

# **Cancer Patient Experience Survey**

2021 Results

## **Cumbria and North East**

Published July 2022

## Executive Summary

### Questions Above Expected Range

	Case Mix Adjusted Scores			England Score
	2021 Score	Lower Expected Range	Upper Expected Range	
Q2. Patient only spoke to primary care professional once or twice before cancer diagnosis	<b>79%</b>	75%	79%	<b>77%</b>
Q12. Patient was told they could have a family member, carer or friend with them when told diagnosis	<b>76%</b>	67%	76%	<b>71%</b>
Q15. Patient was definitely told about their diagnosis in an appropriate place	<b>86%</b>	83%	85%	<b>84%</b>
Q22. Family and/or carers were definitely involved as much as the patient wanted them to be in decisions about treatment options	<b>77%</b>	72%	77%	<b>75%</b>
Q23. Patient could get further advice or a second opinion before making decisions about their treatment options	<b>56%</b>	49%	55%	<b>52%</b>
Q24. Patient was definitely able to have a discussion about their needs or concerns prior to treatment	<b>74%</b>	70%	74%	<b>72%</b>
Q31. Patient had confidence and trust in all of the team looking after them during their stay in hospital	<b>84%</b>	79%	83%	<b>81%</b>
Q39. Patient was always able to discuss worries and fears with hospital staff while being treated as an outpatient or day case	<b>81%</b>	76%	81%	<b>78%</b>
Q44. Possible side effects from treatment were definitely explained in a way the patient could understand	<b>76%</b>	73%	76%	<b>74%</b>
Q45. Patient was always offered practical advice on dealing with any immediate side effects from treatment	<b>73%</b>	68%	72%	<b>70%</b>
Q47. Patient felt possible long-term side effects were definitely explained in a way they could understand in advance of their treatment	<b>63%</b>	57%	62%	<b>60%</b>
Q48. Patient was definitely able to discuss options for managing the impact of any long-term side effects	<b>57%</b>	51%	56%	<b>54%</b>
Q50. During treatment, the patient definitely got enough care and support at home from community or voluntary services	<b>56%</b>	48%	55%	<b>51%</b>
Q53. After treatment, the patient definitely could get enough emotional support at home from community or voluntary services	<b>35%</b>	29%	35%	<b>32%</b>
Q55. Patient was given enough information about the possibility and signs of cancer coming back or spreading	<b>66%</b>	61%	64%	<b>63%</b>

Cumbria and North East has no scores below expected range

## Introduction

The National Cancer Patient Experience Survey 2021 is the 11th iteration of the survey first undertaken in 2010. It has been designed to monitor progress on cancer care; to provide information to drive local quality improvements; to assist commissioners and providers of cancer care; and to inform the work of the various charities and stakeholder groups supporting cancer patients.

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

The 2021 survey involved 134 NHS Trusts. Out of 107,412 people, 59,352 people responded to the survey, yielding a response rate of 55%.

## 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 2021. The fieldwork for the survey was undertaken between October 2021 and February 2022.

As in the previous six 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. Due to the significant changes made to the questionnaire no trend data or year on year comparisons are presented in results, and comparisons to previous years' results should not be made.

### Case-mix adjustment

Both unadjusted and adjusted scores are presented in this report. Case-mix adjusted scores allows 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 ICS is performing given their patient population. The factors taken into account in this case-mix adjustment are gender, age, ethnic group, deprivation, and cancer type.

Unadjusted data should be used to see the actual responses from patients relating to the ICS. 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 presented 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%.

## Suppression

### Question-level suppression

For scores where the base size per question is <11, the score will be suppressed and replaced with an asterisk (\*). The base size does not include non-scored response options.

### Double suppression

If any group within a particular sub-group breakdown (such as the tumour group breakdown) has <11 respondents, then the figure for this particular group is suppressed and replaced with an asterisk (\*).

If there is only one group within the sub-group breakdown that has <11 respondents, and is therefore suppressed, the group with the next lowest number of respondents is also suppressed and replaced with an asterisk (\*) (regardless if it is greater than or less than 11).

### Additional suppression

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

The same rule applies to groups in each subgroup breakdown. For example, if only **one** ICS has the 85+ age group suppressed for Q25 we will need to suppress another ICS'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 ICS scored for each question in the survey compared with national results. It is aimed at helping individual ICSs to understand their performance and identify areas for local and regional improvement. Below is a description of the type of results presented within this report and how to understand them.

### Expected range charts

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

ICSs 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 ICS performs better than what ICSs 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 ICS's size and demographics.

### Comparability tables

The comparability tables show the 2021 unadjusted scores for this ICS for each scored question. The adjusted 2021 score will also be presented for each scored question along with the lower and upper expected range and national score. Scores above the upper limit of the expected range will be highlighted dark blue, scores below the lower limit of the expected range will be highlighted light blue, and scores within the lower and upper limit of the expected ranges will be highlighted grey.

### Sub-group breakdowns

Unadjusted scores are shown for tumour type, gender, age, IMD quintile and ethnicity breakdowns. Unadjusted scores for the same sub-group across different ICSs may not be comparable, as they do not account for the impact that differing patient populations might have on results.

### Tumour type tables

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

### Age group tables

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

### Gender tables

The gender 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.

### **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.

### **Further information**

This research was carried out in accordance with the international standard for organisations conducting social research (accreditation to ISO20252:2012; certificate number GB08/74322). The 2021 survey data has been produced and published in line with the Code of Practice for Official Statistics.

For more information on the methodology, please see the Technical Document. It can be viewed along with the 2021 questionnaire and survey guidance on the website at [www.ncpes.co.uk](http://www.ncpes.co.uk). For all other outputs at ICS level, please see the Excel tables and dashboards at [www.ncpes.co.uk](http://www.ncpes.co.uk).

## Response Rate

### Overall Response Rate

3,811 patients responded out of a total of 6,664 patients, resulting in a response rate of 57%.

	Sample Size	Adjusted Sample	Completed	Response Rate
Overall response rate	7,059	6,664	3,811	57%
National	113,516	107,412	59,352	55%

### Respondents by Survey Type

	Number of Respondents
Paper	3,190
Online	616
Phone	5
Translation Service	0
<b>Total</b>	<b>3,811</b>

### Respondents by Tumour Group

	Number of Respondents
Brain / CNS	22
Breast	858
Colorectal / LGT	542
Gynaecological	159
Haematological	469
Head and Neck	102
Lung	308
Prostate	305
Sarcoma	21
Skin	76
Upper Gastro	195
Urological	341
Other	413
<b>Total</b>	<b>3,811</b>

### Respondents by Age and Gender

Respondents year of birth has been used to determine age. This information has been amalgamated into 8 age bands. The age and gender distribution for the ICS was as follows:

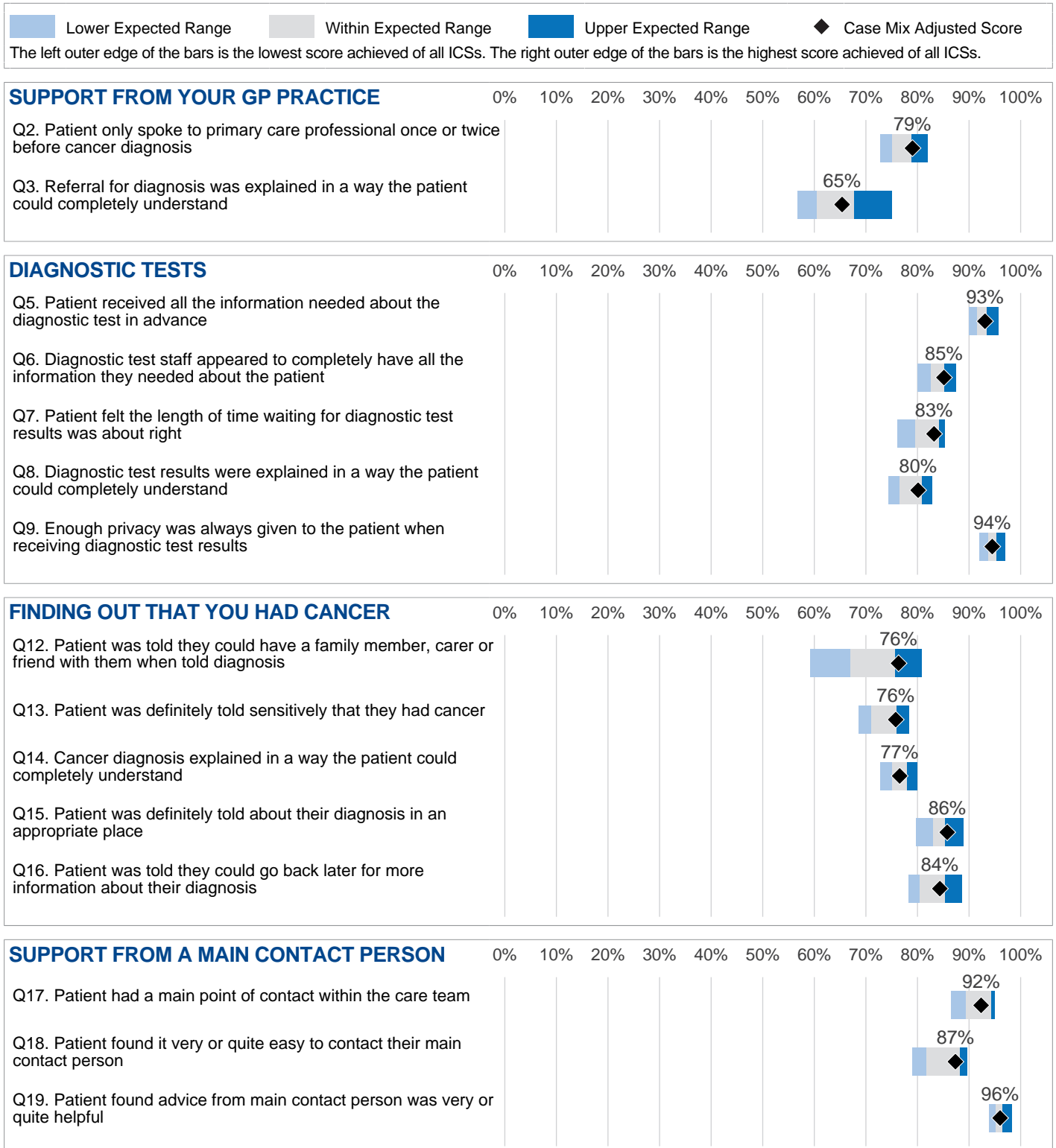
	Age 16-24	Age 25-34	Age 35-44	Age 45-54	Age 55-64	Age 65-74	Age 75-84	Age 85+	Total
Female	4	20	82	239	525	641	407	55	1,973
Male	3	10	22	74	336	686	465	66	1,662
Non-binary	0	0	0	0	0	0	0	0	0
Prefer to self-describe	0	0	0	0	0	0	0	0	0
Prefer not to say	0	0	0	0	1	0	1	0	2
Not given	0	1	3	9	27	64	50	20	174
<b>Total</b>	<b>7</b>	<b>31</b>	<b>107</b>	<b>322</b>	<b>889</b>	<b>1,391</b>	<b>923</b>	<b>141</b>	<b>3,811</b>

