Table S9. Report of trophectoderm genes significantly more highly expressed in young maternal age (YMA) women, along with corresponding knockout mouse studies. Related to Figure 4.

Entrez ID	Gene name	Knockout studies/phenotypes
39	acetyl-CoA acetyltransferase 2(ACAT2)	Fertile and viable mice with abnormal embryo size (day 18.5) (Dickinson et al., 2016, JacksonLaboratory).
		Lower cholesterol levels in blood compared with the
345	apolipoprotein C3(APOC3)	control mice (JacksonLaboratory).
		Repression in the mouse embryonic development
		prior to gastrulation stage (Morris et al., 2002).
	DAB2, clathrin adaptor	Similar to lack of TGF-beta and Nodal pathway
1601	protein(DAB2)	molecules phenotype (JacksonLaboratory).
		Inhibition of early mouse embryo development at
		the morula stage (approximately 2,5 days)
		(JacksonLaboratory). Repression in the mouse
		embryonic development prior to gastrulation stage
		(Morris et al., 2002). Similar to lack of TGF-beta and
	diazepam binding inhibitor, acyl-	Nodal pathway molecules phenotype
1622	CoA binding protein(DBI)	(JacksonLaboratory).
		Inhibition of early mouse embryo development at
	enoyl-CoA hydratase, short chain	the morula stage (approximately 2,5 days)
1892	1(ECHS1)	(JacksonLaboratory).
		GPX4-knockout mice did not survive beyond E8.5
		day (or 16 days in human) (Imai and Nakagawa,
		2003, Yant et al., 2003). Low expression leads to
		DNA fragmentation and programmed cell death
2879	glutathione peroxidase 4(GPX4)	(Imai and Nakagawa, 2003, Borchert et al., 2006).
		Sensitivity to carcinogens and increased
		tumorigenesis, fibroblasts with lack of Hint1
	histidine triad nucleotide binding	expression showed problems in DNA repair and
3094	protein 1(HINT1)	indefinite growth. (JacksonLaboratory)

		LRP2-null mice phenotype shows increased
		perinatal lethality with decreased embryo size at
		E9.5 and increased levels of apoptosis (E9.5) in
		areas corresponding to neural crest
4036	LDL receptor related protein 2(LRP2)	(JacksonLaboratory).
		PRDX2-null mice increase programmed cell death
		and inhibits normal trophectoderm development,
		that could be involved with the accumulation of
		reactive oxygen species (ROS). Significantly lower
		PRDX2 expression has been associated with
		spontaneous abortions during the first trimester
7001	peroxiredoxin 2(PRDX2)	(Wu et al., 2017).
		Allows embryo implantation but not further embryo
		development. Inner cell mass knockout cells did not
5134	programmed cell death 2(PDCD2)	proliferate in vitro (Mu et al., 2010).
		WLS-null mouse phenotypes are lethal between
		E4.5 and E8 (or E9) developmental stage, which
		corresponds to stages between somite and
	wntless Wnt ligand secretion	placenta formation in human (organogenesis).
79971	mediator(WLS)	(JacksonLaboratory)
		GNAS gene mutation is embryonic lethal at the
		stage that follows the embryo implantation. (Turan
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