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Table 1. [illegible] 1.

Principle	Description
[illegible]	[illegible]
[illegible]	[illegible]
[illegible]	[illegible]

[illegible]

Informed consent

[illegible] 2 [illegible] 1 (1).

Elements of informed consent in clinical research

[illegible] (2). [illegible] 4. [illegible] 6 8 [illegible] 4.

Exceptions to the consent process

[illegible]

[illegible] 1.

Use of deception and debriefing in clinical research

[illegible] 5

Table 2. [illegible] 1

Table 3. 10, 52 .

Ability	Description
A	[illegible]

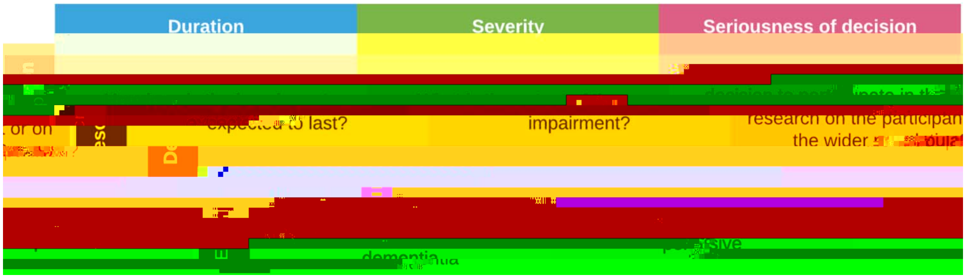

Figure 1.

10 . 12 . 13 . 1 . 14 .

12, 16 .

Participation in research for patients unable to provide self-consent

2 17 .

Impaired capacity

15 . (1). 12 . 12 . 12 .

Vulnerable populations

1 .

Figure 2. 10, 12 .

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(h) 18 . h

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h 1, 19 .

Informed consent considerations for vulnerable populations

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Table 4. [illegible] 1, 40.

	Expressed						Implied
	Written	Verbal	Telephone	Electronic	SDM	Deferred	
Description	[illegible]	[illegible]	[illegible]	[illegible]	[illegible]	[illegible]	[illegible]

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49