## Respondents by Ethnicity

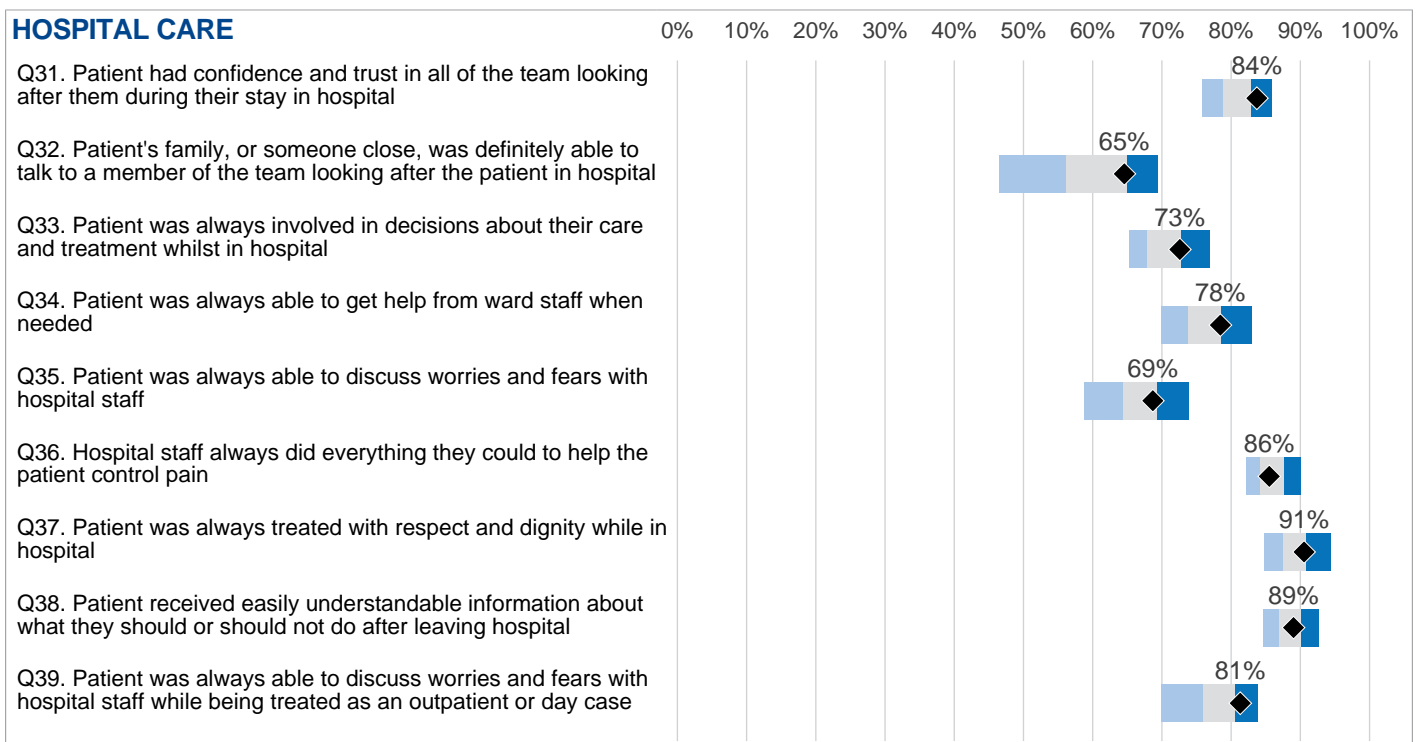
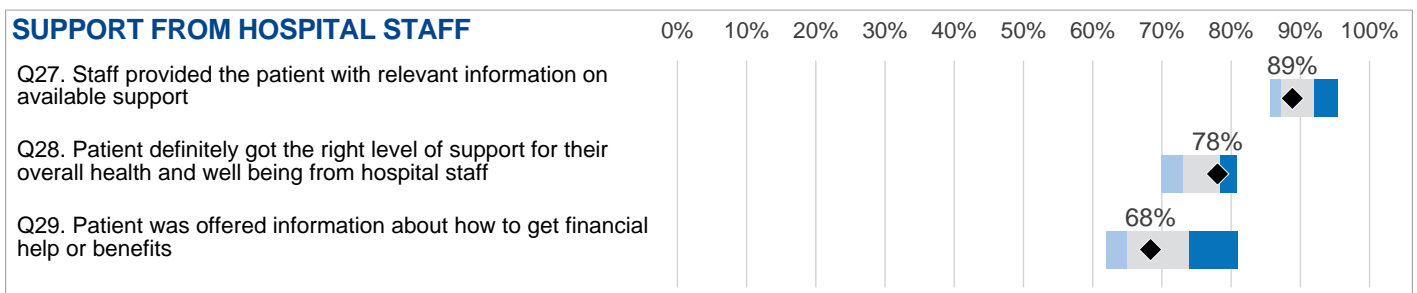
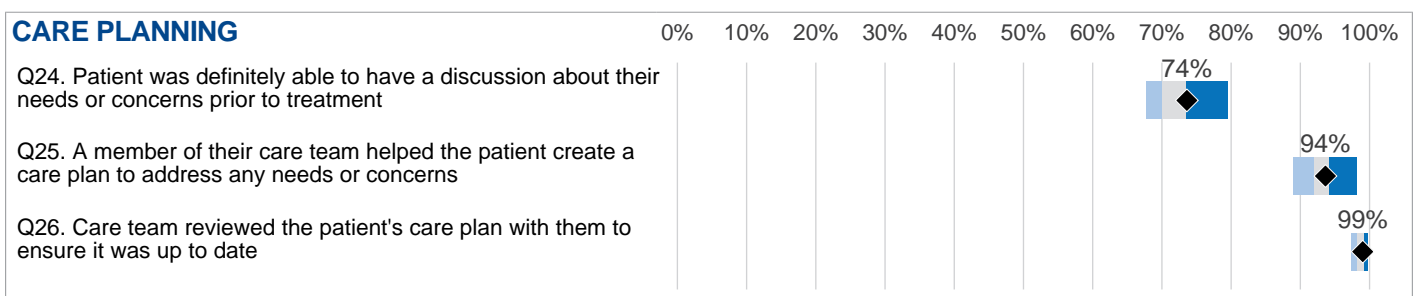
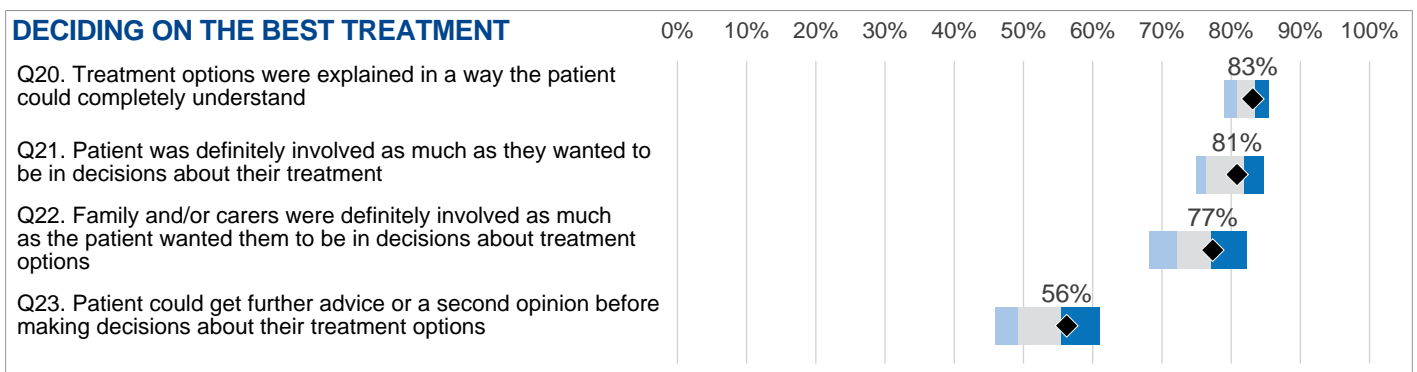
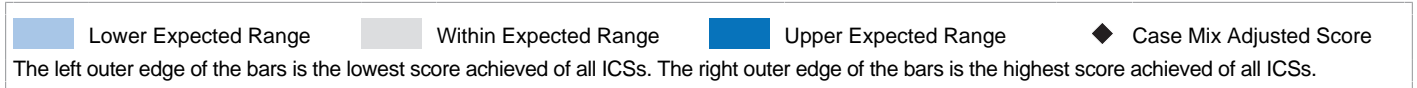
	Number of Respondents
<b>White</b>	
English / Welsh / Scottish / Northern Irish / British	3,470
Irish	12
Gypsy or Irish Traveller	1
Any other White background	44
<b>Mixed / Multiple Ethnic Groups</b>	
White and Black Caribbean	7
White and Black African	1
White and Asian	5
Any other Mixed / multiple ethnic background	6
<b>Asian or Asian British</b>	
Indian	6
Pakistani	9
Bangladeshi	3
Chinese	3
Any other Asian background	7
<b>Black / African / Caribbean / Black British</b>	
African	5
Caribbean	3
Any other Black / African / Caribbean background	1
<b>Other Ethnic Group</b>	
Arab	1
Any other ethnic background	4
<b>Not given</b>	
Not given	223
<b>Total</b>	<b>3,811</b>



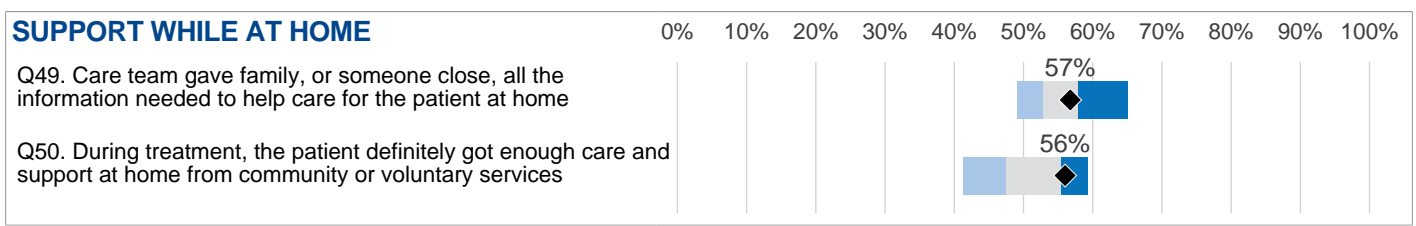
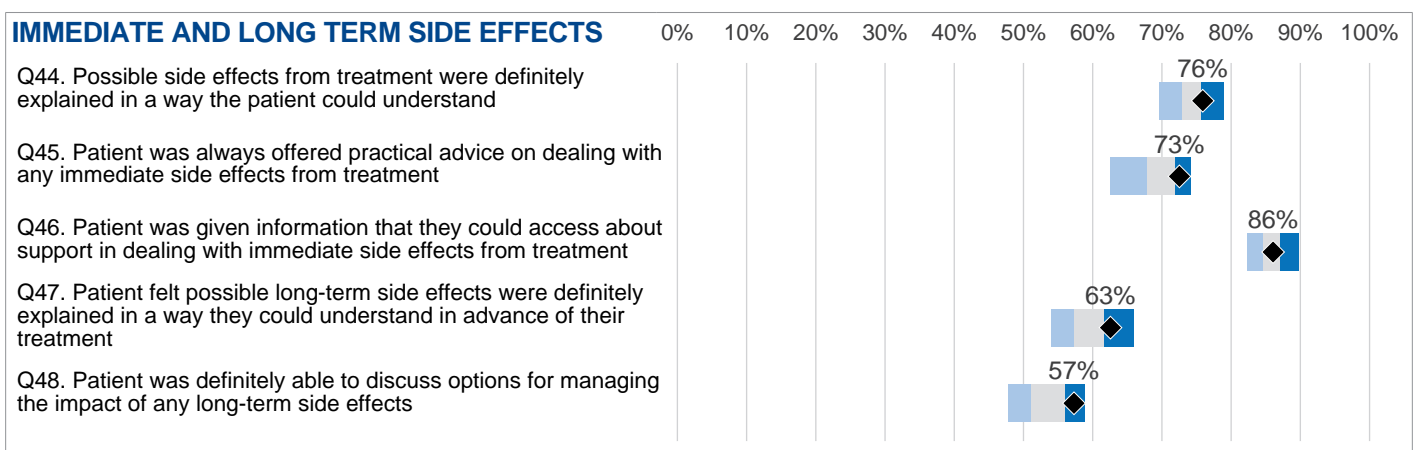
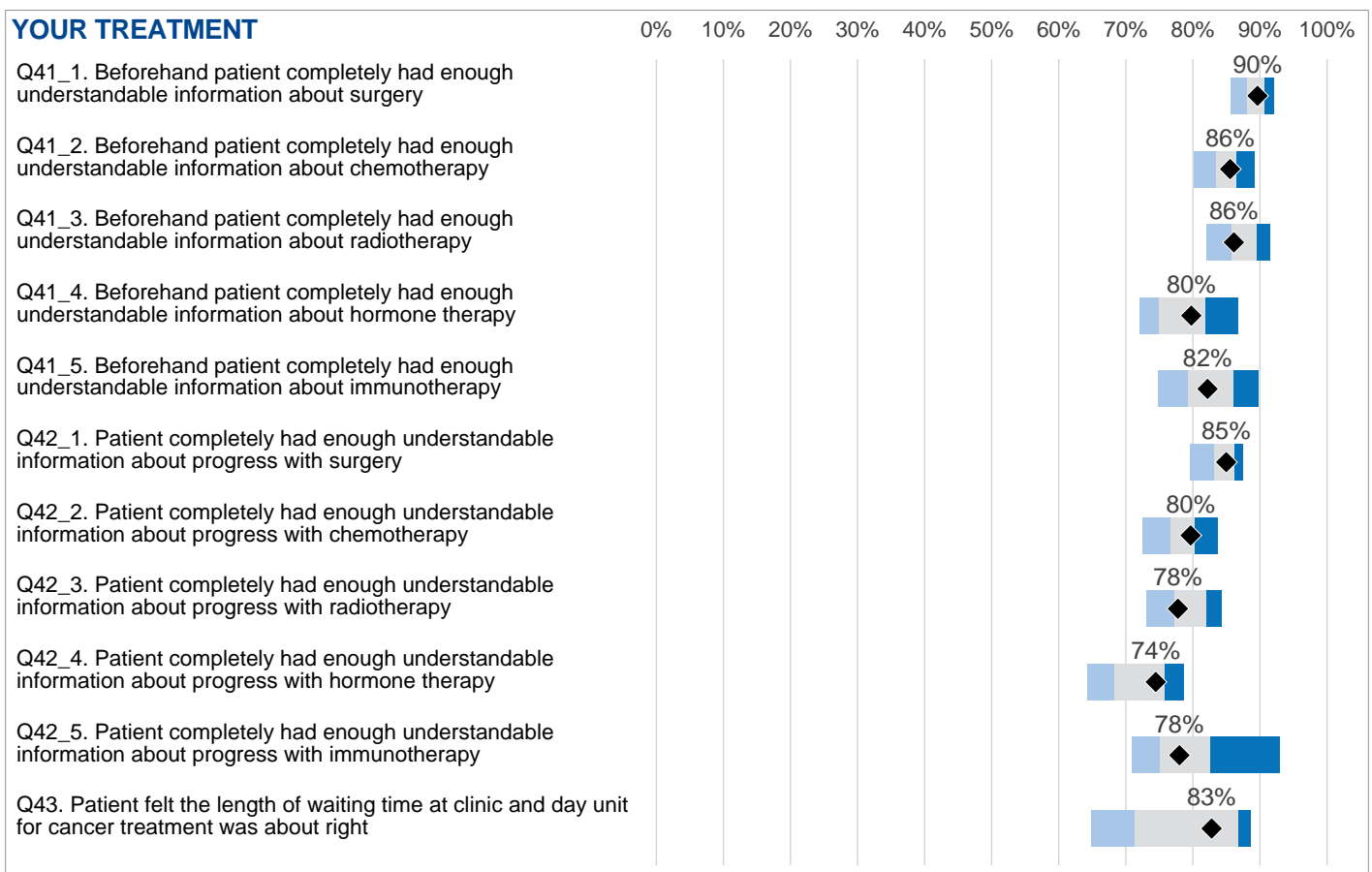
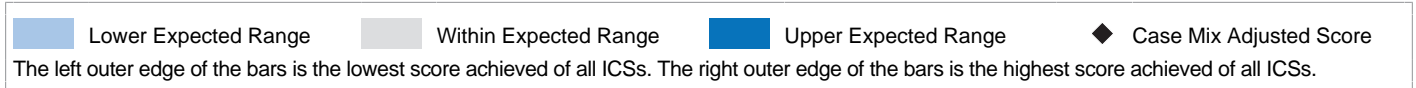
## Expected Range Charts



