## **Supporting Information**

## Dean et al. 10.1073/pnas.1715640115



**Fig. S1.** Fit of the rat–flea model to observed rodent and human mortality during the 1903 plague outbreak in Hong Kong. The observed rat mortality (black dots), the observed human mortality (green dots), and fit (mean and 95% credible interval) of the rat model for plague transmission to both the rat (black) and human (green) mortality. The mortality peak for humans from the model is delayed compared with the observed data. However, the model captures the dynamics of the rat mortality and the relationship between the epizootic and the epidemic well by showing the characteristic higher rat mortality and the delay in the onset of the epidemic in humans.



Fig. 52. Fit of the pneumonic and rat-flea models of plague transmission to mortality during Third Pandemic outbreaks. The observed human mortality data (black dots) for plague outbreaks and the fit (mean and 95% credible interval) of the relevant model for plague transmission in each plague outbreak: pneumonic (blue) and rat-flea (green). Both the rat-flea model of plague transmission and the pneumonic plague transmission are well capable of fitting observed human mortality patterns for plague outbreaks that these models describe.

Table S1.	Summary	/ of the	Third	Pandemic	mortality	/ data

Location	Date, MM/YYYY	Population	Recorded deaths	Transmission mode	Ref.
Sydney, Australia	02/1900–08/1900	400,000	103	Rat–flea	1
Hong Kong, China	01/1903-12/1903	250,000	1,308	Rat–flea	2
Harbin (Fuchiatien), China	12/1910–02/1911	25,000	3,223	Pneumonic	3

The present-day location, dates (month/year), preplague population size, and recorded plague deaths, and known transmission mode for three plague outbreaks during the Third Pandemic.

1. Cumpston JHL, McCallum F (1926) The History of Plague in Australia, 1900–1925 (H. J. Green Govt Printer for Commonwealth of Australia Dept Health, Melbourne). 2. Hunter W (1904) A Research into Epidemic and Epizootic Plague (Noronha and Company, Hong Kong).

3. Anonymous (1912) Report of the International Plague Conference Held at Mukden, April, 1911, ed Strong RP (Bureau of Printing, Manila, Philippines).

## Table S2. Initial conditions for three SIR models of plague transmission

Parameter	Value	Definition
Human ectopa	rasite model	
<b>S</b> <sub>h</sub> ( <b>0</b> )	U(0.001, 1)*population size	Initial susceptible humans
$I_{\text{low}}(0)$	U(1, 10*D <sub>h</sub> (0))	Initial infected (low) humans
$I_{high}(0)$	2*D <sub>h</sub> (0)	Initial infected (high) humans
$R_h(0)$	0	Initial recovered humans
$D_h(0)$	Observed deaths at $T = 0$	Initial dead humans
Pneumonic pla	gue model	
<b>S</b> <sub>h</sub> ( <b>0</b> )	U(0.001,1)*population size	Initial susceptible humans
<i>I<sub>h</sub></i> ( <b>0</b> )	U(1, 10*D <sub>h</sub> (0))	Initial infected humans
$D_h(0)$	Observed deaths at $T = 0$	Initial dead humans
Rat–flea model		
<b>S</b> <sub>r</sub> ( <b>0</b> )	U(0.001, 1)*population size	Initial susceptible rats
$I_r(0)$	U(1, 15*D <sub>h</sub> (0))	Initial infected rats
<b>R</b> <sub>r</sub> ( <b>0</b> )	0	Initial recovered rats
$D_r(0)$	0	Initial dead rats
<b>S</b> <sub>h</sub> ( <b>0</b> )	<b>S</b> <sub>r</sub> ( <b>0</b> )	Initial susceptible humans
<i>I</i> <sub>h</sub> ( <b>0</b> )	1.5*D <sub>h</sub> (0)	Initial infected humans
$R_h(0)$	0	Initial recovered humans
$D_h(0)$	Observed deaths at $T = 0$	Initial dead humans
<b>H</b> ( <b>0</b> )	Kf	Initial fleas on host
<b>F</b> ( <b>0</b> )	$K_f^*D_h(0)$	Initial free infected fleas

Single numbers are fixed values and distributions (U = uniform) are priors.

Parameter	Parameter value/prior distribution	Posterior estimate, mean [95% highest posterior density interval]
<b>S</b> <sub>h</sub> ( <b>0</b> )	S <sub>r</sub> (0)	Fixed
$I_h(0)$	5.0	Fixed
<i>R</i> <sub><i>h</i></sub> ( <b>0</b> )	0	Fixed
$D_h(0)$	Observed deaths at $T = 0$	Fixed
$\beta_h$	U(0.001, 1)	0.11 [0.10, 0.12]
$S_r(0)$	U(0.001, 1)*population size	0.018 [0.017, 0.018] * 250,000
$I_r(0)$	U(1, 23)	22.8 [22.6, 23]
$R_r(0)$	0	Fixed
$D_r(0)$	Observed deaths at $T = 0$	Fixed
β <sub>r</sub>	<i>U</i> (0.001, 1)	0.053 [0.053, 0.053]

Table S3.	Initial parameter values and posterior estimates for the rat-flea model fitted rat and
human mo	ortality in Hong Kong

Single numbers are fixed values, and distributions (U = uniform) are priors.

