

Table S2. Natural transformation and survival of *C. jejuni* following DNA damage.

DNA damaging agent	Concentration /intensity	Transformants/CFU^a	Recovery incubation^b	CFU/ml
Negative control (MitC)	-	2.1x10 ⁻⁶	-	5.3x10 ⁸
Mitomycin C	10 ng/ml	2.9x10 ⁻⁶	-	4.9x10 ⁸
Mitomycin C	50 ng/ml	1.5x10 ⁻⁶	-	2.3x10 ⁸
Mitomycin C	100 ng/ml	1.6x10 ⁻⁶	-	1.9x10 ⁸
Mitomycin C	200 ng/ml	3.1x10 ⁻⁶	-	1.1x10 ⁸
Mitomycin C	500 ng/ml	3.4x10 ⁻⁶	-	5.9x10 ⁶
Negative control (UV)	0 J/m ²	-	2 h	9.8x10 ⁸ ± 3.5x10 ⁷
Negative control (UV)	0 J/m ²	3.0x10 ⁻⁴ ± 3.0x10 ⁻⁵	2 h + DNA	9.5x10 ⁸ ± 2.1x10 ⁸
UV	10 J/m ²	-	2 h	2.3x10 ⁸ ± 1.1x10 ⁷
UV	10 J/m ²	3.6x10 ⁻⁴ ± 3.9x10 ⁻⁵	2 h + DNA	1.9x10 ⁸ ± 2.5x10 ⁷
UV	15 J/m ²	-	2 h	1.7x10 ⁸ ± 3.2x10 ⁷
	15 J/m ²	2.0x10 ⁻⁴ ± 8.2x10 ⁻⁶	2 h + DNA	1.6x10 ⁸ ± 3.5x10 ⁶

UV	30 J/m ²	-	2 h	4.6x10 ⁷ ± 5.3x10 ⁶
UV	30 J/m ²	2.5x10 ⁻⁴ ± 5.2x10 ⁻⁵	2 h + DNA	4.5x10 ⁷ ± 1.1x10 ⁷
UV	60 J/m ²	-	2 h	4.3x10 ⁶ ± 0
UV	60 J/m ²	1.4x10 ⁻⁴ ± 2.5x10 ⁻⁵	2 h + DNA	4.5x10 ⁶ ± 1.8x10 ⁶
UV	80 J/m ²	-	2 h	1.0x10 ⁶ ± 1.2x10 ⁵
UV	80 J/m ²	1.4x10 ⁻⁴ ± 2.4x10 ⁻⁵	2 h + DNA	7.8x10 ⁵ ± 7.1x10 ⁴

^aCam^R transformants per CFU.

^bFollowing UV treatment, one volume of BHI and when indicated 2 µg/ml isogenic chromosomal DNA carrying a Cam^R marker was added to the culture. The bacteria were allowed to recover from the UV treatment for 2 h at 37° C in a microaerobic environment before scoring transformants and CFU/ml.

Results are representative from several independent experiments.