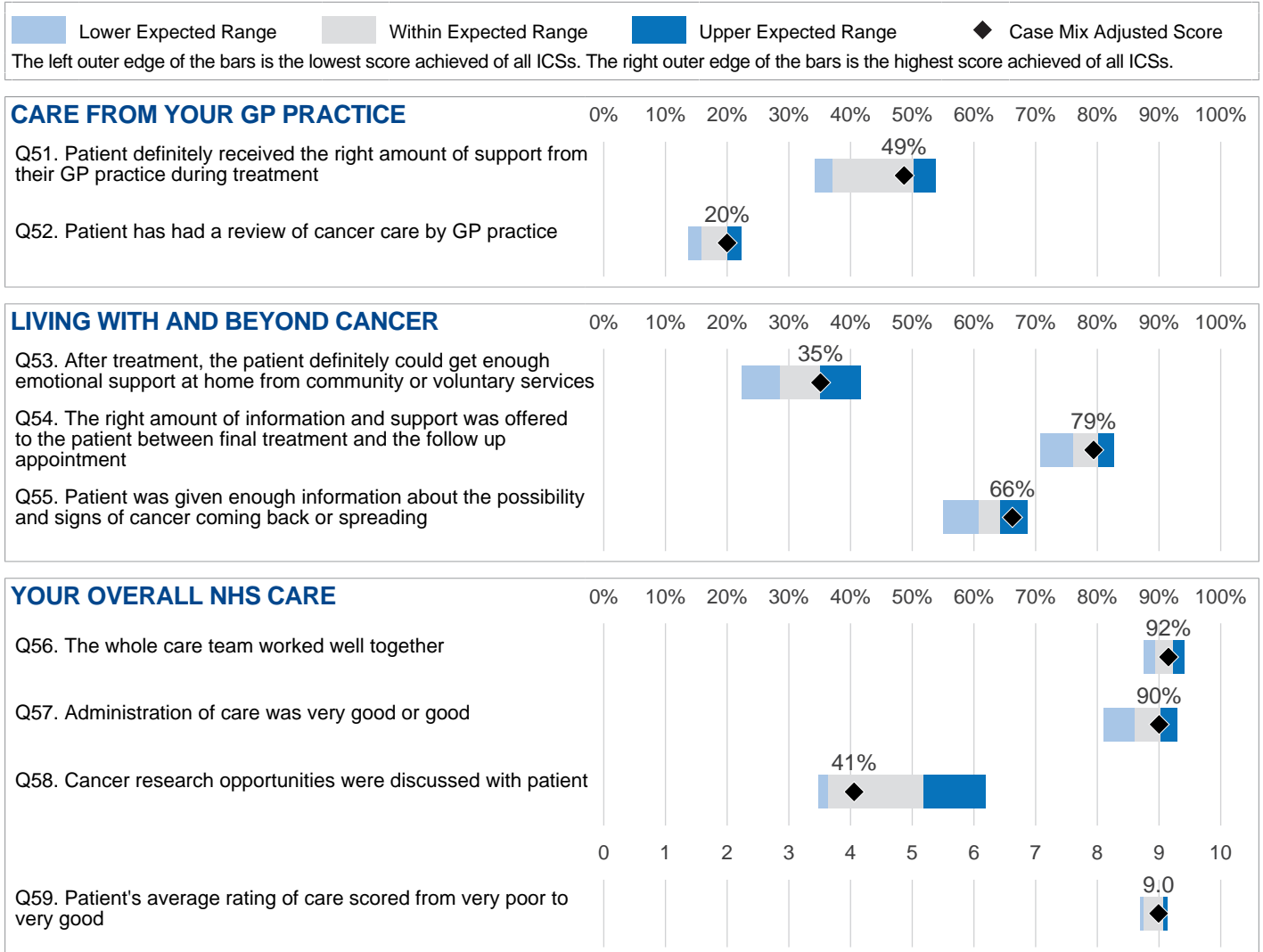
## Expected Range Charts



## Expected Range Charts






## Expected Range Charts



## Comparability tables

\* Indicates where a score has been suppressed to prevent individuals and their responses being identifiable in the data.

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

SUPPORT FROM YOUR GP PRACTICE	Unadjusted Scores		Case Mix Adjusted Scores			England Score
	2021 n	2021 Score	2021 Score	Lower Expected Range	Upper Expected Range	
Q2. Patient only spoke to primary care professional once or twice before cancer diagnosis	1851	<b>78%</b>	<b>79%</b>	75%	79%	<b>77%</b>
Q3. Referral for diagnosis was explained in a way the patient could completely understand	2504	<b>64%</b>	<b>65%</b>	61%	68%	<b>64%</b>




DIAGNOSTIC TESTS	Unadjusted Scores		Case Mix Adjusted Scores			England Score
	2021 n	2021 Score	2021 Score	Lower Expected Range	Upper Expected Range	
Q5. Patient received all the information needed about the diagnostic test in advance	3025	<b>93%</b>	<b>93%</b>	92%	93%	<b>93%</b>
Q6. Diagnostic test staff appeared to completely have all the information they needed about the patient	3182	<b>85%</b>	<b>85%</b>	83%	85%	<b>84%</b>
Q7. Patient felt the length of time waiting for diagnostic test results was about right	3195	<b>84%</b>	<b>83%</b>	80%	84%	<b>82%</b>
Q8. Diagnostic test results were explained in a way the patient could completely understand	3207	<b>80%</b>	<b>80%</b>	77%	81%	<b>79%</b>
Q9. Enough privacy was always given to the patient when receiving diagnostic test results	3201	<b>94%</b>	<b>94%</b>	94%	95%	<b>94%</b>

FINDING OUT THAT YOU HAD CANCER	Unadjusted Scores		Case Mix Adjusted Scores			England Score
	2021 n	2021 Score	2021 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	3530	<b>77%</b>	<b>76%</b>	67%	76%	<b>71%</b>
Q13. Patient was definitely told sensitively that they had cancer	3766	<b>76%</b>	<b>76%</b>	71%	76%	<b>73%</b>
Q14. Cancer diagnosis explained in a way the patient could completely understand	3775	<b>76%</b>	<b>77%</b>	75%	78%	<b>76%</b>
Q15. Patient was definitely told about their diagnosis in an appropriate place	3751	<b>86%</b>	<b>86%</b>	83%	85%	<b>84%</b>
Q16. Patient was told they could go back later for more information about their diagnosis	3346	<b>84%</b>	<b>84%</b>	81%	85%	<b>83%</b>

SUPPORT FROM A MAIN CONTACT PERSON	Unadjusted Scores		Case Mix Adjusted Scores			England Score
	2021 n	2021 Score	2021 Score	Lower Expected Range	Upper Expected Range	
Q17. Patient had a main point of contact within the care team	3638	<b>92%</b>	<b>92%</b>	89%	94%	<b>92%</b>
Q18. Patient found it very or quite easy to contact their main contact person	3074	<b>87%</b>	<b>87%</b>	82%	88%	<b>85%</b>
Q19. Patient found advice from main contact person was very or quite helpful	3212	<b>96%</b>	<b>96%</b>	95%	97%	<b>96%</b>

## Comparability tables

\* Indicates where a score has been suppressed to prevent individuals and their responses being identifiable in the data.

	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
	2021 n	2021 Score	2021 Score	Lower Expected Range	Upper Expected Range	
Q20. Treatment options were explained in a way the patient could completely understand	3579	<b>83%</b>	<b>83%</b>	81%	83%	<b>82%</b>
Q21. Patient was definitely involved as much as they wanted to be in decisions about their treatment	3732	<b>81%</b>	<b>81%</b>	76%	82%	<b>79%</b>
Q22. Family and/or carers were definitely involved as much as the patient wanted them to be in decisions about treatment options	3173	<b>77%</b>	<b>77%</b>	72%	77%	<b>75%</b>
Q23. Patient could get further advice or a second opinion before making decisions about their treatment options	1731	<b>57%</b>	<b>56%</b>	49%	55%	<b>52%</b>




CARE PLANNING	Unadjusted Scores		Case Mix Adjusted Scores			England Score
	2021 n	2021 Score	2021 Score	Lower Expected Range	Upper Expected Range	
Q24. Patient was definitely able to have a discussion about their needs or concerns prior to treatment	3410	<b>74%</b>	<b>74%</b>	70%	74%	<b>72%</b>
Q25. A member of their care team helped the patient create a care plan to address any needs or concerns	2033	<b>94%</b>	<b>94%</b>	92%	94%	<b>93%</b>
Q26. Care team reviewed the patient's care plan with them to ensure it was up to date	1666	<b>99%</b>	<b>99%</b>	98%	99%	<b>99%</b>

SUPPORT FROM HOSPITAL STAFF	Unadjusted Scores		Case Mix Adjusted Scores			England Score
	2021 n	2021 Score	2021 Score	Lower Expected Range	Upper Expected Range	
Q27. Staff provided the patient with relevant information on available support	3022	<b>89%</b>	<b>89%</b>	87%	92%	<b>90%</b>
Q28. Patient definitely got the right level of support for their overall health and well being from hospital staff	3736	<b>78%</b>	<b>78%</b>	73%	78%	<b>76%</b>
Q29. Patient was offered information about how to get financial help or benefits	2076	<b>68%</b>	<b>68%</b>	65%	74%	<b>69%</b>

HOSPITAL CARE	Unadjusted Scores		Case Mix Adjusted Scores			England Score
	2021 n	2021 Score	2021 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	1613	<b>84%</b>	<b>84%</b>	79%	83%	<b>81%</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	1307	<b>65%</b>	<b>65%</b>	56%	65%	<b>61%</b>
Q33. Patient was always involved in decisions about their care and treatment whilst in hospital	1582	<b>73%</b>	<b>73%</b>	68%	73%	<b>70%</b>
Q34. Patient was always able to get help from ward staff when needed	1574	<b>79%</b>	<b>78%</b>	74%	79%	<b>76%</b>
Q35. Patient was always able to discuss worries and fears with hospital staff	1539	<b>69%</b>	<b>69%</b>	64%	69%	<b>67%</b>
Q36. Hospital staff always did everything they could to help the patient control pain	1427	<b>85%</b>	<b>86%</b>	84%	88%	<b>86%</b>
Q37. Patient was always treated with respect and dignity while in hospital	1604	<b>91%</b>	<b>91%</b>	88%	91%	<b>89%</b>
Q38. Patient received easily understandable information about what they should or should not do after leaving hospital	1575	<b>89%</b>	<b>89%</b>	87%	90%	<b>89%</b>
Q39. Patient was always able to discuss worries and fears with hospital staff while being treated as an outpatient or day case	3264	<b>81%</b>	<b>81%</b>	76%	81%	<b>78%</b>

## Comparability tables

\* Indicates where a score has been suppressed to prevent individuals and their responses being identifiable in the data.