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Table S4.	Posterior means and 95% highest density posterior intervals for estimated parameters in three plague models for Second and
Third Panc	demic outbreaks

Location	Model	Population at risk (proportion)	Initial infected $[I_{low}(0), I_{h}(0), I_{r}(0)]$	Transmission rate $(\beta_{low}, \beta_{high}, \beta_{p}, \beta_{r}, \beta_{h})$
Givry (1348)	EP	0.75 [0.69, 0.81]	2.21 [2, 2.61]	0.04 [0.02, 0.05]
				0.39 [0.32, 0.53]
	PP	0.42 [0.38, 0.45]	1.85 [1.41, 2.32]	0.44 [0.43, 0.44]
	RP	0.73 [0.64, 0.81]	28.81 [26.60, 29.99]	0.06 [0.06, 0.06]
				0.19 [0.18, 0.2]
Florence (1400)	EP	0.36 [0.35, 0.36]	79.65 [78.99, 80]	0.049 [0.04, 0.05]
				0.32 [0.31, 0.38]
	PP	0.17 [0.17, 0.17]	79.79 [79.39, 79.99]	0.42 [0.42, 0.42]
	RP	0.19 [0.19, 0.19]	119.91 [119.76, 120.0]	0.084 [0.083, 0.085]
				0.2 [0.199, 0.2]
Barcelona (1490)	EP	0.28 [0.27, 0.28]	8.68 [7.54, 9.97]	0.032 [0.007, 0.05]
				0.49 [0.35, 0.67]
	PP	0.14 [0.13, 0.14]	9.90 [9.73, 10.0]	0.43 [0.43, 0.43]
	RP	0.14 [0.13, 0.14]	14.95 [14.87, 15.0]	0.08 [0.08, 0.08]
				0.2 [0.19, 0.2]
London (1563)	EP	0.42 [0.41, 0.42]	32.45 [29.68, 35.62]	0.04 [0.04, 0.05]
				0.27 [0.26, 0.28]
	PP	0.21 [0.20, 0.21]	50.85 [48.81, 52.99]	0.43 [0.43, 0.43]
	RP	0.30 [0.30, 0.31]	254.80 [254.43, 255]	0.06 [0.059, 0.06]
				0.2 [0.2, 0.2]
Eyam (1666)	EP	0.97 [0.92, 1.0]	3.76 [3, 4.97]	0.032 [0.01, 0.05]
				0.32 [0.2, 0.5]
	PP	0.56 [0.48, 0.63]	3.80 [3, 4.82]	0.41 [0.41, 0.42]
	RP	0.96 [0.90, 1.0]	38.08 [29.53, 44.97]	0.04 [0.04, 0.05]
				0.19 [0.18, 0.2]
Gdansk (1709)	EP	0.93 [0.92, 0.94]	51.3 [49, 54.6]	0.049 [0.046, 0.05]
				0.28 [0.26, 0.3]
	PP	0.46 [0.46, 0.47]	79.11 [76.56, 81.95]	0.42 [0.42, 0.42]
	RP	0.92 [0.90, 0.93]	734.48 [733.36, 735]	0.04 [0.04, 0.05]
				0.2 [0.2, 0.2]
Stockholm (1710)	EP	0.42 [0.41, 0.42]	159.63 [153.01, 168.35]	0.04 [0.03, 0.05]
				0.33 [0.30, 0.38]
	PP	0.22 [0.21, 0.22]	145.36 [139.14, 151.28]	0.42 [0.42, 0.42]
	RP	0.36 [0.35, 0.36]	2,290.65 [2,282.25, 2,294.99]	0.069 [0.069, 0.069]
				0.2 [0.2, 0.2]
Moscow (1771)	EP	0.34 [0.34, 0.35]	157.41 [150.41, 164.44]	0.04 [0.04, 0.05]
				0.34 [0.32, 0.39]
	PP	0.17 [0.17, 0.18]	148.31 [144.46, 152.12]	0.43 [0.43, 0.43]
	RP	0.20 [0.20, 0.21]	659.86 [659.57, 660.0]	0.069 [0.069, 0.069]
				0.2 [0.2, 0.2]
Malta (1813)	EP	0.09 [0.09, 0.09]	18.09 [16.47, 19.9]	0.04 [0.04, 0.05]
				0.26 [0.23, 0.31]
	PP	0.04 [0.04, 0.04]	9.96 [9.90, 10.0]	0.43 [0.43, 0.43]
	RP	0.045 [0.044, 0.046]	14.98 [14.939, 15.0]	0.06 [0.06, 0.06]
				0.2 [0.2, 0.2]
Sydney (1900)	EP	0.49 [0.003, 0.95]	7.49 [5.48, 9.77]	0.024 [0.0, 0.04]
				0.15 [0.0, 0.3]
	PP	0.001 [0.0, 0.001]	1.46 [1, 2.06]	0.42 [0.41, 0.42]
	RP	0.001 [0.0, 0.001]	13.559 [10.637, 15.0]	0.05 [0.04, 0.05]
				0.18 [0.14, 0.2]
Hong Kong (1903)	EP	0.011 [0.011, 0.012]	3.05 [3, 3.17]	0.048 [0.044, 0.05]
				0.24 [0.22, 0.26]
	PP	0.01 [0.01, 0.01]	2.88 [2.41, 3.35]	0.42 [0.42, 0.42]
	RP	0.011 [0.009, 0.013]	36.66 [27.63, 44.99]	0.05 [ 0.05, 0.05]
				0.16 [0.13, 0.2]
Harbin (1910)	EP	0.02 [0.02, 0.021]	33.93 [27.09, 41.58]	0.03 [0.01, 0.05]
				0.88 [0.76, 1.]
	PP	0.12 [0.12, 0.13]	16.99 [14.9, 18.98]	0.48 [ 0.48, 0.48]
	RP	0.11 [ 0.11, 0.11]	119.25 [117.66, 119.99]	0.14 [0.13, 0.14]
				0.19 [0.19, 0.2]

