Description of Additional Supplementary Files

File Name: Supplementary Data 1

Description: Cofactors Associated with Oral Rotavirus Vaccine Shedding After Dose 1. Logistic

regression was used to explore the relationship between each covariate and ORV shedding 1 week after

dose 1. Variables with a univariate p value of <0.05 were explored in multivariate models. Measurements

obtained after dose 1 (i.e. after week of life 6) were excluded given that the aim was to identify predictors

of ORV shedding rather than shedding-induced changes. a1AG, a1 acid glycoprotein; a1AT, a1-

antitrypsin; bOPV, bivalent oral poliovirus vaccine; GMC, geometric mean concentration; IND, India;

IPV, inactivated poliovirus vaccine; LRT, likelihood ratio test for fit of model with vs without polio

vaccine schedule; MLW, Malawi; MPO, myeloperoxidase; neo+, infected with rotavirus neonatally

(defined by detection of rotavirus shedding in week of life 1 or baseline seropositivity); neo-, uninfected

with rotavirus neonatally; ORV, oral rotavirus vaccine; RR, relative risk; RV, rotavirus; tOPV, trivalent

oral poliovirus vaccine; * log-transformed to approximate normality in statistical models; † excluded to

minimise multicollinearity.

File Name: Supplementary Data 2

Description: Cofactors Associated with Oral Rotavirus Vaccine Seroconversion. Logistic regression

was used to explore the relationship between each covariate and seroconversion. Variables with a

univariate p value of <0.05 were retained in multivariate models. α1AG, α1 acid glycoprotein; α1AT, α1-

antitrypsin; bOPV, bivalent oral poliovirus vaccine; GMC, geometric mean concentration; IND, India;

IPV, inactivated poliovirus vaccine; LRT, likelihood ratio test for fit of model with vs without polio

vaccine schedule; MLW, Malawi; MPO, myeloperoxidase; neo+, infected with rotavirus neonatally

(defined by detection of rotavirus shedding in week of life 1 or baseline seropositivity); neo-, uninfected

with rotavirus neonatally; ORV, oral rotavirus vaccine; RR, relative risk; RV, rotavirus; tOPV, trivalent

oral poliovirus vaccine; * log-transformed to approximate normality in statistical models; † excluded to

minimise multicollinearity; § +2 weeks in UK due to later vaccination schedule.

File Name: Supplementary Data 3

Description: Cofactors Associated with Post-vaccination Rotavirus-specific IgA Concentration. Linear regression was used to explore the relationship between each covariate and post-vaccination RV-IgA (log-transformed). Variables with a univariate p value of <0.05 were retained in multivariate models. α1AG, α1 acid glycoprotein; α1AT, α1-antitrypsin; bOPV, bivalent oral poliovirus vaccine; GMC, geometric mean concentration; IND, India; IPV, inactivated poliovirus vaccine; LRT, likelihood ratio test for fit of model with vs without polio vaccine schedule; MLW, Malawi; MPO, myeloperoxidase; neo+, infected with rotavirus neonatally (defined by detection of rotavirus shedding in week of life 1 or baseline seropositivity); neo-, uninfected with rotavirus neonatally; ORV, oral rotavirus vaccine; RV, rotavirus; tOPV, trivalent oral poliovirus vaccine; * log-transformed to approximate normality in statistical models; †excluded to minimise multicollinearity; \$ +2 weeks in UK due to later vaccination schedule.

File Name: Supplementary Data 4

Description: Geographic Discrepancies in Genus Composition. Data are presented for (A) Malawi vs India, (B) India vs UK, and (C) Malawi vs UK. Genera are included if they had an FDR-adjusted p value of <0.05 based on either two-sided Fisher's exact test (differences in prevalence) or Aldex2 with two-sided Wilcoxon rank sum test (differences in abundance). Genera were tested if they were present in at least 5% of samples in one of the cohorts being compared. FDR, false discovery rates.