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

YOUR TREATMENT	Unadjusted Scores		Case Mix Adjusted Scores			England Score
	2021 n	2021 Score	2021 Score	Lower Expected Range	Upper Expected Range	
Q41_1. Beforehand patient completely had enough understandable information about surgery	2044	<b>90%</b>	<b>90%</b>	88%	91%	<b>89%</b>
Q41_2. Beforehand patient completely had enough understandable information about chemotherapy	1976	<b>86%</b>	<b>86%</b>	83%	87%	<b>85%</b>
Q41_3. Beforehand patient completely had enough understandable information about radiotherapy	1103	<b>86%</b>	<b>86%</b>	86%	90%	<b>88%</b>
Q41_4. Beforehand patient completely had enough understandable information about hormone therapy	552	<b>80%</b>	<b>80%</b>	75%	82%	<b>78%</b>
Q41_5. Beforehand patient completely had enough understandable information about immunotherapy	476	<b>83%</b>	<b>82%</b>	79%	86%	<b>83%</b>
Q42_1. Patient completely had enough understandable information about progress with surgery	2031	<b>85%</b>	<b>85%</b>	83%	86%	<b>85%</b>
Q42_2. Patient completely had enough understandable information about progress with chemotherapy	1949	<b>80%</b>	<b>80%</b>	77%	80%	<b>79%</b>
Q42_3. Patient completely had enough understandable information about progress with radiotherapy	1099	<b>78%</b>	<b>78%</b>	77%	82%	<b>80%</b>
Q42_4. Patient completely had enough understandable information about progress with hormone therapy	548	<b>75%</b>	<b>74%</b>	68%	76%	<b>72%</b>
Q42_5. Patient completely had enough understandable information about progress with immunotherapy	470	<b>79%</b>	<b>78%</b>	75%	83%	<b>79%</b>
Q43. Patient felt the length of waiting time at clinic and day unit for cancer treatment was about right	3685	<b>83%</b>	<b>83%</b>	71%	87%	<b>79%</b>




IMMEDIATE AND LONG TERM SIDE EFFECTS	Unadjusted Scores		Case Mix Adjusted Scores			England Score
	2021 n	2021 Score	2021 Score	Lower Expected Range	Upper Expected Range	
Q44. Possible side effects from treatment were definitely explained in a way the patient could understand	3602	<b>76%</b>	<b>76%</b>	73%	76%	<b>74%</b>
Q45. Patient was always offered practical advice on dealing with any immediate side effects from treatment	3429	<b>73%</b>	<b>73%</b>	68%	72%	<b>70%</b>
Q46. Patient was given information that they could access about support in dealing with immediate side effects from treatment	2773	<b>86%</b>	<b>86%</b>	85%	87%	<b>86%</b>
Q47. Patient felt possible long-term side effects were definitely explained in a way they could understand in advance of their treatment	3366	<b>63%</b>	<b>63%</b>	57%	62%	<b>60%</b>
Q48. Patient was definitely able to discuss options for managing the impact of any long-term side effects	2961	<b>58%</b>	<b>57%</b>	51%	56%	<b>54%</b>

SUPPORT WHILE AT HOME	Unadjusted Scores		Case Mix Adjusted Scores			England Score
	2021 n	2021 Score	2021 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	2520	<b>57%</b>	<b>57%</b>	53%	58%	<b>55%</b>
Q50. During treatment, the patient definitely got enough care and support at home from community or voluntary services	1579	<b>56%</b>	<b>56%</b>	48%	55%	<b>51%</b>

CARE FROM YOUR GP PRACTICE	Unadjusted Scores		Case Mix Adjusted Scores			England Score
	2021 n	2021 Score	2021 Score	Lower Expected Range	Upper Expected Range	
Q51. Patient definitely received the right amount of support from their GP practice during treatment	2248	<b>48%</b>	<b>49%</b>	37%	50%	<b>44%</b>
Q52. Patient has had a review of cancer care by GP practice	3605	<b>20%</b>	<b>20%</b>	16%	20%	<b>18%</b>

## Comparability tables

\* Indicates where a score has been suppressed to prevent individuals and their responses being identifiable in the data.

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

LIVING WITH AND BEYOND CANCER	Unadjusted Scores		Case Mix Adjusted Scores			England Score
	2021 n	2021 Score	2021 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	820	<b>36%</b>	<b>35%</b>	29%	35%	<b>32%</b>
Q54. The right amount of information and support was offered to the patient between final treatment and the follow up appointment	1598	<b>79%</b>	<b>79%</b>	76%	80%	<b>78%</b>
Q55. Patient was given enough information about the possibility and signs of cancer coming back or spreading	2936	<b>66%</b>	<b>66%</b>	61%	64%	<b>63%</b>

YOUR OVERALL NHS CARE	Unadjusted Scores		Case Mix Adjusted Scores			England Score
	2021 n	2021 Score	2021 Score	Lower Expected Range	Upper Expected Range	
Q56. The whole care team worked well together	3606	<b>92%</b>	<b>92%</b>	89%	92%	<b>91%</b>
Q57. Administration of care was very good or good	3713	<b>90%</b>	<b>90%</b>	86%	90%	<b>88%</b>
Q58. Cancer research opportunities were discussed with patient	2045	<b>41%</b>	<b>41%</b>	36%	52%	<b>44%</b>
Q59. Patient's average rating of care scored from very poor to very good	3621	<b>9.0</b>	<b>9.0</b>	8.8	9.1	<b>8.9</b>



## Tumour type tables

\* Indicates where a score has been suppressed because there are less than 11 responses. n.a. Indicates that there were no respondents for that tumour group.

	Tumour Type													
	Brain / CNS	Breast	Colorectal / LGT	Gynaecological	Haematological	Head and Neck	Lung	Prostate	Sarcoma	Skin	Upper Gastro	Urological	Other	All Cancers
Q2. Patient only spoke to primary care professional once or twice before cancer diagnosis	55%	93%	78%	81%	63%	75%	71%	80%	69%	90%	68%	75%	74%	<b>78%</b>
Q3. Referral for diagnosis was explained in a way the patient could completely understand	*	83%	66%	66%	54%	60%	51%	67%	*	66%	47%	55%	63%	<b>64%</b>

	Tumour Type													
	Brain / CNS	Breast	Colorectal / LGT	Gynaecological	Haematological	Head and Neck	Lung	Prostate	Sarcoma	Skin	Upper Gastro	Urological	Other	All Cancers
Q5. Patient received all the information needed about the diagnostic test in advance	77%	94%	94%	95%	93%	93%	93%	90%	100%	97%	90%	93%	92%	<b>93%</b>
Q6. Diagnostic test staff appeared to completely have all the information they needed about the patient	60%	86%	86%	83%	86%	86%	88%	86%	79%	82%	82%	83%	84%	<b>85%</b>
Q7. Patient felt the length of time waiting for diagnostic test results was about right	73%	82%	87%	82%	86%	90%	84%	85%	89%	72%	84%	82%	78%	<b>84%</b>
Q8. Diagnostic test results were explained in a way the patient could completely understand	44%	84%	81%	80%	79%	85%	80%	80%	79%	80%	76%	78%	78%	<b>80%</b>
Q9. Enough privacy was always given to the patient when receiving diagnostic test results	81%	96%	95%	92%	96%	96%	93%	96%	95%	89%	91%	93%	93%	<b>94%</b>

	Tumour Type													
	Brain / CNS	Breast	Colorectal / LGT	Gynaecological	Haematological	Head and Neck	Lung	Prostate	Sarcoma	Skin	Upper Gastro	Urological	Other	All Cancers
Q12. Patient was told they could have a family member, carer or friend with them when told diagnosis	95%	81%	80%	61%	75%	76%	80%	77%	70%	51%	83%	67%	77%	<b>77%</b>
Q13. Patient was definitely told sensitively that they had cancer	67%	81%	77%	69%	77%	83%	77%	74%	67%	69%	72%	66%	74%	<b>76%</b>
Q14. Cancer diagnosis explained in a way the patient could completely understand	50%	81%	80%	76%	70%	85%	78%	74%	71%	74%	73%	72%	74%	<b>76%</b>
Q15. Patient was definitely told about their diagnosis in an appropriate place	71%	90%	86%	82%	85%	88%	86%	89%	81%	78%	81%	81%	84%	<b>86%</b>
Q16. Patient was told they could go back later for more information about their diagnosis	75%	89%	82%	82%	86%	92%	84%	89%	78%	87%	80%	75%	81%	<b>84%</b>

## Tumour type tables

\* Indicates where a score has been suppressed because there are less than 11 responses. n.a. Indicates that there were no respondents for that tumour group.

	Tumour Type													
	Brain / CNS	Breast	Colorectal / LGT	Gynaecological	Haematological	Head and Neck	Lung	Prostate	Sarcoma	Skin	Upper Gastro	Urological	Other	All Cancers
Q17. Patient had a main point of contact within the care team	100%	96%	91%	97%	90%	96%	95%	91%	90%	89%	90%	85%	92%	92%
Q18. Patient found it very or quite easy to contact their main contact person	95%	88%	89%	83%	90%	95%	90%	82%	95%	86%	86%	84%	86%	87%
Q19. Patient found advice from main contact person was very or quite helpful	95%	95%	98%	96%	97%	99%	97%	93%	100%	94%	94%	95%	96%	96%

	Tumour Type													
	Brain / CNS	Breast	Colorectal / LGT	Gynaecological	Haematological	Head and Neck	Lung	Prostate	Sarcoma	Skin	Upper Gastro	Urological	Other	All Cancers
Q20. Treatment options were explained in a way the patient could completely understand	64%	85%	85%	89%	83%	85%	86%	78%	81%	86%	81%	79%	82%	83%
Q21. Patient was definitely involved as much as they wanted to be in decisions about their treatment	73%	80%	82%	82%	83%	88%	82%	81%	80%	84%	83%	74%	80%	81%
Q22. Family and/or carers were definitely involved as much as the patient wanted them to be in decisions about treatment options	60%	75%	78%	70%	81%	87%	85%	77%	83%	67%	78%	73%	78%	77%
Q23. Patient could get further advice or a second opinion before making decisions about their treatment options	*	54%	58%	49%	64%	76%	62%	49%	*	48%	60%	54%	53%	57%

	Tumour Type													
	Brain / CNS	Breast	Colorectal / LGT	Gynaecological	Haematological	Head and Neck	Lung	Prostate	Sarcoma	Skin	Upper Gastro	Urological	Other	All Cancers
Q24. Patient was definitely able to have a discussion about their needs or concerns prior to treatment	50%	74%	74%	73%	78%	81%	74%	71%	84%	74%	78%	66%	73%	74%
Q25. A member of their care team helped the patient create a care plan to address any needs or concerns	92%	92%	95%	96%	97%	97%	92%	93%	100%	91%	91%	96%	91%	94%
Q26. Care team reviewed the patient's care plan with them to ensure it was up to date	100%	98%	99%	97%	100%	100%	100%	98%	100%	100%	99%	100%	98%	99%

	Tumour Type													
	Brain / CNS	Breast	Colorectal / LGT	Gynaecological	Haematological	Head and Neck	Lung	Prostate	Sarcoma	Skin	Upper Gastro	Urological	Other	All Cancers
Q27. Staff provided the patient with relevant information on available support	84%	89%	89%	93%	91%	98%	89%	91%	100%	88%	86%	83%	84%	89%
Q28. Patient definitely got the right level of support for their overall health and well being from hospital staff	73%	75%	79%	75%	86%	84%	83%	74%	86%	74%	82%	75%	76%	78%
Q29. Patient was offered information about how to get financial help or benefits	64%	70%	60%	72%	72%	75%	80%	61%	85%	76%	77%	48%	66%	68%

