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With more than 20 000 cases reported, the outbreak of Ebola virus disease (EVD) in west Africa is by far the largest in recorded history. Despite the scale of the current outbreak, EVD is often perceived as a "small-scale killer".¹ By comparison, malaria caused an estimated 854 000 deaths worldwide in 2013.² However, although limited at the global level, the impact of EVD on mortality could be substantial in countries with intense transmission. We thus aimed to compare EVD with other causes of death in Liberia, Sierra Leone, and Guinea in 2014.

We did an uncertainty analysis of EVD mortality (see appendix), based on two parameters: the extent of underreporting of EVD cases and the case fatality rate (CFR)—ie, the proportion of EVD cases who die. Similar to other analyses of EVD spread,³ we hypothesised that there were up to 2.5 times more EVD cases than reported. This factor derives from a mathematical model, which compared the reported number of EVD cases to the number of beds in use in Ebola treatment units in August, 2014.⁴ We assumed that the CFR varied between 60% and 85%. The lower rate corresponds to CFRs seen among hospitalised EVD patients with known disease outcomes.⁵ Lower CFRs have been documented, but only in Ebola treatment units that implement non-standard treatment protocols.6 The upper rate corresponds to CFRs seen in non-hospitalised EVD patients.⁵

We estimated the number of EVD deaths as the product of (1) the reported number of EVD cases, (2) the under-reporting factor and (3) the CFR. Based solely on confirmed and probable EVD cases, the number of EVD deaths in 2014 ranged from 2928 to 10372 in Liberia, from 4468 to 15824 in Sierra Leone, and from 1739 to 5548 in Guinea.

We used the most recent (2013) national estimates of non-EVD mortality,²⁷ together with projections of population growth, to calculate the expected number of deaths from non-EVD causes in Liberia, Sierra Leone, and Guinea in 2014. For all combinations of model parameters, we mapped how the estimated number of EVD deaths ranked relative to the expected number of deaths from non-EVD causes. In Liberia, for virtually all model parameters, EVD deaths exceeded the expected number of deaths due to the leading non-EVD cause of death (figure). In Sierra Leone, a broad range of model parameters also indicated that EVD might have killed more people in 2014 than the leading non-EVD cause of death (ie, malaria). In other sets of model parameters, EVD still killed more people than the second

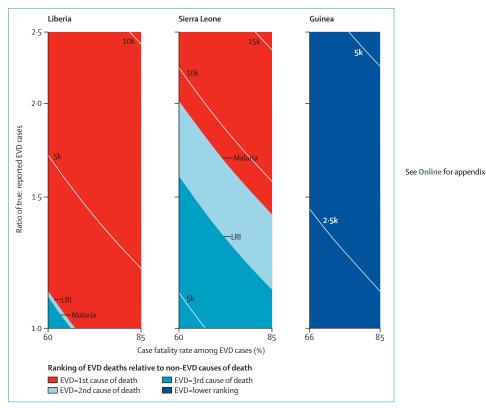


Figure: Comparison of Ebola virus disease (EVD) deaths and expected deaths from non-EVD causes in Liberia, Sierra Leone, and Guinea in 2014

LRI=lower respiratory infections. White contours represent estimated number of EVD deaths in 2014 for specific combinations of model parameters (where 2.5k = 2500 deaths, etc). Calculations of number of EVD deaths, and comparisons with other causes of death, are described in the appendix. Range of possible case fatality rates (CFRs) is narrower in Guinea than in the other two countries because a higher percentage of EVD deaths was recorded in that country among reported EVD cases. Lower bound of CFR is thus set at level estimated by surveillance data in that country. To aid figure interpretation, we illustrate the case of Sierra Leone. In the red area, there are more EVD deaths than we expect deaths from the leading non-EVD cause of death in that country-ie, malaria. Hence, in this region of the parameter space, EVD would be the first cause of death. In the light blue area, there are fewer EVD deaths than deaths from the leading non-EVD cause of death, but there are more EVD deaths than deaths from the second non-EVD cause of death (ie, LRI). Hence EVD would be the second cause of death in that region of the parameter space. In the slightly darker blue area at the bottom of the graph, there are fewer EVD deaths than deaths from LRI, but more than deaths from HIV/ AIDS—ie, the third leading non-EVD cause of death (see appendix). EVD would thus be the third leading cause of death in that region of the parameter space. The boundary between the red and light blue areas (marked "malaria") represents combinations of model parameters where the number of EVD deaths is equal to the expected number of malaria deaths. The boundary between the light blue and the darker blue areas (marked "LRI") represents combinations for which the number of EVD deaths is equal to the expected number of LRI deaths. In Guinea, the entire graph is dark blue because the number of EVD deaths is lower than the expected number of deaths from the third non-EVD cause of death in the country for all model parameters.

(ie, lower respiratory infections) or the third (ie, HIV/AIDS) leading causes of death. In Guinea, EVD never ranked higher than the top three non-EVD causes of death (figure). The highest estimate of EVD deaths was slightly lower than the expected number of deaths from diarrhoeal diseases—ie, the fifth leading non-EVD cause of death in that country (see appendix).

Our analyses have limitations. First, we do not account for increases in non-EVD deaths due to lower healthcare use during the outbreak.8 Second, the estimates of non-EVD mortality are based largely on imprecise survey or census data.² Despite statistical corrections,⁹ they might underestimate the number of non-EVD deaths.¹⁰ Third. the range of model parameters we considered might be too broad. In one study of EVD viral sequences collected in June, 2014, in Sierra Leone, there were only up to 1.7 times more cases than reported.¹¹ Case reporting might also have improved in the autumn of 2014, as new Ebola treatment units were opened. Fourth, we only include confirmed and probable EVD cases, whereas EVD deaths might also be common among suspected cases (ie, cases without laboratory or clinical data).

Nonetheless, our analyses identify a large discrepancy between the high impact of EVD on mortality in Liberia and Sierra Leone (and to a lesser extent Guinea), and its low impact on mortality at global or continental levels. Since assessments of the burden of disease should inform the allocation of health investments,12 this discrepancy could create conflicting priorities among actors of the health sector in these countries. External actors operating across multiple countries and regions (eq, donors, non-governmental organisations) might emphasise tackling diseases more common throughout sub-Saharan Africa than EVD.13 National governments, on the other hand, might stress the need to prevent future EVD outbreaks. The emergence of EVD will thus require

increased coordination between national and external actors to ensure that important components of the disease burden do not go unaddressed. Focusing on building high-quality health systems, capable of addressing a broad range of diseases including EVD, could help overcome this tension.¹⁴

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