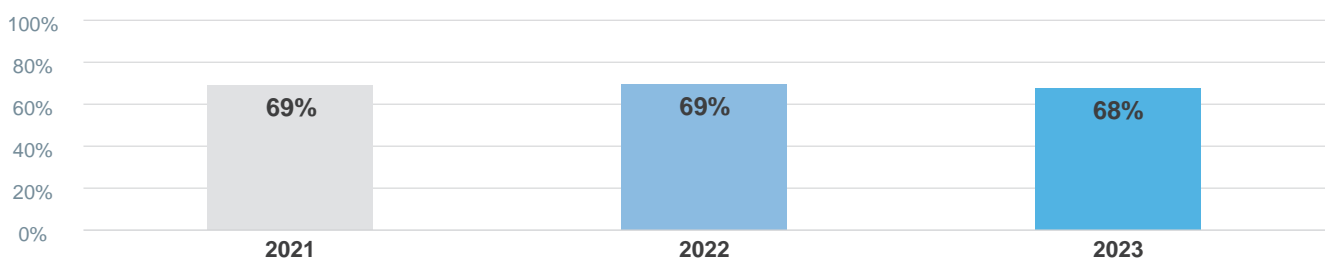
Q33. Patient was always involved in decisions about their care and treatment whilst in hospital



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



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



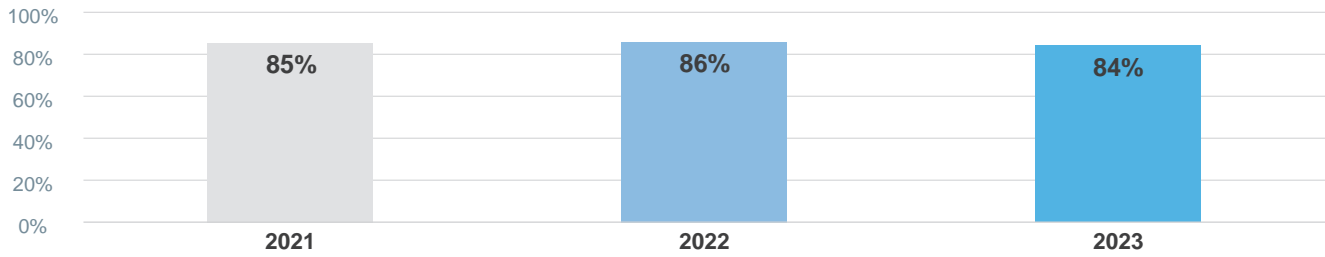
## Year on year charts

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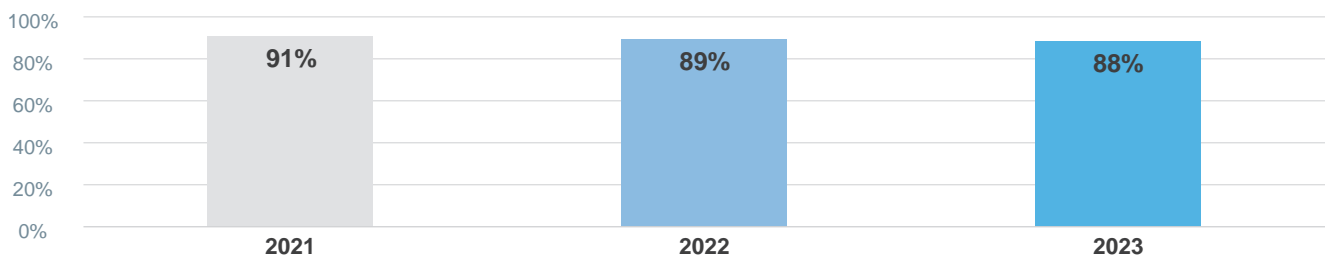
\*\* No score available for these years.

