Applying National Cancer Control Indicators to Regional and Remote Cancer Patients' Experiences Jeff Dunn, Belinda C Goodwin (belindagoodwin@cancerqld.org.au), Joanne F. Aitken, Sonja March, Fiona Crawford-Williams, Michael J. Ireland, Nicholas Ralph, Leah Zajdlewicz, Arlen Rowe, and Suzanne K. Chambers.

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Supplementary Table 1. Frequencies and percentages of responses to NCCI – Patient Experience Items and grouping of binary responses, n (%).

	Yes, I completely understood it	Yes, I understood some of it	No	I can't remember		Missing
When you were told you had cancer, did you understand the explanation of what was wrong with you?	345 (66.7%)	153 (29.4%)	14 (2.7%)	5 (1.0%)		1 (0.19%)
	Yes, and it was easy to understand	Yes, and it was difficult to understand	No	I did not need written information	I don't know/ can't remember	Missing
When you were told you had cancer, were you given written information about the type of cancer you had?	262 (51.3%)	54 (10.6%)	134 (26.2%)	44 (8.6%)	17 (3/3%)	7 (1.4%)
	Yes	No I	I don't know/ can't remember			Missing
Have you been offered a written assessment and care plan	159 (31.0%)	265 (51.7%)	89 (17.3%)			5 (1.0%)
	Yes	No	I don't know/ can't remember			Missing
Were you given the name of a Clinical Nurse Specialist who would support you through your treatment?	317 (61.8%)	143 (27.9%)	53 (10.2%)			5 (1.0%)
	Yes, definitely	Yes, to some extent	No, but I would have liked to be more involved	I don't know/can't remember		Missing
Were you involved as much as you wanted to be in decisions about your care and treatment?	305 (59.2%)	161 (31.3%)	36 (6.9%)	13 (2.5%)		3 (0.6%)

[^]valid percentage reported except for missing column, calculated based on non-missing data ==== = coded as yes, ==== = coded as no, no border = treated as missing

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Supplementary Table 1. (Cont.)

	Yes, definitely	Yes, to some extent	No	I didn't know my treatment was being discussed	I don't know/ can't remember	Missing
Do you think your views were taken into account when the team of doctors and nurses caring for you were discussing which treatment you should have?	294 (57.8%)	152 (29.2%)	31 (6.1%)	17 (2.9%)	15 (2.8%)	9 (1.7%)
	Yes, definitely	Yes, to some extent	No	I did not need an explanation	I don't know/ can't remember	Missing
Were the possible side effects of treatment(s) explained in a way you could understand?	338 (65.5%)	136 (26.4%)	30 (5.8%)	2 (0.4%)	10 (1.9%)	2 (0.4%)
	Yes, and it was easy to understand	Yes, and it was difficult to understand	No	I don't know/ can't remember		Missing
Before you started your treatment, were you given written information about the side effects of treatment(s)?	354 (69.4%)	54 (10.6%)	73 (14.3%)	29 (5.7%)		8 (1.5%)

[^]valid percentage reported except for missing column, calculated based on non-missing data ---- = coded as yes, ---- = coded as no, no border = treated as missing

Note: To assess whether time since diagnosis was associated with non-response (i.e., no response or I don't know/ don't remember response), point bi-serial correlations were conducted between non-response and days since diagnosis. One weak association whereby those who were further past diagnosis were less likely to recall/provide a response to the NCCI item "Were you involved as much as you wanted to be in decisions about your care and treatment" (r = .130, p = .004)