## Tumour type tables

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	Tumour Type													All Cancers
	Brain/ CNS	Breast	Colorectal/ LGT	Gynaecological	Haematological	Head and Neck	Lung	Prostate	Sarcoma	Skin	Upper Gastro	Urological	Other	
Q31. Patient had confidence and trust in all of the team looking after them during their stay in hospital	69%	80%	87%	81%	87%	87%	83%	82%	84%	94%	86%	84%	82%	<b>84%</b>
Q32. Patient's family, or someone close, was definitely able to talk to a member of the team looking after the patient in hospital	*	58%	65%	68%	74%	67%	73%	44%	81%	*	74%	53%	70%	<b>65%</b>
Q33. Patient was always involved in decisions about their care and treatment whilst in hospital	50%	73%	70%	67%	77%	69%	75%	70%	74%	88%	72%	77%	74%	<b>73%</b>
Q34. Patient was always able to get help from ward staff when needed	50%	77%	77%	73%	84%	78%	84%	73%	84%	100%	80%	81%	78%	<b>79%</b>
Q35. Patient was always able to discuss worries and fears with hospital staff	*	61%	70%	70%	78%	73%	73%	62%	79%	*	76%	66%	69%	<b>69%</b>
Q36. Hospital staff always did everything they could to help the patient control pain	77%	84%	88%	78%	94%	82%	86%	82%	82%	92%	87%	85%	83%	<b>85%</b>
Q37. Patient was always treated with respect and dignity while in hospital	69%	89%	90%	84%	95%	93%	90%	88%	89%	100%	92%	95%	90%	<b>91%</b>
Q38. Patient received easily understandable information about what they should or should not do after leaving hospital	69%	90%	88%	88%	94%	92%	89%	78%	95%	94%	90%	91%	88%	<b>89%</b>
Q39. Patient was always able to discuss worries and fears with hospital staff while being treated as an outpatient or day case	46%	79%	80%	80%	87%	85%	87%	82%	83%	80%	80%	80%	80%	<b>81%</b>

	Tumour Type													All Cancers
	Brain/ CNS	Breast	Colorectal/ LGT	Gynaecological	Haematological	Head and Neck	Lung	Prostate	Sarcoma	Skin	Upper Gastro	Urological	Other	
Q41_1. Beforehand patient completely had enough understandable information about surgery	67%	92%	90%	94%	87%	91%	92%	84%	90%	85%	91%	87%	90%	<b>90%</b>
Q41_2. Beforehand patient completely had enough understandable information about chemotherapy	65%	84%	83%	95%	87%	92%	88%	82%	*	*	87%	81%	89%	<b>86%</b>
Q41_3. Beforehand patient completely had enough understandable information about radiotherapy	60%	87%	83%	91%	89%	87%	89%	84%	*	*	83%	73%	90%	<b>86%</b>
Q41_4. Beforehand patient completely had enough understandable information about hormone therapy	*	81%	*	*	*	*	*	81%	n.a.	n.a.	*	*	72%	<b>80%</b>
Q41_5. Beforehand patient completely had enough understandable information about immunotherapy	n.a.	80%	73%	*	75%	*	91%	*	*	77%	*	83%	87%	<b>83%</b>
Q42_1. Patient completely had enough understandable information about progress with surgery	60%	87%	86%	88%	84%	86%	87%	82%	100%	80%	91%	81%	79%	<b>85%</b>
Q42_2. Patient completely had enough understandable information about progress with chemotherapy	67%	78%	78%	86%	84%	85%	78%	80%	*	*	84%	78%	78%	<b>80%</b>
Q42_3. Patient completely had enough understandable information about progress with radiotherapy	60%	82%	74%	90%	69%	89%	74%	67%	*	*	80%	65%	76%	<b>78%</b>
Q42_4. Patient completely had enough understandable information about progress with hormone therapy	*	72%	*	*	*	*	*	79%	n.a.	n.a.	*	*	76%	<b>75%</b>
Q42_5. Patient completely had enough understandable information about progress with immunotherapy	n.a.	72%	80%	*	79%	*	83%	*	*	75%	*	83%	80%	<b>79%</b>
Q43. Patient felt the length of waiting time at clinic and day unit for cancer treatment was about right	68%	79%	87%	83%	85%	88%	84%	88%	81%	80%	81%	87%	77%	<b>83%</b>

## Tumour type tables

\* Indicates where a score has been suppressed because there are less than 11 responses. n.a. Indicates that there were no respondents for that tumour group.

	Tumour Type													
	Brain/ CNS	Breast	Colorectal/ LGT	Gynaecological	Haematological	Head and Neck	Lung	Prostate	Sarcoma	Skin	Upper Gastro	Urological	Other	All Cancers
Q44. Possible side effects from treatment were definitely explained in a way the patient could understand	50%	74%	77%	82%	75%	80%	81%	75%	85%	81%	78%	74%	75%	<b>76%</b>
Q45. Patient was always offered practical advice on dealing with any immediate side effects from treatment	50%	71%	75%	74%	76%	85%	80%	66%	68%	76%	71%	73%	71%	<b>73%</b>
Q46. Patient was given information that they could access about support in dealing with immediate side effects from treatment	79%	85%	86%	88%	90%	87%	90%	85%	100%	80%	86%	84%	85%	<b>86%</b>
Q47. Patient felt possible long-term side effects were definitely explained in a way they could understand in advance of their treatment	48%	61%	62%	65%	63%	76%	68%	67%	70%	71%	66%	59%	58%	<b>63%</b>
Q48. Patient was definitely able to discuss options for managing the impact of any long-term side effects	50%	52%	61%	55%	65%	76%	63%	59%	61%	67%	57%	54%	54%	<b>58%</b>

	Tumour Type													
	Brain/ CNS	Breast	Colorectal/ LGT	Gynaecological	Haematological	Head and Neck	Lung	Prostate	Sarcoma	Skin	Upper Gastro	Urological	Other	All Cancers
Q49. Care team gave family, or someone close, all the information needed to help care for the patient at home	47%	51%	57%	55%	64%	77%	62%	56%	67%	55%	61%	53%	54%	<b>57%</b>
Q50. During treatment, the patient definitely got enough care and support at home from community or voluntary services	50%	49%	56%	54%	58%	75%	62%	54%	64%	61%	60%	53%	54%	<b>56%</b>

	Tumour Type													
	Brain/ CNS	Breast	Colorectal/ LGT	Gynaecological	Haematological	Head and Neck	Lung	Prostate	Sarcoma	Skin	Upper Gastro	Urological	Other	All Cancers
Q51. Patient definitely received the right amount of support from their GP practice during treatment	75%	48%	45%	56%	47%	69%	44%	53%	55%	45%	42%	43%	47%	<b>48%</b>
Q52. Patient has had a review of cancer care by GP practice	15%	19%	23%	20%	17%	33%	19%	20%	30%	14%	22%	18%	20%	<b>20%</b>

## Tumour type tables

\* Indicates where a score has been suppressed because there are less than 11 responses. n.a. Indicates that there were no respondents for that tumour group.

	Tumour Type													All Cancers
	Brain / CNS	Breast	Colorectal / LGT	Gynaecological	Haematological	Head and Neck	Lung	Prostate	Sarcoma	Skin	Upper Gastro	Urological	Other	
Q53. After treatment, the patient definitely could get enough emotional support at home from community or voluntary services	*	30%	45%	41%	38%	51%	42%	28%	*	*	21%	32%	30%	<b>36%</b>
Q54. The right amount of information and support was offered to the patient between final treatment and the follow up appointment	*	76%	76%	87%	85%	96%	83%	70%	*	81%	84%	73%	79%	<b>79%</b>
Q55. Patient was given enough information about the possibility and signs of cancer coming back or spreading	53%	59%	62%	60%	79%	74%	69%	67%	70%	85%	60%	68%	67%	<b>66%</b>

	Tumour Type													All Cancers
	Brain / CNS	Breast	Colorectal / LGT	Gynaecological	Haematological	Head and Neck	Lung	Prostate	Sarcoma	Skin	Upper Gastro	Urological	Other	
Q56. The whole care team worked well together	89%	92%	92%	95%	95%	99%	93%	86%	100%	91%	89%	91%	87%	<b>92%</b>
Q57. Administration of care was very good or good	95%	92%	90%	94%	93%	96%	91%	86%	95%	79%	89%	87%	88%	<b>90%</b>
Q58. Cancer research opportunities were discussed with patient	43%	28%	43%	47%	48%	65%	56%	43%	53%	36%	56%	30%	37%	<b>41%</b>
Q59. Patient's average rating of care scored from very poor to very good	8.5	9.0	8.9	9.1	9.3	9.4	9.1	8.8	9.3	9.1	9.0	8.9	8.8	<b>9.0</b>

## Age group tables

\* Indicates where a score has been suppressed because there are less than 11 responses. n.a. Indicates that there were no respondents for that age group.

<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	*	*	83%	76%	72%	79%	84%	88%	<b>78%</b>
Q3. Referral for diagnosis was explained in a way the patient could completely understand	*	*	79%	76%	67%	60%	62%	66%	<b>64%</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	*	*	89%	93%	94%	93%	92%	94%	<b>93%</b>
Q6. Diagnostic test staff appeared to completely have all the information they needed about the patient	*	*	79%	82%	84%	85%	87%	90%	<b>85%</b>
Q7. Patient felt the length of time waiting for diagnostic test results was about right	*	*	67%	75%	80%	86%	87%	90%	<b>84%</b>
Q8. Diagnostic test results were explained in a way the patient could completely understand	*	*	73%	74%	78%	82%	83%	81%	<b>80%</b>
Q9. Enough privacy was always given to the patient when receiving diagnostic test results	*	*	96%	92%	94%	95%	95%	94%	<b>94%</b>

<b>FINDING OUT THAT YOU HAD CANCER</b>	Age								
	16 - 24	25 - 34	35 - 44	45 - 54	55 - 64	65 - 74	75 - 84	85+	All
Q12. Patient was told they could have a family member, carer or friend with them when told diagnosis	*	*	64%	68%	73%	78%	82%	83%	<b>77%</b>
Q13. Patient was definitely told sensitively that they had cancer	*	*	67%	72%	73%	76%	80%	84%	<b>76%</b>
Q14. Cancer diagnosis explained in a way the patient could completely understand	*	*	67%	73%	75%	77%	80%	80%	<b>76%</b>
Q15. Patient was definitely told about their diagnosis in an appropriate place	*	*	79%	81%	83%	86%	89%	92%	<b>86%</b>
Q16. Patient was told they could go back later for more information about their diagnosis	*	*	87%	86%	85%	85%	81%	82%	<b>84%</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	*	*	93%	95%	93%	92%	92%	93%	<b>92%</b>
Q18. Patient found it very or quite easy to contact their main contact person	*	*	80%	82%	88%	88%	89%	87%	<b>87%</b>
Q19. Patient found advice from main contact person was very or quite helpful	*	*	96%	94%	96%	96%	97%	97%	<b>96%</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	*	*	78%	78%	83%	83%	85%	85%	<b>83%</b>
Q21. Patient was definitely involved as much as they wanted to be in decisions about their treatment	*	*	73%	73%	80%	82%	84%	78%	<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	*	*	65%	71%	76%	77%	83%	79%	<b>77%</b>
Q23. Patient could get further advice or a second opinion before making decisions about their treatment options	*	*	60%	48%	56%	55%	62%	54%	<b>57%</b>

## Age group tables

\* Indicates where a score has been suppressed because there are less than 11 responses. n.a. Indicates that there were no respondents for that age group.