Posterior estimates for initial conditions for different plague models and outbreaks. Models are designated as human ectoparasite (EP), primary pneumonic plague (PP), and rat and rat–flea (RP). Posterior estimates (mean [95% highest density posterior interval]) for the proportion of the initial population at risk, the initial number of infected [I(0)], and the transmission rate ( $\beta$ ).

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Table S5.Comparison of transmission models and estimates forthe basic reproduction number for different plague models andThird Pandemic outbreaks

Location	Model	BIC	$\Delta BIC$	R <sub>o</sub>
Sydney (1900)	EP	235	46	0.86 [0.86, 0.87]
	PP	196	7	1.05 [1.05,1.05]
	RP	189	0	1.36 [1.36,1.36]
Hong Kong (1903)	EP	611	107	1.52 [1.52, 1.52]
	PP	900	396	1.06 [1.06,1.06]
	RP	504	0	1.41 [1.41,1.41]
Harbin (1910)	EP	851	31	2.98 [2.98, 2.98]
	PP	820	0	1.21 [1.21,1.21]
	RP	1,606	786	3.62 [3.62,3.62]

The models are designated as human ectoparasite (EP), primary pneumonic plague (PP), and rat and rat–flea (RP). Values in bold represent the best-fitting models that were within 10 points of the lowest BIC. The  $R_0$ (mean [95% confidence interval]) was estimated for each model using the next-generation matrix.

Table S6.	Comparison	of transmission	models with	different	levels of	underreporting

			BIC	
Location	Model	10% underreporting	25% underreporting	50% underreporting
Givry (1348)	EP	1,288	1,280	1,395
	PP	1,333	1,333	1,331
	RP	1,292	1,370	1,439
Florence (1400)	EP	2,729	2,876	3,392
	PP	4,668	4,928	5,877
	RP	10,568	11,264	12,752
Barcelona (1490)	EP	1,942	1,951	2,121
	PP	2,418	2,453	2,610
	RP	3,482	3,640	3,991
London (1563)	EP	1,582	1,577	1,575
	PP	4,630	4,629	4,629
	RP	4,256	4,954	6,743
Eyam (1666)	EP	1,176	1,175	1,243
	PP	1,174	1,174	1,238
	RP	1,210	1,228	1,304
Gdansk (1709)	EP	825	1,803	No convergence
	PP	3,817	3,817	3,817
	RP	2,176	4,447	No convergence
Stockholm (1710)	EP	718	709	688
	PP	2,180	2,109	2,110
	RP	1,238	1,612	2,759
Moscow (1771)	EP	3,916	3,916	3,931
	PP	6,790	6,790	6,790
	RP	17,604	22,612	No convergence
Malta (1813)	EP	2,760	2,775	2,864
	PP	3,653	3,850	4,244
	RP	6,632	6,953	7,656

The models are designated as human ectoparasite (EP), primary pneumonic plague (PP), and rat and rat-flea (RP). Values in bold represent the best-fitting models that were within 10 points of the lowest BIC.

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