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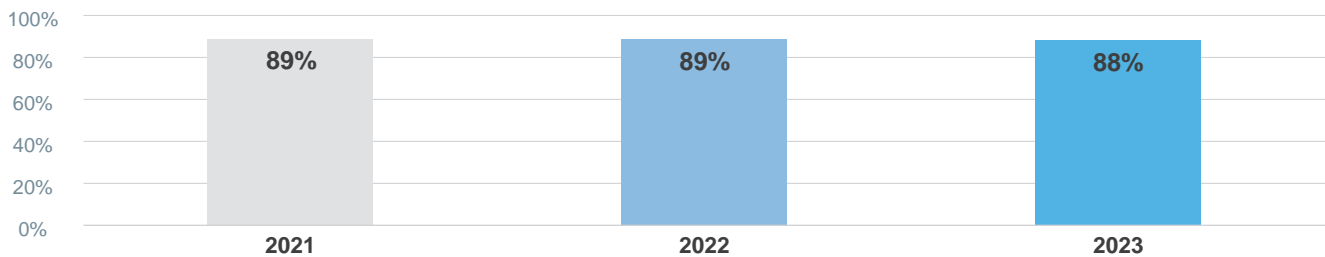
### Q36. Hospital staff always did everything they could to help the patient control pain



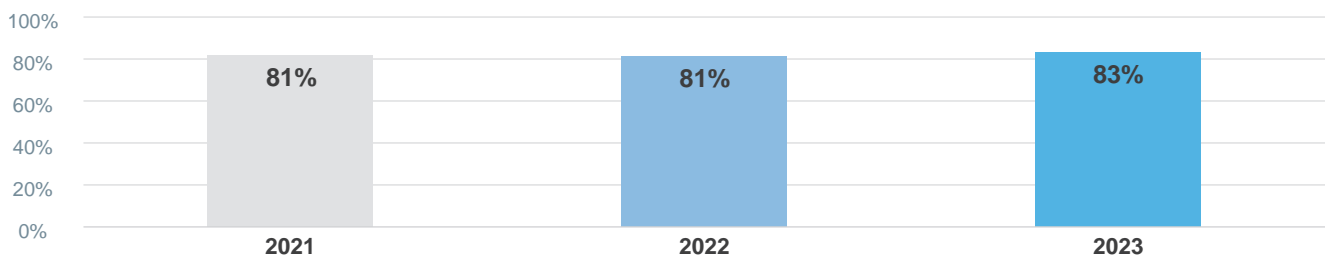
### Q37. Patient was always treated with respect and dignity while in hospital



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

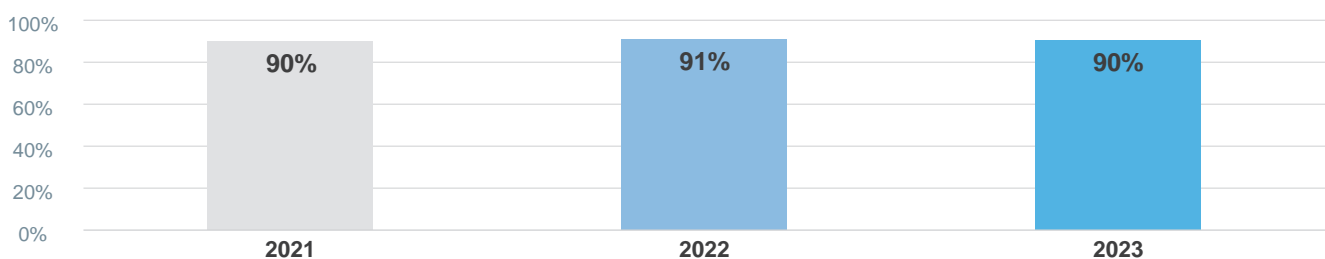


### Q39. Patient was always able to discuss worries and fears with hospital staff while being treated as an outpatient or day case



