

SUPPLEMENTARY TABLE. Sensitivity analysis* of estimated nirsevimab effectiveness against medically attended acute respiratory illness and hospitalization from respiratory syncytial virus, overall and by child's respiratory syncytial virus season — Yukon-Kuskokwim Region, Alaska, October 23, 2023–June 30, 2024

Outcome/RSV season (age on Oct 1, 2023)	Nirsevimab dosage pattern	Primary analysis n = 472		Sensitivity analysis with SARS-CoV-2- and influenza-positive children removed			
				6 case-patients removed n = 466		6 case-patients and 86 controls removed n = 380	
		Total no.	Adjusted effectiveness, % (95% CI)*	Total no.	Adjusted effectiveness, % (95% CI)*	Total no.	Adjusted effectiveness, % (95% CI)*
Medically attended ARI							
Overall	No nirsevimab doses	245	Ref	240	Ref	191	Ref
	Nirsevimab receipt ≥7 days earlier	227	82 (62–91)	226	81 (58–91)	189	83 (63–92)
1st season (<8 mos)	No nirsevimab doses	131	Ref	126	Ref	102	Ref
	Nirsevimab receipt ≥7 days earlier	161	76 (42–90)	160	73 (31–90)	129	74 (33–90)
2nd season (8–19 mos)	No nirsevimab doses	114	Ref	114	Ref	89	Ref
	Nirsevimab receipt ≥7 days earlier	66	88 (48–97)	66	88 (48–97)	60	91 (60–98)
Hospitalization							
Overall	No nirsevimab doses	35	Ref	32	Ref	28	Ref
	Nirsevimab receipt ≥7 days earlier	29	93 (64–99)	29	91 (53–98)	25	93 (53–99)

Abbreviations: Ref = reference group; RSV = respiratory syncytial virus.

* Cases or cases and controls with positive results for SARS-CoV-2 or influenza viruses were excluded. Effectiveness was calculated as $(1 - \text{adjusted odds ratio}) \times 100\%$. Odds ratios were calculated using multivariable logistic regression, adjusted by age in months at medical visit (continuous), sex, calendar month of medical visit, residence community type, and presence of a high-risk underlying condition.