Table 3 **Participants' agreement with the type of information they should receive prior to enrolment in research (*n* = 202)**

Item	Response[a]	
	No.	No.
Participants should know the risks and side-effects of the research	197	98.5
Participants should be given an explanation of anticipated benefits from the research	191	95.5
Participants should be given an explanation of the procedures and any drugs that will be used	181	90.0
Participants should be told that enrolment in research is voluntary	179	89.5
Participants should know the purpose of the research	177	88.1
Patients should be provided with contact information if questions, concerns, or complaints about the research were to occur	176	88.0
Participants should know the alternatives of medical treatment they can receive outside of the study	157	78.9

[a]*Percentage of those who strongly agreed or agreed.*