## YOUR TREATMENT

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



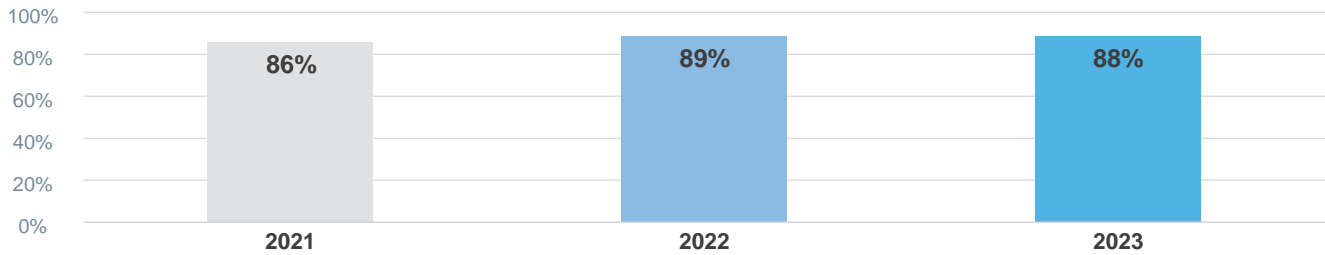
## Year on year charts

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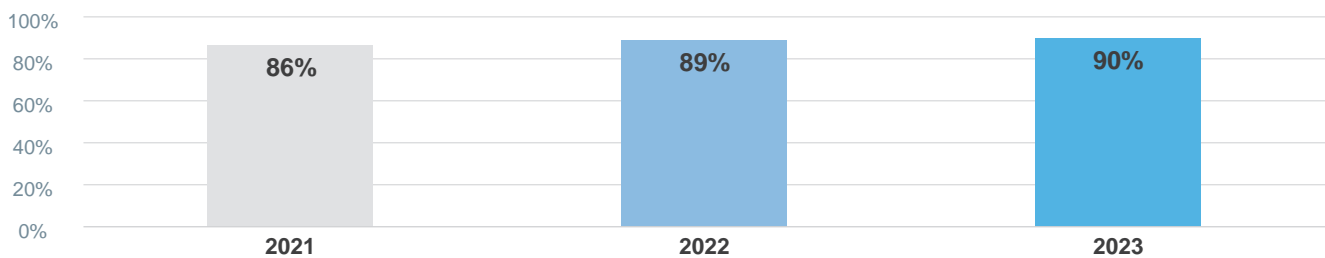
\*\* No score available for these years.

The scores are unadjusted and based on England scores only.

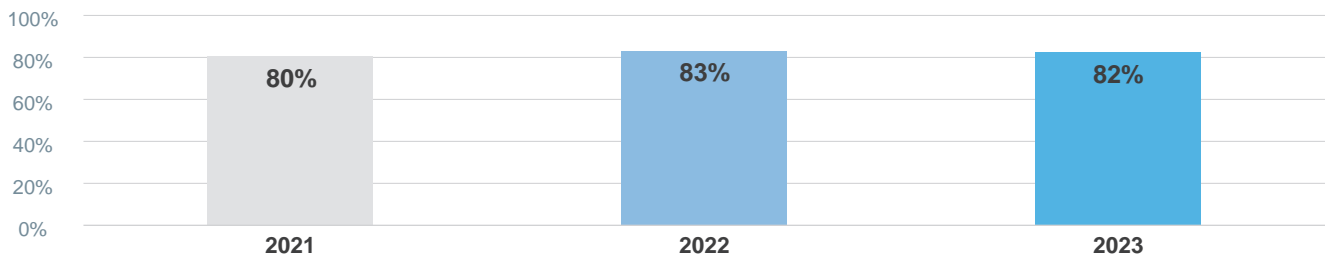
Q41\_2. Beforehand patient completely had enough understandable information about chemotherapy



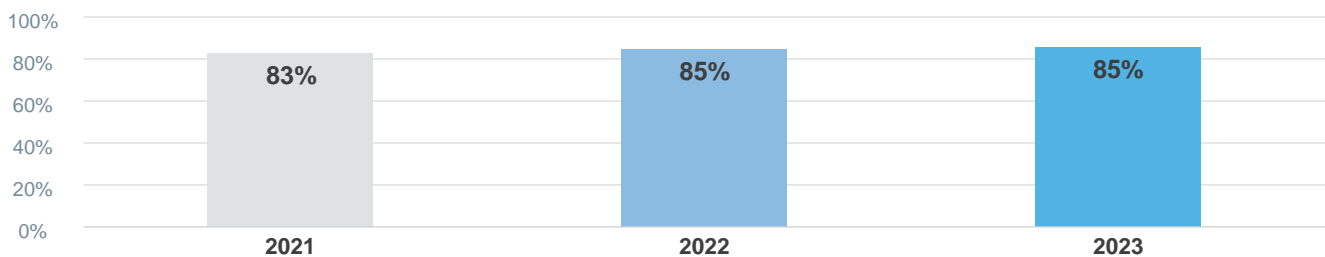
Q41\_3. Beforehand patient completely had enough understandable information about radiotherapy



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



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



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



## Year on year charts

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\*\* No score available for these years.

