Increased BMI and reduced efficacy of oral emergency contraception - how many women might be

affected?

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Oral emergency contraception (EC) is available as levonorgestrel and ulipristal acetate. In England, EC is available through a variety of settings and can be obtained without a prescription. In 2014, evidence from clinical studies prompted a review of available evidence by the European Medicine Agency's Committee for Medicinal Products for Human Use to assess whether increased bodyweight reduces the efficacy of EC. The review concluded that available data were inconclusive. However, if future research were to substantiate a relationship of reduced EC efficacy in women with higher bodyweight, it would be important to understand the potential impact at a population level. We used data from the Health Survey for England (HSE) 2010<sup>3</sup> to explore this.

HSE is an annually-conducted, nationally-representative probabilistic household survey of the general population resident in England. Details of HSE methodology are reported elsewhere. Demographic data are collected using face to face interviews and self-completed questionnaire booklets. Bodyweight is measured at a nurse visit. In 2010, the survey incorporated questions regarding sexual behaviour and contraceptive use, including EC use in the 12 months preceding the interview. We estimated EC use in 1,508 sexually-experienced (reporting at least one sexual partner over the lifetime to the date of the interview) 16 to 44 year-old women by bodyweight. Data were analysed in Stata 12.1 accounting for weighting, clustering and stratification of the data.

EC use in the last year was reported by 9% (95% CI 7%-11%) of all sexually-experienced 16-44 year-old women and was more frequently reported among younger women (Table 1). Among those who reported EC use in the last year, 33% (95% CI 25%-43%) had a bodyweight >75kg and 22% (95% CI 15%-31%) had a bodyweight >80kg (Figure 1).

Table 1: Percentage of women who reported oral emergency contraception use in the last year (sexually-experienced women aged 16-44 years)

	Used EC in last year		
	(95% confidence interval)		
Overall	9% (7%-11%)		
By age group (years)			
16-24	20% (16%-26%)		
25-34	8% (6%-11%)		
35-44	3% (2%-5%)		

Current guidance advises EC use regardless of bodyweight, although available data remain inconclusive with regard to the relationship with EC efficacy. Our analysis demonstrates that EC could be less effective in a substantial proportion of current EC users if efficacy were found to be lower among those with higher bodyweight. If future provision of EC is reliant on some measure of bodyweight, this would have serious implications in the delivery of EC, especially in settings where the threshold for access to EC is low, such as pharmacies. Younger women would be particularly affected given the higher rates of EC use in this group. Research is urgently needed to determine the relationship between bodyweight and EC efficacy.

Figure 1: Bodyweight of EC users by age group (sexually-experienced women aged 16-44 years)

## **Competing interests**

KAG has previously advised and given lectures for the manufacturers of ellaOne (HRA Pharma). The other authors declare no competing interests.

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