Appendix 2. The Desirability of Categories of Genetic Information

Category	Preference - Pero				
	Definitely Not	Probably Not	Probably	Definitely	No Response
Non-Medical	26.6% (147)	32.2% (178)	25.5% (141)	14.5% (80)	1.3% (7)
Common					
Untreatable	13.7% (76)	19.9% (110)	30.2% (167)	35.3% (195)	0.9% (5)
Fatal Adult-					
Onset	11.2% (62)	18.4% (102)	28.6% (158)	40.3% (223)	1.4% (8)
Serious					
Treatable					
Adult-Onset	8.3% (46)	14.1% (78)	32.4% (179)	43.9% (243)	1.3% (7)
Common					
Treatable	6.7% (37)	11% (61)	32.2% (178)	49% (271)	1.1% (6)
Devel. Delay,					
Learning					
Disability	5.4% (30)	10.5% (58)	30% (166)	53% (293)	1.1% (6)
Fatal					
Congenital	9.4% (52)	6.1% (34)	25.1% (139)	58.2% (322)	1.1% (6)
Serious					
Treatable					
Child-Onset	3.8% (21)	5.6% (31)	26.2% (145)	63.5% (351)	0.9% (5)

Appendix 3. P-Values Resulting From Pairwise Comparisons Between the 8 Different Categories of Genetic Information

Non-Medical Traits	Common Untreatable Conditions <.0001	Fatal Adult Onset Conditions <.0001	Serious Treatable Adult-Onset Conditions	Common Treatable Conditions <.0001	Developmental Delay/Learning Disability <.0001	Fatal Congenital Conditions <.0001	Serious Treatable Childhood-Onset Conditions <.0001
Common Untreatable		.0543	<.0001	<.0001	<.0001	<.0001	<.0001
Conditions Fatal Adult-Onset Conditions			.0001	<.0001	<.0001	<.0001	<.0001
Serious Treatable Adult-Onset Conditions				.0078	<.0001	.0002	<.0001
Common Treatable Conditions					.2575	.2482	<.0001
Developmental Delay/Learning Disability						.8348	<.0001
Fatal Congenital Conditions							.0001

Note: Bolding represents non-significant p-values (p>0.05). Blank columns are duplicate values.

Appendix 4. Reasons Respondents Wanted or Did Not Want Categories of Genetic Information

				Percent	(frequency) $n = 553$					
	Reasons respondents wanted information				Reasons res	Reasons respondents did not want information				
	Prepare Financially Medically etc	Decisions about Future Children	Consider Ending Pregnancy	Just to Know	Would Not End Pregnancy	Increase Stress	Protect Child's Privacy	Retain Mystery	Not Important	No Response
Non- Medical	11.9% (66)	4.9% (27)	1.6% (9)	21.2% (117)	15% (83)	1.8% (10)	5.8% (32)	9.4% (52)	26.2% (145)	2.2% (12)
Common Untreatable	43.6% (241)	4.9% (27)	5.1% (28)	11.4% (63)	7.6% (42)	13.2% (73)	8% (44)	0.5% (3)	4% (22)	1.8% (10)
Fatal Adult- Onset	41.6% (230)	8.1% (45)	9.4% (52)	9% (50)	8% (44)	12.8% (71)	6% (33)	1.3% (7)	1.6% (9)	2.2% (12)
Serious Treatable Adult- Onset	52.8% (292)	8.5% (47)	3.8% (21)	10.1% (56)	4.3% (24)	8% (44)	7.8% (43)	0% (0)	1.8% (10)	2.9% (16)
Common Treatable	65.1% (360)	7.6% (42)	2.4% (13)	5.8% (32)	6.7% (37)	6.5% (36)	3.1% (17)	0.2% (1)	0.9% (5)	1.8% (10)
Devel. Delay Learning Disability	67.5% (373)	5.1% (28)	6.3% (35)	3.1% (17)	6% (33)	5.6% (31)	1.6% (9)	1.3% (7)	1.4% (8)	2.2% (12)

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Fatal	35.4% (196)	9% (50)	30.9%	7.2% (40)	5.2% (29)	5.8% (32)	2.2% (12)	0.4% (2)	1.8% (10)	2% (11)
Congenital			(171)							
Serious	71.1% (393)	7.2% (40)	7.1% (39)	3.6% (20)	3.3% (18)	4.3% (24)	0.7% (4)	0.5% (3)	0.4% (2)	1.8% (10)
Treatable										
Child-										
Onset										

Appendix 5. Participant Agreement With Different Statements About Prenatal Whole Genome Sequencing