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



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



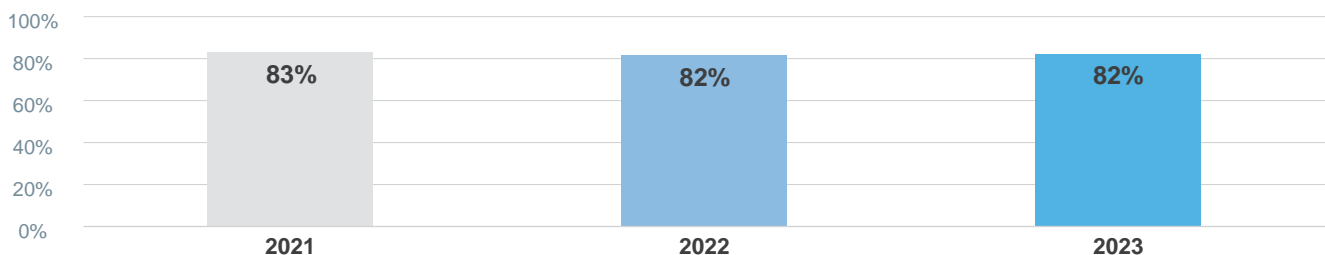
Q42\_4. Patient completely had enough understandable information about their response to hormone therapy



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



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



## Year on year charts

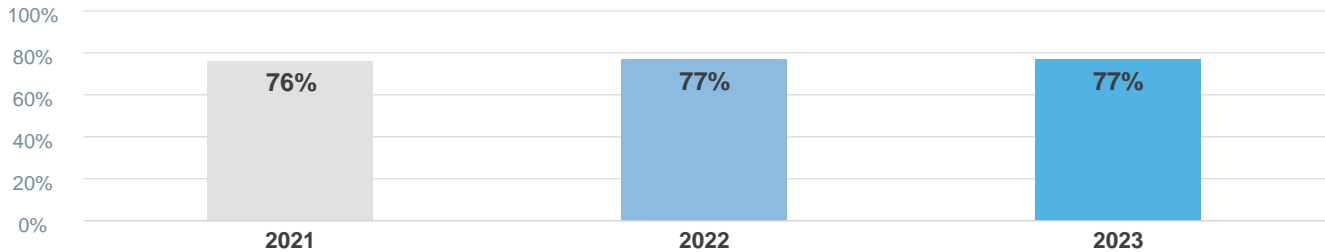
\* Indicates where a score is not available due to suppression or a low base size.

\*\* No score available for these years.

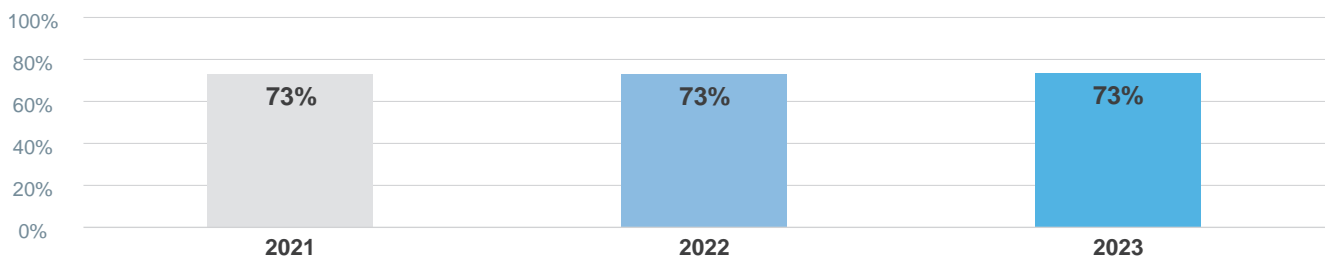
The scores are unadjusted and based on England scores only.

### IMMEDIATE AND LONG TERM SIDE EFFECTS

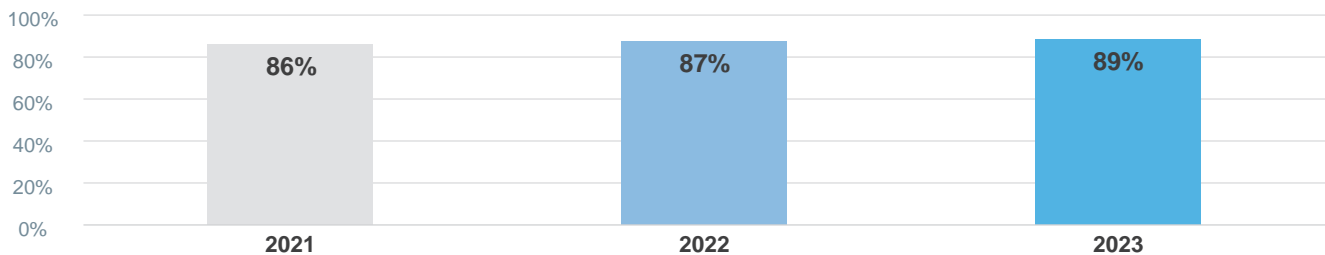
Q44. Possible side effects from treatment were definitely explained in a way the patient could understand