CARE PLANNING	Age								
	16 - 24	25 - 34	35 - 44	45 - 54	55 - 64	65 - 74	75 - 84	85+	All
Q24. Patient was definitely able to have a discussion about their needs or concerns prior to treatment	*	*	64%	68%	74%	75%	74%	74%	<b>74%</b>
Q25. A member of their care team helped the patient create a care plan to address any needs or concerns	*	*	89%	91%	93%	95%	94%	96%	<b>94%</b>
Q26. Care team reviewed the patient's care plan with them to ensure it was up to date	*	*	98%	100%	98%	99%	99%	100%	<b>99%</b>

SUPPORT FROM HOSPITAL STAFF	Age								
	16 - 24	25 - 34	35 - 44	45 - 54	55 - 64	65 - 74	75 - 84	85+	All
Q27. Staff provided the patient with relevant information on available support	*	*	82%	85%	89%	89%	90%	86%	<b>89%</b>
Q28. Patient definitely got the right level of support for their overall health and well being from hospital staff	*	*	60%	70%	75%	79%	85%	80%	<b>78%</b>
Q29. Patient was offered information about how to get financial help or benefits	*	*	66%	73%	71%	69%	64%	51%	<b>68%</b>

HOSPITAL CARE	Age								
	16 - 24	25 - 34	35 - 44	45 - 54	55 - 64	65 - 74	75 - 84	85+	All
Q31. Patient had confidence and trust in all of the team looking after them during their stay in hospital	*	*	73%	75%	82%	86%	89%	86%	<b>84%</b>
Q32. Patient's family, or someone close, was definitely able to talk to a member of the team looking after the patient in hospital	*	*	50%	59%	61%	66%	72%	71%	<b>65%</b>
Q33. Patient was always involved in decisions about their care and treatment whilst in hospital	*	*	56%	71%	71%	73%	76%	73%	<b>73%</b>
Q34. Patient was always able to get help from ward staff when needed	*	*	69%	69%	76%	80%	83%	87%	<b>79%</b>
Q35. Patient was always able to discuss worries and fears with hospital staff	*	*	53%	56%	70%	70%	74%	73%	<b>69%</b>
Q36. Hospital staff always did everything they could to help the patient control pain	*	*	73%	83%	82%	87%	90%	91%	<b>85%</b>
Q37. Patient was always treated with respect and dignity while in hospital	*	*	83%	85%	90%	91%	95%	93%	<b>91%</b>
Q38. Patient received easily understandable information about what they should or should not do after leaving hospital	*	*	87%	88%	87%	90%	90%	87%	<b>89%</b>
Q39. Patient was always able to discuss worries and fears with hospital staff while being treated as an outpatient or day case	*	*	73%	71%	81%	82%	86%	79%	<b>81%</b>

## Age group tables

\* Indicates where a score has been suppressed because there are less than 11 responses. n.a. Indicates that there were no respondents for that age group.

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	*	*	83%	87%	88%	91%	93%	91%	<b>90%</b>
Q41_2. Beforehand patient completely had enough understandable information about chemotherapy	*	*	77%	79%	86%	85%	91%	85%	<b>86%</b>
Q41_3. Beforehand patient completely had enough understandable information about radiotherapy	n.a.	*	81%	86%	84%	87%	89%	*	<b>86%</b>
Q41_4. Beforehand patient completely had enough understandable information about hormone therapy	n.a.	*	65%	69%	84%	84%	83%	*	<b>80%</b>
Q41_5. Beforehand patient completely had enough understandable information about immunotherapy	*	*	77%	72%	80%	84%	90%	75%	<b>83%</b>
Q42_1. Patient completely had enough understandable information about progress with surgery	*	*	72%	80%	82%	89%	87%	88%	<b>85%</b>
Q42_2. Patient completely had enough understandable information about progress with chemotherapy	*	*	73%	73%	80%	80%	85%	74%	<b>80%</b>
Q42_3. Patient completely had enough understandable information about progress with radiotherapy	n.a.	*	76%	76%	78%	78%	80%	*	<b>78%</b>
Q42_4. Patient completely had enough understandable information about progress with hormone therapy	n.a.	*	68%	65%	71%	80%	84%	*	<b>75%</b>
Q42_5. Patient completely had enough understandable information about progress with immunotherapy	*	*	69%	77%	72%	82%	83%	75%	<b>79%</b>
Q43. Patient felt the length of waiting time at clinic and day unit for cancer treatment was about right	*	*	68%	74%	84%	84%	86%	84%	<b>83%</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	*	*	78%	74%	77%	77%	73%	78%	<b>76%</b>
Q45. Patient was always offered practical advice on dealing with any immediate side effects from treatment	*	*	65%	67%	73%	75%	73%	71%	<b>73%</b>
Q46. Patient was given information that they could access about support in dealing with immediate side effects from treatment	*	*	82%	82%	85%	89%	88%	71%	<b>86%</b>
Q47. Patient felt possible long-term side effects were definitely explained in a way they could understand in advance of their treatment	*	*	58%	58%	63%	65%	62%	60%	<b>63%</b>
Q48. Patient was definitely able to discuss options for managing the impact of any long-term side effects	*	*	53%	55%	56%	58%	62%	61%	<b>58%</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	*	*	41%	51%	55%	56%	65%	64%	<b>57%</b>
Q50. During treatment, the patient definitely got enough care and support at home from community or voluntary services	*	*	33%	59%	54%	54%	61%	64%	<b>56%</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	*	*	54%	51%	49%	44%	50%	48%	<b>48%</b>
Q52. Patient has had a review of cancer care by GP practice	*	*	23%	21%	22%	17%	21%	23%	<b>20%</b>



## Age group tables

\* Indicates where a score has been suppressed because there are less than 11 responses. n.a. Indicates that there were no respondents for that age group.

	Age								All
	16 - 24	25 - 34	35 - 44	45 - 54	55 - 64	65 - 74	75 - 84	85+	
Q53. After treatment, the patient definitely could get enough emotional support at home from community or voluntary services	*	*	22%	31%	33%	35%	44%	31%	<b>36%</b>
Q54. The right amount of information and support was offered to the patient between final treatment and the follow up appointment	*	*	65%	75%	77%	81%	84%	75%	<b>79%</b>
Q55. Patient was given enough information about the possibility and signs of cancer coming back or spreading	*	*	55%	57%	64%	69%	69%	76%	<b>66%</b>

	Age								All
	16 - 24	25 - 34	35 - 44	45 - 54	55 - 64	65 - 74	75 - 84	85+	
Q56. The whole care team worked well together	*	*	90%	89%	91%	91%	93%	95%	<b>92%</b>
Q57. Administration of care was very good or good	*	*	89%	89%	89%	90%	90%	92%	<b>90%</b>
Q58. Cancer research opportunities were discussed with patient	*	*	26%	34%	41%	43%	45%	32%	<b>41%</b>
Q59. Patient's average rating of care scored from very poor to very good	*	*	8.5	8.8	9.0	9.1	9.1	8.9	<b>9.0</b>

## Gender tables

\* Indicates where a score has been suppressed because there are less than 11 responses. n.a. Indicates that there were no respondents for that gender group.

	Gender						All
	Female	Male	Non-binary	Prefer to self-describe	Prefer not to say	Not given	
<b>SUPPORT FROM YOUR GP PRACTICE</b>							
Q2. Patient only spoke to primary care professional once or twice before cancer diagnosis	80%	76%	n.a.	n.a.	n.a.	82%	<b>78%</b>
Q3. Referral for diagnosis was explained in a way the patient could completely understand	69%	60%	n.a.	n.a.	*	*	<b>64%</b>

	Gender						All
	Female	Male	Non-binary	Prefer to self-describe	Prefer not to say	Not given	
<b>DIAGNOSTIC TESTS</b>							
Q5. Patient received all the information needed about the diagnostic test in advance	93%	92%	n.a.	n.a.	*	*	<b>93%</b>
Q6. Diagnostic test staff appeared to completely have all the information they needed about the patient	84%	86%	n.a.	n.a.	*	*	<b>85%</b>
Q7. Patient felt the length of time waiting for diagnostic test results was about right	81%	86%	n.a.	n.a.	*	*	<b>84%</b>
Q8. Diagnostic test results were explained in a way the patient could completely understand	81%	79%	n.a.	n.a.	*	*	<b>80%</b>
Q9. Enough privacy was always given to the patient when receiving diagnostic test results	94%	95%	n.a.	n.a.	*	*	<b>94%</b>

	Gender						All
	Female	Male	Non-binary	Prefer to self-describe	Prefer not to say	Not given	
<b>FINDING OUT THAT YOU HAD CANCER</b>							
Q12. Patient was told they could have a family member, carer or friend with them when told diagnosis	76%	77%	n.a.	n.a.	*	*	<b>77%</b>
Q13. Patient was definitely told sensitively that they had cancer	76%	76%	n.a.	n.a.	*	*	<b>76%</b>
Q14. Cancer diagnosis explained in a way the patient could completely understand	77%	76%	n.a.	n.a.	*	*	<b>76%</b>
Q15. Patient was definitely told about their diagnosis in an appropriate place	85%	87%	n.a.	n.a.	*	*	<b>86%</b>
Q16. Patient was told they could go back later for more information about their diagnosis	84%	84%	n.a.	n.a.	*	*	<b>84%</b>

	Gender						All
	Female	Male	Non-binary	Prefer to self-describe	Prefer not to say	Not given	
<b>SUPPORT FROM A MAIN CONTACT PERSON</b>							
Q17. Patient had a main point of contact within the care team	93%	91%	n.a.	n.a.	*	*	<b>92%</b>
Q18. Patient found it very or quite easy to contact their main contact person	87%	88%	n.a.	n.a.	*	*	<b>87%</b>
Q19. Patient found advice from main contact person was very or quite helpful	96%	96%	n.a.	n.a.	*	*	<b>96%</b>

## Gender tables

\* Indicates where a score has been suppressed because there are less than 11 responses. n.a. Indicates that there were no respondents for that gender group.

	Gender						All
	Female	Male	Non-binary	Prefer to self-describe	Prefer not to say	Not given	
Q20. Treatment options were explained in a way the patient could completely understand	84%	83%	n.a.	n.a.	*	*	<b>83%</b>
Q21. Patient was definitely involved as much as they wanted to be in decisions about their treatment	80%	82%	n.a.	n.a.	*	*	<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	75%	80%	n.a.	n.a.	*	*	<b>77%</b>
Q23. Patient could get further advice or a second opinion before making decisions about their treatment options	55%	57%	n.a.	n.a.	*	*	<b>57%</b>

	Gender						All
	Female	Male	Non-binary	Prefer to self-describe	Prefer not to say	Not given	
Q24. Patient was definitely able to have a discussion about their needs or concerns prior to treatment	72%	75%	n.a.	n.a.	*	*	<b>74%</b>
Q25. A member of their care team helped the patient create a care plan to address any needs or concerns	93%	95%	n.a.	n.a.	*	*	<b>94%</b>
Q26. Care team reviewed the patient's care plan with them to ensure it was up to date	99%	99%	n.a.	n.a.	*	*	<b>99%</b>

	Gender						All
	Female	Male	Non-binary	Prefer to self-describe	Prefer not to say	Not given	
Q27. Staff provided the patient with relevant information on available support	87%	91%	n.a.	n.a.	*	*	<b>89%</b>
Q28. Patient definitely got the right level of support for their overall health and well being from hospital staff	75%	82%	n.a.	n.a.	*	*	<b>78%</b>
Q29. Patient was offered information about how to get financial help or benefits	69%	67%	n.a.	n.a.	*	*	<b>68%</b>

## Gender tables

\* Indicates where a score has been suppressed because there are less than 11 responses. n.a. Indicates that there were no respondents for that gender group.

	Gender						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	81%	86%	n.a.	n.a.	*	*	<b>84%</b>
Q32. Patient's family, or someone close, was definitely able to talk to a member of the team looking after the patient in hospital	62%	67%	n.a.	n.a.	*	*	<b>65%</b>
Q33. Patient was always involved in decisions about their care and treatment whilst in hospital	70%	75%	n.a.	n.a.	*	*	<b>73%</b>
Q34. Patient was always able to get help from ward staff when needed	74%	82%	n.a.	n.a.	*	*	<b>79%</b>
Q35. Patient was always able to discuss worries and fears with hospital staff	65%	73%	n.a.	n.a.	*	*	<b>69%</b>
Q36. Hospital staff always did everything they could to help the patient control pain	83%	88%	n.a.	n.a.	*	*	<b>85%</b>
Q37. Patient was always treated with respect and dignity while in hospital	88%	93%	n.a.	n.a.	*	*	<b>91%</b>
Q38. Patient received easily understandable information about what they should or should not do after leaving hospital	88%	89%	n.a.	n.a.	*	*	<b>89%</b>
Q39. Patient was always able to discuss worries and fears with hospital staff while being treated as an outpatient or day case	79%	85%	n.a.	n.a.	*	*	<b>81%</b>

	Gender						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%	n.a.	n.a.	*	*	<b>90%</b>
Q41_2. Beforehand patient completely had enough understandable information about chemotherapy	85%	87%	n.a.	n.a.	n.a.	87%	<b>86%</b>
Q41_3. Beforehand patient completely had enough understandable information about radiotherapy	87%	85%	n.a.	n.a.	n.a.	95%	<b>86%</b>
Q41_4. Beforehand patient completely had enough understandable information about hormone therapy	79%	82%	n.a.	n.a.	n.a.	89%	<b>80%</b>
Q41_5. Beforehand patient completely had enough understandable information about immunotherapy	83%	82%	n.a.	n.a.	n.a.	89%	<b>83%</b>
Q42_1. Patient completely had enough understandable information about progress with surgery	85%	84%	n.a.	n.a.	*	*	<b>85%</b>
Q42_2. Patient completely had enough understandable information about progress with chemotherapy	80%	80%	n.a.	n.a.	*	*	<b>80%</b>
Q42_3. Patient completely had enough understandable information about progress with radiotherapy	81%	72%	n.a.	n.a.	*	*	<b>78%</b>
Q42_4. Patient completely had enough understandable information about progress with hormone therapy	73%	79%	n.a.	n.a.	n.a.	78%	<b>75%</b>
Q42_5. Patient completely had enough understandable information about progress with immunotherapy	78%	79%	n.a.	n.a.	n.a.	94%	<b>79%</b>
Q43. Patient felt the length of waiting time at clinic and day unit for cancer treatment was about right	80%	87%	n.a.	n.a.	*	*	<b>83%</b>

## Gender tables

\* Indicates where a score has been suppressed because there are less than 11 responses. n.a. Indicates that there were no respondents for that gender group.

