

Supplementary Table I~IV

Table I Extended survival of *Brcal* mutant embryos by inactivation of *Chk2*

Crosses	<i>Brcal</i> ^{+Δ} <i>Chk2</i> ^{+/-}	x	<i>Brcal</i> ^{+Δ} <i>Chk2</i> ^{+/-}
Total offspring			221
Genotypes	<i>Brcal</i> ^{ΔΔ} <i>Chk2</i> ^{+/+}		<i>Brcal</i> ^{ΔΔ} <i>Chk2</i> ^{+/-} <i>Brcal</i> ^{ΔΔ} <i>Chk2</i> ^{-/-}
Predicted number	14		28 14
Observed number	0		9 15

Table II Extended survival of *Brcal* mutant embryos by inactivation of *Chk1*

Crosses	<i>Brcal</i> ^{+Δ} <i>Chk1</i> ^{+/-}	x	<i>Brcal</i> ^{+Δ} <i>Chk1</i> ^{+/-}
Total offspring			264
Genotypes	<i>Brcal</i> ^{ΔΔ} <i>Chk1</i> ^{+/+}		<i>Brcal</i> ^{ΔΔ} <i>Chk1</i> ^{+/-} <i>Brcal</i> ^{ΔΔ} <i>Chk1</i> ^{-/-}
Predicted number	17		33 17
Observed number	2		27 ^a 0

a: Died 24 hrs after birth

Table III Aging and tumorigenesis of *Brcal*^{Δ/Δ}*p53*^{+/-} and *Brcal*^{Δ/Δ}*Chk2*^{-/-} mice

Phenotypes	Mice	
	<i>Brcal</i> ^{Δ/Δ} <i>p53</i> ^{+/-}	<i>Brcal</i> ^{Δ/Δ} <i>Chk2</i> ^{-/-}
Aging-related changes: ^a		
Decreasing of Skin thickness	75%	<10%
Decreasing of Bone density	70%	<10%
Intestinal villi atrophy	80%	<10%
Tumorigenesis:		
Spectrum	Mammary, Thymus, Spleen, Lung, Liver, Brain, Soft tissue	Mammary, Ovarian, Liver, Thymus
Frequency	High in mammary and thymus	High in mammary and ovarian
Histology	Highly diverse histologic morphology	Less diverse
Cytogenetic	Aneuploid (% high)	Aneuploid (% Low)

a: Histological analysis was performed on 8 months old mice. At 18 months of age 47% males and 12.5% of females of *Brcal*^{Δ/Δ}*Chk2*^{-/-} mice exhibited premature aging.

Table IV Extended survival of *Brcal* mutant embryos by inactivation of ATM

Crosses	<i>Brcal</i> ^{+Δ} <i>Atm</i> ^{+/-}	x	<i>Brcal</i> ^{+Δ} <i>Atm</i> ^{+/-}
Total offspring		180	
Genotypes	<i>Brcal</i> ^{ΔΔ} <i>Atm</i> ^{+/+}	<i>Brcal</i> ^{ΔΔ} <i>Atm</i> ^{+/-}	<i>Brcal</i> ^{ΔΔ} <i>Atm</i> ^{-/-}
Predicted number	11	23	11
Observed number	0	17 (3) ^a	10

a: Died 24 hrs after birth