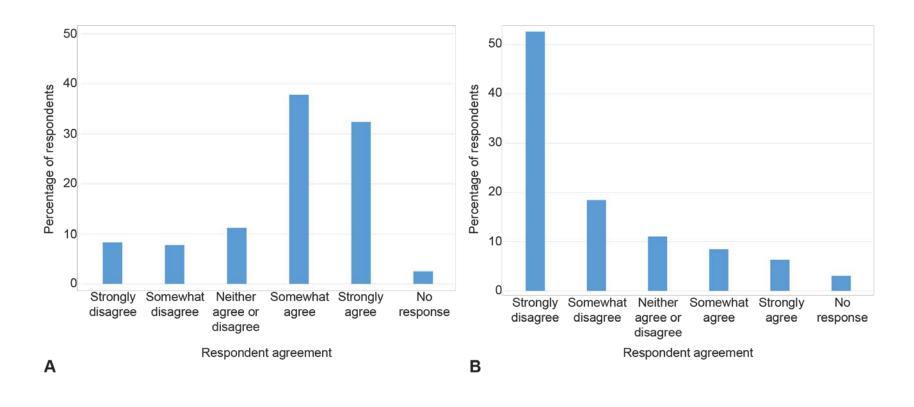
Statement	Response	Percent (Frequency) n = 553
	Strongly Disagree	6.7% (37)
	Somewhat Disagree	9.6% (53)
The more information parents have about their	Neither Agree or Disagree	9.6% (53)
baby's genes, the better.	Somewhat Agree	31.7% (175)
	Strongly Agree	40.3% (223)
	No Response	2.2% (12)
	Strongly Disagree	5.2% (29)
	Somewhat Disagree	2.7% (15)
Parents should be able to access all medically	Neither Agree or Disagree	4% (22)
relevant genetic information that they want to know.	Somewhat Agree	22.6% (125)
KHOW.	Strongly Agree	64.4% (356)
	No Response	1.1% (6)
	Strongly Disagree	17% (94)
Parents should be able to access all non-medical	Somewhat Disagree	16.8% (93)
genetic information that they want to know. Non-	Neither Agree or Disagree	18.1% (100)
medical information might include eye color or	Somewhat Agree	20.1% (111)
height.	Strongly Agree	21.9% (121)
	No Response	6.2% (34)
	Strongly Disagree	8.3% (46)
	Somewhat Disagree	7.8% (43)
It is appropriate for physicians to provide their	Neither Agree or Disagree	11.2% (62)
opinion about the kinds of genetic information that parents should learn about their babies.	Somewhat Agree	37.8% (209)
parents should learn about their bables.	Strongly Agree	32.4% (179)
	No Response	2.5% (14)

The authors provided this information as a supplement to their article.

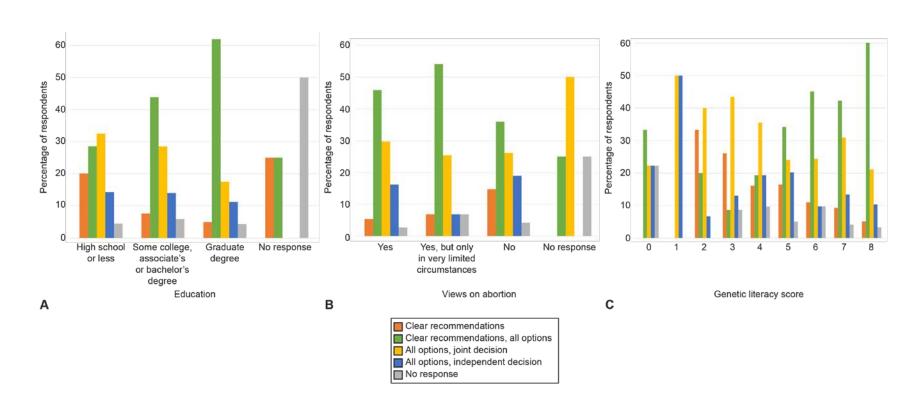
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	Strongly Disagree	52.6% (291)
	Somewhat Disagree	18.4% (102)
The state or federal government should decide what categories of fetal genetic information can and	Neither Agree or Disagree	11.0% (61)
cannot be returned.	Somewhat Agree	8.5% (47)
	Strongly Agree	6.3% (35)
	No Response	3.1% (17)

Appendix 6. Respondent agreement with the statement, "it is appropriate for physicians to provide their opinion about the kinds of genetic information that parents should learn about their babies," (A) and, "the state or federal government should decide what categories of fetal genetic information can and cannot be returned" (B).



Appendix 7. Respondent preferences for receiving help from a doctor, when making a decision about which categories of information to receive by education (A), views on abortion (B), and genetic literacy (C).



Appendix 8. Respondent preference for receiving scientifically uncertain genetic information about autism.