	Gender						All
	Female	Male	Non-binary	Prefer to self-describe	Prefer not to say	Not given	
<b>IMMEDIATE AND LONG TERM SIDE EFFECTS</b>							
Q44. Possible side effects from treatment were definitely explained in a way the patient could understand	75%	78%	n.a.	n.a.	*	*	<b>76%</b>
Q45. Patient was always offered practical advice on dealing with any immediate side effects from treatment	72%	74%	n.a.	n.a.	*	*	<b>73%</b>
Q46. Patient was given information that they could access about support in dealing with immediate side effects from treatment	85%	87%	n.a.	n.a.	*	*	<b>86%</b>
Q47. Patient felt possible long-term side effects were definitely explained in a way they could understand in advance of their treatment	61%	65%	n.a.	n.a.	*	*	<b>63%</b>
Q48. Patient was definitely able to discuss options for managing the impact of any long-term side effects	54%	62%	n.a.	n.a.	*	*	<b>58%</b>

	Gender						All
	Female	Male	Non-binary	Prefer to self-describe	Prefer not to say	Not given	
<b>SUPPORT WHILE AT HOME</b>							
Q49. Care team gave family, or someone close, all the information needed to help care for the patient at home	53%	62%	n.a.	n.a.	*	*	<b>57%</b>
Q50. During treatment, the patient definitely got enough care and support at home from community or voluntary services	52%	59%	n.a.	n.a.	*	*	<b>56%</b>

	Gender						All
	Female	Male	Non-binary	Prefer to self-describe	Prefer not to say	Not given	
<b>CARE FROM YOUR GP PRACTICE</b>							
Q51. Patient definitely received the right amount of support from their GP practice during treatment	47%	49%	n.a.	n.a.	*	*	<b>48%</b>
Q52. Patient has had a review of cancer care by GP practice	19%	20%	n.a.	n.a.	*	*	<b>20%</b>

	Gender						All
	Female	Male	Non-binary	Prefer to self-describe	Prefer not to say	Not given	
<b>LIVING WITH AND BEYOND CANCER</b>							
Q53. After treatment, the patient definitely could get enough emotional support at home from community or voluntary services	32%	40%	n.a.	n.a.	*	*	<b>36%</b>
Q54. The right amount of information and support was offered to the patient between final treatment and the follow up appointment	77%	82%	n.a.	n.a.	*	*	<b>79%</b>
Q55. Patient was given enough information about the possibility and signs of cancer coming back or spreading	61%	71%	n.a.	n.a.	*	*	<b>66%</b>

## Gender tables

\* Indicates where a score has been suppressed because there are less than 11 responses. n.a. Indicates that there were no respondents for that gender group.

YOUR OVERALL NHS CARE	Gender						All
	Female	Male	Non-binary	Prefer to self-describe	Prefer not to say	Not given	
Q56. The whole care team worked well together	91%	92%	n.a.	n.a.	*	*	<b>92%</b>
Q57. Administration of care was very good or good	90%	90%	n.a.	n.a.	*	*	<b>90%</b>
Q58. Cancer research opportunities were discussed with patient	35%	47%	n.a.	n.a.	*	*	<b>41%</b>
Q59. Patient's average rating of care scored from very poor to very good	8.9	9.1	n.a.	n.a.	*	*	<b>9.0</b>

## Ethnicity tables

\* Indicates where a score has been suppressed because there are less than 11 responses. n.a. Indicates that there were no respondents for that ethnicity group.

	Ethnic group						
	White	Mixed	Asian	Black	Other	Not given	All
<b>SUPPORT FROM YOUR GP PRACTICE</b>							
Q2. Patient only spoke to primary care professional once or twice before cancer diagnosis	78%	55%	93%	*	*	82%	<b>78%</b>
Q3. Referral for diagnosis was explained in a way the patient could completely understand	65%	57%	71%	*	*	55%	<b>64%</b>

	Ethnic group						
	White	Mixed	Asian	Black	Other	Not given	All
<b>DIAGNOSTIC TESTS</b>							
Q5. Patient received all the information needed about the diagnostic test in advance	93%	94%	86%	*	*	94%	<b>93%</b>
Q6. Diagnostic test staff appeared to completely have all the information they needed about the patient	85%	81%	56%	*	*	85%	<b>85%</b>
Q7. Patient felt the length of time waiting for diagnostic test results was about right	83%	69%	83%	*	*	90%	<b>84%</b>
Q8. Diagnostic test results were explained in a way the patient could completely understand	80%	75%	60%	*	*	81%	<b>80%</b>
Q9. Enough privacy was always given to the patient when receiving diagnostic test results	94%	87%	84%	*	*	96%	<b>94%</b>

	Ethnic group						
	White	Mixed	Asian	Black	Other	Not given	All
<b>FINDING OUT THAT YOU HAD CANCER</b>							
Q12. Patient was told they could have a family member, carer or friend with them when told diagnosis	77%	76%	72%	*	*	77%	<b>77%</b>
Q13. Patient was definitely told sensitively that they had cancer	76%	74%	74%	*	*	75%	<b>76%</b>
Q14. Cancer diagnosis explained in a way the patient could completely understand	76%	74%	70%	*	*	76%	<b>76%</b>
Q15. Patient was definitely told about their diagnosis in an appropriate place	86%	78%	81%	*	*	87%	<b>86%</b>
Q16. Patient was told they could go back later for more information about their diagnosis	84%	76%	88%	*	*	83%	<b>84%</b>

	Ethnic group						
	White	Mixed	Asian	Black	Other	Not given	All
<b>SUPPORT FROM A MAIN CONTACT PERSON</b>							
Q17. Patient had a main point of contact within the care team	92%	84%	93%	*	*	93%	<b>92%</b>
Q18. Patient found it very or quite easy to contact their main contact person	88%	81%	76%	*	*	88%	<b>87%</b>
Q19. Patient found advice from main contact person was very or quite helpful	96%	94%	88%	*	*	97%	<b>96%</b>

	Ethnic group						
	White	Mixed	Asian	Black	Other	Not given	All
<b>DECIDING ON THE BEST TREATMENT</b>							
Q20. Treatment options were explained in a way the patient could completely understand	84%	83%	81%	*	*	79%	<b>83%</b>
Q21. Patient was definitely involved as much as they wanted to be in decisions about their treatment	81%	68%	77%	*	*	78%	<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	77%	71%	88%	*	*	78%	<b>77%</b>
Q23. Patient could get further advice or a second opinion before making decisions about their treatment options	56%	58%	53%	*	*	67%	<b>57%</b>

## Ethnicity tables

\* Indicates where a score has been suppressed because there are less than 11 responses. n.a. Indicates that there were no respondents for that ethnicity group.

<b>CARE PLANNING</b>	Ethnic group						
	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%	65%	64%	*	*	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%	83%	100%	*	*	95%	<b>94%</b>
Q26. Care team reviewed the patient's care plan with them to ensure it was up to date	99%	*	93%	*	*	98%	<b>99%</b>

<b>SUPPORT FROM HOSPITAL STAFF</b>	Ethnic group						
	White	Mixed	Asian	Black	Other	Not given	All
Q27. Staff provided the patient with relevant information on available support	89%	83%	87%	*	*	88%	<b>89%</b>
Q28. Patient definitely got the right level of support for their overall health and well being from hospital staff	79%	58%	63%	*	*	78%	<b>78%</b>
Q29. Patient was offered information about how to get financial help or benefits	68%	56%	59%	*	*	74%	<b>68%</b>

<b>HOSPITAL CARE</b>	Ethnic group						
	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	84%	*	50%	*	*	88%	<b>84%</b>
Q32. Patient's family, or someone close, was definitely able to talk to a member of the team looking after the patient in hospital	65%	*	55%	*	*	75%	<b>65%</b>
Q33. Patient was always involved in decisions about their care and treatment whilst in hospital	72%	*	75%	*	*	77%	<b>73%</b>
Q34. Patient was always able to get help from ward staff when needed	79%	*	58%	*	*	82%	<b>79%</b>
Q35. Patient was always able to discuss worries and fears with hospital staff	69%	*	73%	*	*	75%	<b>69%</b>
Q36. Hospital staff always did everything they could to help the patient control pain	85%	*	83%	*	*	90%	<b>85%</b>
Q37. Patient was always treated with respect and dignity while in hospital	91%	*	92%	*	*	92%	<b>91%</b>
Q38. Patient received easily understandable information about what they should or should not do after leaving hospital	89%	*	92%	*	*	93%	<b>89%</b>
Q39. Patient was always able to discuss worries and fears with hospital staff while being treated as an outpatient or day case	82%	68%	57%	*	*	84%	<b>81%</b>



## Ethnicity tables

\* Indicates where a score has been suppressed because there are less than 11 responses. n.a. Indicates that there were no respondents for that ethnicity group.

YOUR TREATMENT	Ethnic group						
	White	Mixed	Asian	Black	Other	Not given	All
Q41_1. Beforehand patient completely had enough understandable information about surgery	90%	92%	90%	*	*	88%	<b>90%</b>
Q41_2. Beforehand patient completely had enough understandable information about chemotherapy	86%	83%	93%	*	*	87%	<b>86%</b>
Q41_3. Beforehand patient completely had enough understandable information about radiotherapy	86%	*	77%	*	*	92%	<b>86%</b>
Q41_4. Beforehand patient completely had enough understandable information about hormone therapy	81%	*	*	*	*	76%	<b>80%</b>
Q41_5. Beforehand patient completely had enough understandable information about immunotherapy	83%	*	n.a.	*	n.a.	80%	<b>83%</b>
Q42_1. Patient completely had enough understandable information about progress with surgery	85%	83%	70%	*	*	88%	<b>85%</b>
Q42_2. Patient completely had enough understandable information about progress with chemotherapy	80%	83%	79%	*	*	85%	<b>80%</b>
Q42_3. Patient completely had enough understandable information about progress with radiotherapy	79%	*	69%	*	*	74%	<b>78%</b>
Q42_4. Patient completely had enough understandable information about progress with hormone therapy	75%	*	*	*	*	74%	<b>75%</b>
Q42_5. Patient completely had enough understandable information about progress with immunotherapy	79%	*	n.a.	*	n.a.	83%	<b>79%</b>
Q43. Patient felt the length of waiting time at clinic and day unit for cancer treatment was about right	83%	61%	65%	*	*	86%	<b>83%</b>

IMMEDIATE AND LONG TERM SIDE EFFECTS	Ethnic group						
	White	Mixed	Asian	Black	Other	Not given	All
Q44. Possible side effects from treatment were definitely explained in a way the patient could understand	76%	72%	74%	*	*	76%	<b>76%</b>
Q45. Patient was always offered practical advice on dealing with any immediate side effects from treatment	73%	68%	52%	*	*	73%	<b>73%</b>
Q46. Patient was given information that they could access about support in dealing with immediate side effects from treatment	86%	71%	74%	*	*	91%	<b>86%</b>
Q47. Patient felt possible long-term side effects were definitely explained in a way they could understand in advance of their treatment	63%	61%	70%	*	*	63%	<b>63%</b>
Q48. Patient was definitely able to discuss options for managing the impact of any long-term side effects	58%	59%	48%	*	*	61%	<b>58%</b>

SUPPORT WHILE AT HOME	Ethnic group						
	White	Mixed	Asian	Black	Other	Not given	All
Q49. Care team gave family, or someone close, all the information needed to help care for the patient at home	57%	54%	50%	*	*	61%	<b>57%</b>
Q50. During treatment, the patient definitely got enough care and support at home from community or voluntary services	56%	57%	63%	*	*	54%	<b>56%</b>

CARE FROM YOUR GP PRACTICE	Ethnic group						
	White	Mixed	Asian	Black	Other	Not given	All
Q51. Patient definitely received the right amount of support from their GP practice during treatment	48%	53%	41%	*	*	47%	<b>48%</b>
Q52. Patient has had a review of cancer care by GP practice	20%	26%	30%	*	*	23%	<b>20%</b>

## Ethnicity tables

\* Indicates where a score has been suppressed because there are less than 11 responses. n.a. Indicates that there were no respondents for that ethnicity group.