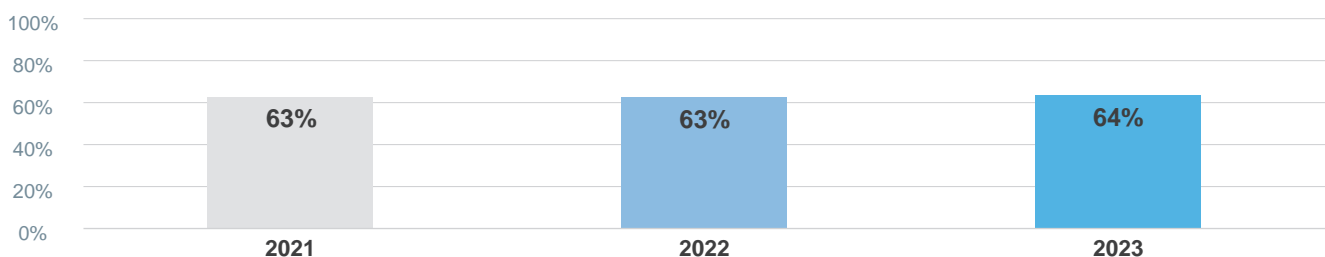
Q45. Patient was always offered practical advice on dealing with any immediate side effects from treatment



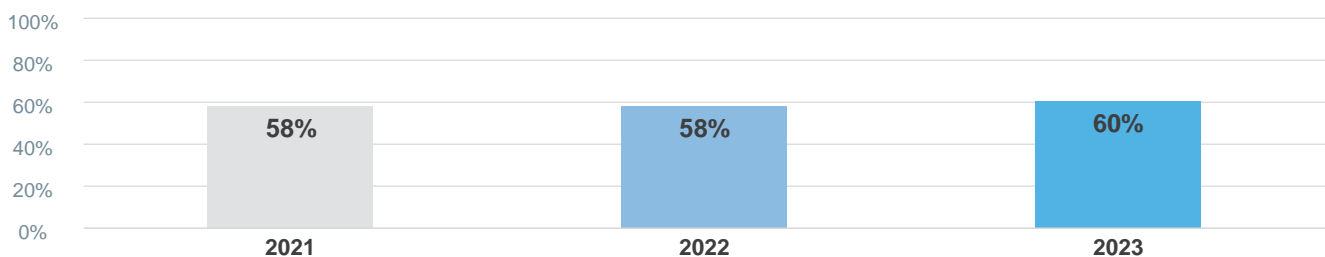
Q46. Patient was given information that they could access about support in dealing with immediate side effects from treatment



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



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



## Year on year charts

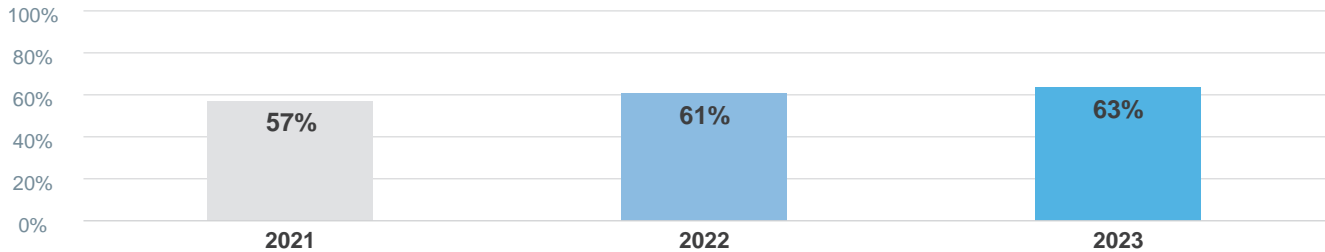
\* Indicates where a score is not available due to suppression or a low base size.

\*\* No score available for these years.

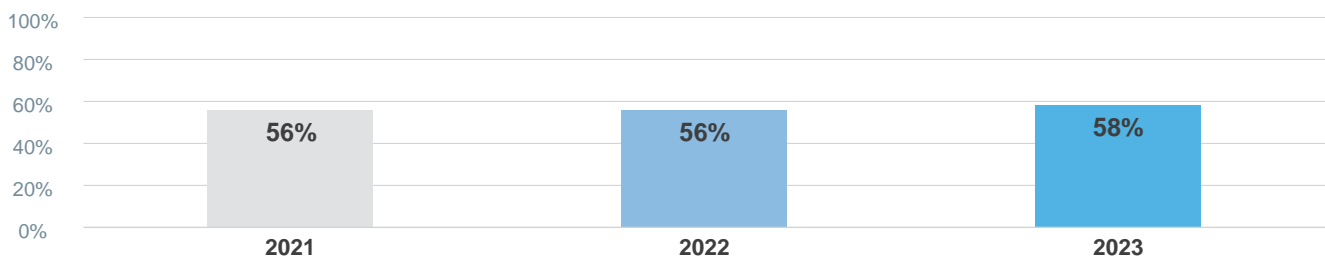
The scores are unadjusted and based on England scores only.

### SUPPORT WHILE AT HOME

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

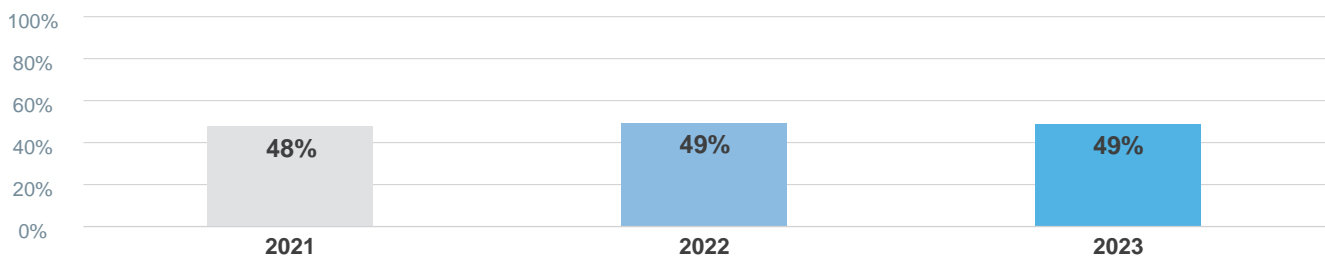


Q50. During treatment, the patient definitely got enough care and support at home from community or voluntary services