	Ethnic group						
	White	Mixed	Asian	Black	Other	Not given	All
<b>LIVING WITH AND BEYOND CANCER</b>							
Q53. After treatment, the patient definitely could get enough emotional support at home from community or voluntary services	36%	*	38%	*	*	37%	<b>36%</b>
Q54. The right amount of information and support was offered to the patient between final treatment and the follow up appointment	79%	*	76%	*	*	78%	<b>79%</b>
Q55. Patient was given enough information about the possibility and signs of cancer coming back or spreading	66%	69%	60%	*	*	71%	<b>66%</b>

	Ethnic group						
	White	Mixed	Asian	Black	Other	Not given	All
<b>YOUR OVERALL NHS CARE</b>							
Q56. The whole care team worked well together	91%	79%	96%	*	*	94%	<b>92%</b>
Q57. Administration of care was very good or good	90%	74%	100%	*	*	92%	<b>90%</b>
Q58. Cancer research opportunities were discussed with patient	40%	67%	54%	*	*	48%	<b>41%</b>
Q59. Patient's average rating of care scored from very poor to very good	9.0	7.9	8.5	*	*	9.0	<b>9.0</b>

## IMD quintile tables

\* Indicates where a score has been suppressed because there are less than 11 responses. n.a. Indicates that there were no respondents for that quintile group.

	IMD Quintile						
	1 (most deprived)	2	3	4	5 (least deprived)	Non-England	All
<b>SUPPORT FROM YOUR GP PRACTICE</b>							
Q2. Patient only spoke to primary care professional once or twice before cancer diagnosis	74%	75%	81%	83%	82%	n.a.	<b>78%</b>
Q3. Referral for diagnosis was explained in a way the patient could completely understand	58%	64%	68%	67%	66%	n.a.	<b>64%</b>

	IMD Quintile						
	1 (most deprived)	2	3	4	5 (least deprived)	Non-England	All
<b>DIAGNOSTIC TESTS</b>							
Q5. Patient received all the information needed about the diagnostic test in advance	93%	93%	92%	94%	93%	n.a.	<b>93%</b>
Q6. Diagnostic test staff appeared to completely have all the information they needed about the patient	83%	86%	88%	82%	87%	n.a.	<b>85%</b>
Q7. Patient felt the length of time waiting for diagnostic test results was about right	84%	86%	83%	82%	82%	n.a.	<b>84%</b>
Q8. Diagnostic test results were explained in a way the patient could completely understand	78%	81%	81%	82%	79%	n.a.	<b>80%</b>
Q9. Enough privacy was always given to the patient when receiving diagnostic test results	94%	94%	95%	94%	96%	n.a.	<b>94%</b>

	IMD Quintile						
	1 (most deprived)	2	3	4	5 (least deprived)	Non-England	All
<b>FINDING OUT THAT YOU HAD CANCER</b>							
Q12. Patient was told they could have a family member, carer or friend with them when told diagnosis	78%	78%	76%	75%	76%	n.a.	<b>77%</b>
Q13. Patient was definitely told sensitively that they had cancer	74%	78%	74%	76%	76%	n.a.	<b>76%</b>
Q14. Cancer diagnosis explained in a way the patient could completely understand	76%	77%	75%	78%	76%	n.a.	<b>76%</b>
Q15. Patient was definitely told about their diagnosis in an appropriate place	86%	86%	84%	87%	85%	n.a.	<b>86%</b>
Q16. Patient was told they could go back later for more information about their diagnosis	84%	84%	83%	85%	86%	n.a.	<b>84%</b>

	IMD Quintile						
	1 (most deprived)	2	3	4	5 (least deprived)	Non-England	All
<b>SUPPORT FROM A MAIN CONTACT PERSON</b>							
Q17. Patient had a main point of contact within the care team	93%	94%	91%	92%	91%	n.a.	<b>92%</b>
Q18. Patient found it very or quite easy to contact their main contact person	88%	90%	88%	86%	86%	n.a.	<b>87%</b>
Q19. Patient found advice from main contact person was very or quite helpful	95%	97%	96%	97%	95%	n.a.	<b>96%</b>

## IMD quintile tables

\* Indicates where a score has been suppressed because there are less than 11 responses. n.a. Indicates that there were no respondents for that quintile group.

	IMD Quintile						
	1 (most deprived)	2	3	4	5 (least deprived)	Non-England	All
Q20. Treatment options were explained in a way the patient could completely understand	82%	85%	83%	84%	83%	n.a.	<b>83%</b>
Q21. Patient was definitely involved as much as they wanted to be in decisions about their treatment	81%	83%	79%	82%	80%	n.a.	<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	78%	79%	75%	77%	78%	n.a.	<b>77%</b>
Q23. Patient could get further advice or a second opinion before making decisions about their treatment options	62%	59%	54%	56%	50%	n.a.	<b>57%</b>

	IMD Quintile						
	1 (most deprived)	2	3	4	5 (least deprived)	Non-England	All
Q24. Patient was definitely able to have a discussion about their needs or concerns prior to treatment	73%	76%	72%	74%	74%	n.a.	<b>74%</b>
Q25. A member of their care team helped the patient create a care plan to address any needs or concerns	93%	94%	92%	94%	94%	n.a.	<b>94%</b>
Q26. Care team reviewed the patient's care plan with them to ensure it was up to date	99%	99%	99%	99%	99%	n.a.	<b>99%</b>

	IMD Quintile						
	1 (most deprived)	2	3	4	5 (least deprived)	Non-England	All
Q27. Staff provided the patient with relevant information on available support	88%	88%	88%	89%	91%	n.a.	<b>89%</b>
Q28. Patient definitely got the right level of support for their overall health and well being from hospital staff	79%	80%	78%	78%	77%	n.a.	<b>78%</b>
Q29. Patient was offered information about how to get financial help or benefits	67%	66%	69%	72%	69%	n.a.	<b>68%</b>

	IMD Quintile						
	1 (most deprived)	2	3	4	5 (least deprived)	Non-England	All
Q31. Patient had confidence and trust in all of the team looking after them during their stay in hospital	84%	83%	83%	87%	83%	n.a.	<b>84%</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%	64%	65%	67%	63%	n.a.	<b>65%</b>
Q33. Patient was always involved in decisions about their care and treatment whilst in hospital	71%	73%	72%	72%	75%	n.a.	<b>73%</b>
Q34. Patient was always able to get help from ward staff when needed	79%	81%	77%	77%	78%	n.a.	<b>79%</b>
Q35. Patient was always able to discuss worries and fears with hospital staff	72%	70%	66%	69%	68%	n.a.	<b>69%</b>
Q36. Hospital staff always did everything they could to help the patient control pain	85%	84%	84%	87%	87%	n.a.	<b>85%</b>
Q37. Patient was always treated with respect and dignity while in hospital	92%	91%	89%	90%	91%	n.a.	<b>91%</b>
Q38. Patient received easily understandable information about what they should or should not do after leaving hospital	89%	91%	84%	90%	90%	n.a.	<b>89%</b>
Q39. Patient was always able to discuss worries and fears with hospital staff while being treated as an outpatient or day case	82%	83%	80%	81%	81%	n.a.	<b>81%</b>

## IMD quintile tables

\* Indicates where a score has been suppressed because there are less than 11 responses.

n.a. Indicates that there were no respondents for that quintile group.

YOUR TREATMENT	IMD Quintile						
	1 (most deprived)	2	3	4	5 (least deprived)	Non-England	All
Q41_1. Beforehand patient completely had enough understandable information about surgery	89%	92%	87%	90%	90%	n.a.	<b>90%</b>
Q41_2. Beforehand patient completely had enough understandable information about chemotherapy	88%	85%	84%	88%	83%	n.a.	<b>86%</b>
Q41_3. Beforehand patient completely had enough understandable information about radiotherapy	87%	87%	83%	90%	84%	n.a.	<b>86%</b>
Q41_4. Beforehand patient completely had enough understandable information about hormone therapy	81%	87%	78%	78%	77%	n.a.	<b>80%</b>
Q41_5. Beforehand patient completely had enough understandable information about immunotherapy	85%	80%	85%	80%	83%	n.a.	<b>83%</b>
Q42_1. Patient completely had enough understandable information about progress with surgery	85%	87%	83%	84%	85%	n.a.	<b>85%</b>
Q42_2. Patient completely had enough understandable information about progress with chemotherapy	81%	83%	79%	78%	78%	n.a.	<b>80%</b>
Q42_3. Patient completely had enough understandable information about progress with radiotherapy	78%	81%	75%	80%	76%	n.a.	<b>78%</b>
Q42_4. Patient completely had enough understandable information about progress with hormone therapy	74%	84%	71%	73%	71%	n.a.	<b>75%</b>
Q42_5. Patient completely had enough understandable information about progress with immunotherapy	81%	80%	76%	78%	82%	n.a.	<b>79%</b>
Q43. Patient felt the length of waiting time at clinic and day unit for cancer treatment was about right	82%	86%	82%	85%	80%	n.a.	<b>83%</b>

IMMEDIATE AND LONG TERM SIDE EFFECTS	IMD Quintile						
	1 (most deprived)	2	3	4	5 (least deprived)	Non-England	All
Q44. Possible side effects from treatment were definitely explained in a way the patient could understand	78%	77%	77%	73%	77%	n.a.	<b>76%</b>
Q45. Patient was always offered practical advice on dealing with any immediate side effects from treatment	75%	74%	71%	71%	73%	n.a.	<b>73%</b>
Q46. Patient was given information that they could access about support in dealing with immediate side effects from treatment	86%	88%	86%	85%	86%	n.a.	<b>86%</b>
Q47. Patient felt possible long-term side effects were definitely explained in a way they could understand in advance of their treatment	64%	64%	62%	62%	62%	n.a.	<b>63%</b>
Q48. Patient was definitely able to discuss options for managing the impact of any long-term side effects	59%	60%	58%	54%	58%	n.a.	<b>58%</b>

SUPPORT WHILE AT HOME	IMD Quintile						
	1 (most deprived)	2	3	4	5 (least deprived)	Non-England	All
Q49. Care team gave family, or someone close, all the information needed to help care for the patient at home	58%	62%	56%	57%	51%	n.a.	<b>57%</b>
Q50. During treatment, the patient definitely got enough care and support at home from community or voluntary services	59%	53%	52%	60%	54%	n.a.	<b>56%</b>

CARE FROM YOUR GP PRACTICE	IMD Quintile						
	1 (most deprived)	2	3	4	5 (least deprived)	Non-England	All
Q51. Patient definitely received the right amount of support from their GP practice during treatment	44%	46%	51%	48%	51%	n.a.	<b>48%</b>
Q52. Patient has had a review of cancer care by GP practice	22%	19%	23%	18%	18%	n.a.	<b>20%</b>

## IMD quintile tables

\* Indicates where a score has been suppressed because there are less than 11 responses. n.a. Indicates that there were no respondents for that quintile group.

	IMD Quintile						
	1 (most deprived)	2	3	4	5 (least deprived)	Non-England	All
Q53. After treatment, the patient definitely could get enough emotional support at home from community or voluntary services	41%	30%	34%	34%	38%	n.a.	<b>36%</b>
Q54. The right amount of information and support was offered to the patient between final treatment and the follow up appointment	78%	80%	80%	79%	79%	n.a.	<b>79%</b>
Q55. Patient was given enough information about the possibility and signs of cancer coming back or spreading	65%	68%	64%	67%	66%	n.a.	<b>66%</b>

	IMD Quintile						
	1 (most deprived)	2	3	4	5 (least deprived)	Non-England	All
Q56. The whole care team worked well together	92%	92%	91%	91%	91%	n.a.	<b>92%</b>
Q57. Administration of care was very good or good	90%	91%	90%	90%	89%	n.a.	<b>90%</b>
Q58. Cancer research opportunities were discussed with patient	46%	38%	38%	38%	45%	n.a.	<b>41%</b>
Q59. Patient's average rating of care scored from very poor to very good	9.0	9.1	8.9	9.0	9.0	n.a.	<b>9.0</b>