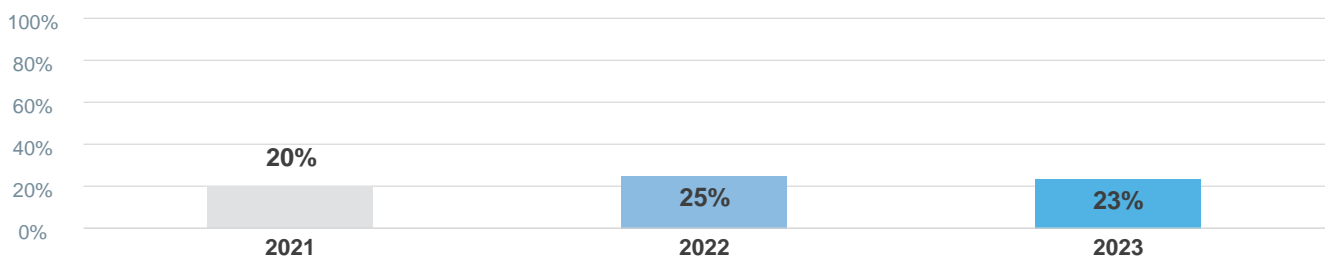


### CARE FROM YOUR GP PRACTICE

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

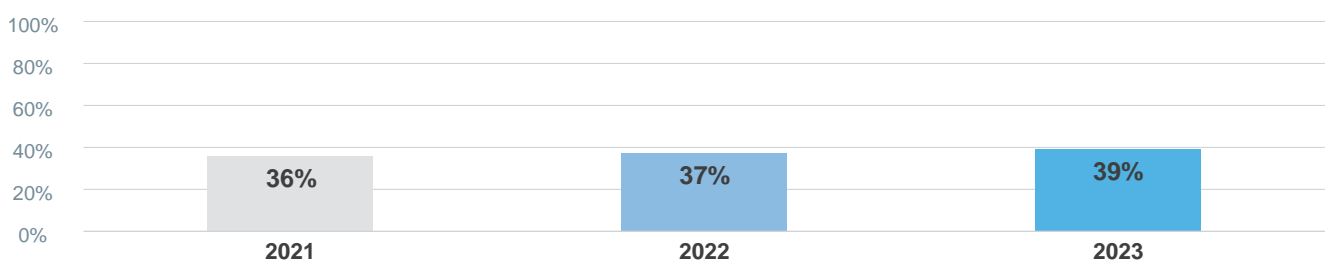


Q52. Patient has had a review of cancer care by GP practice



### LIVING WITH AND BEYOND CANCER

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



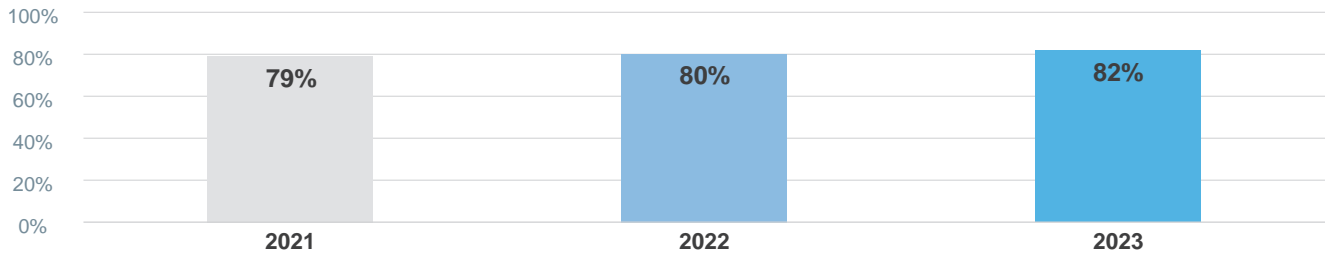
## Year on year charts

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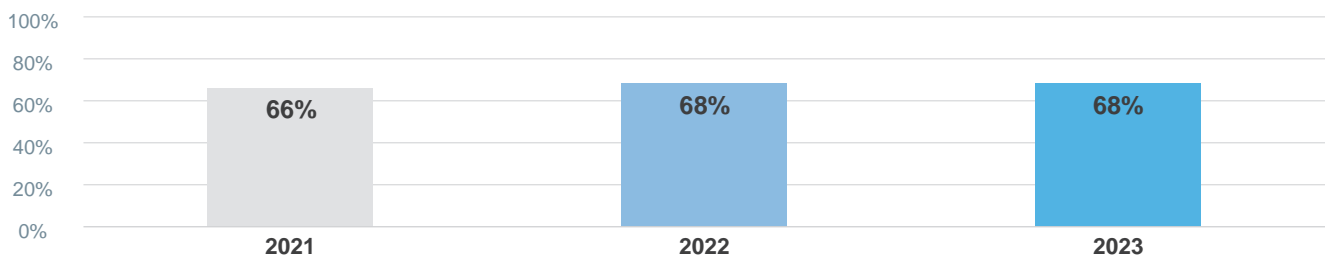
\*\* No score available for these years.

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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

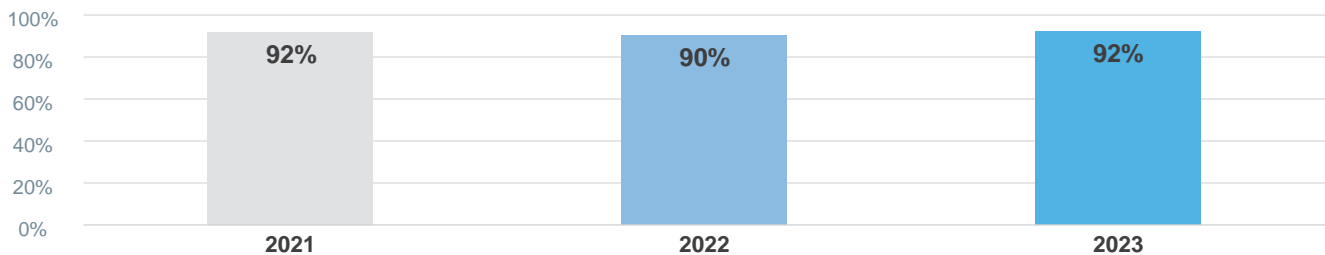


Q55. Patient was given enough information about the possibility and signs of cancer coming back or spreading

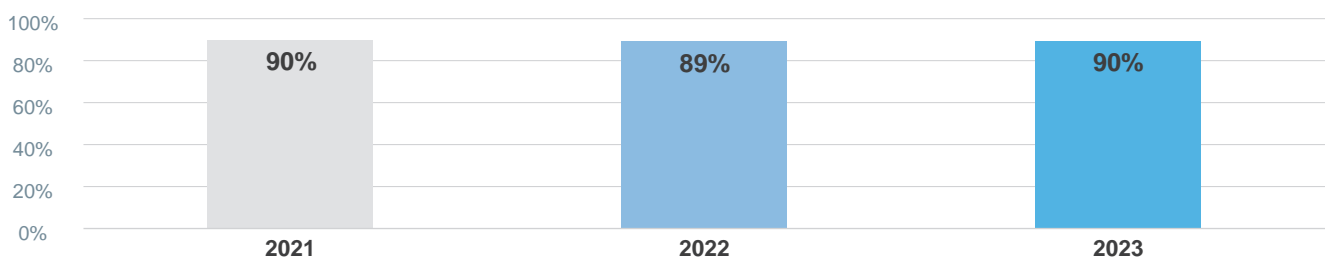


### YOUR OVERALL NHS CARE

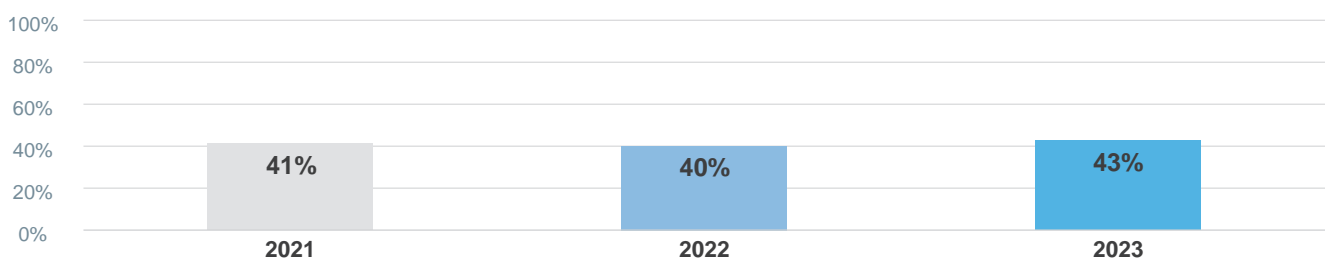
Q56. The whole care team worked well together



Q57. Administration of care was very good or good



Q58. Cancer research opportunities were discussed with patient



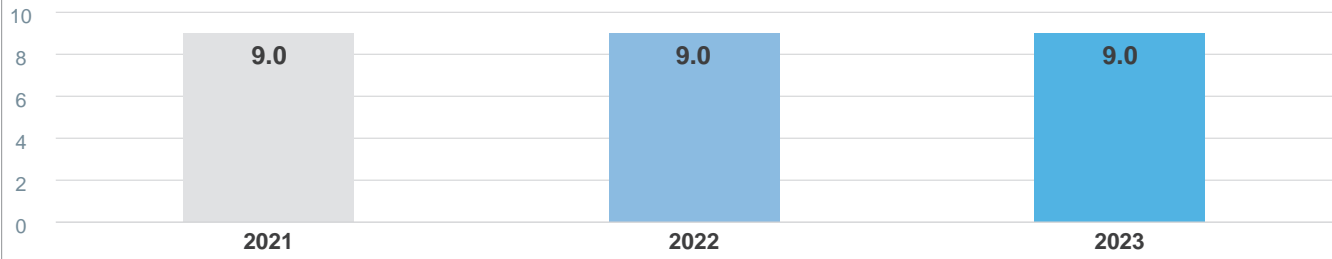
## Year on year charts

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\*\* No score available for these years.




The scores are unadjusted and based on England scores only.

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








## Trust expected range summary

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

Trust		Expected range classification		
RR7	Gateshead Health NHS Foundation Trust	1	35	25
RTF	Northumbria Healthcare NHS Foundation Trust	1	37	23
R0B	South Tyneside and Sunderland NHS Foundation Trust	1	46	14
RTD	The Newcastle Upon Tyne Hospitals NHS Foundation Trust		54	7
RXP	County Durham and Darlington NHS Foundation Trust	1	55	5
RNN	North Cumbria Integrated Care NHS Foundation Trust	1	58	2
RTR	South Tees Hospitals NHS Foundation Trust	2	56	3
RVW	North Tees and Hartlepool NHS Foundation Trust	3	57	1

## ICB expected range summary

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

ICB		Expected range classification	
QHM	NHS North East and North Cumbria Integrated Care Board	44	17