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Kennedy w 3 minuty odkrywa szokującą prawdę o szczepionkach

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Robert F. Kennedy Jr. został nazwany przez Anthony'ego Fauciego kłamcą, gdy twierdził, że żadna z 72 szczepionek dla dzieci w Stanach Zjednoczonych nigdy nie została przetestowana pod kątem bezpieczeństwa. Po rocznej batalii prawnej prawnicy Fauciego przyznali, że Kennedy miał rację.



Szerokie przebudzenie mediów

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Został nazwany kłamcą przez Anthony'ego Fauciego za stwierdzenie, że „żadna z 72 szczepionek zalecanych dla dzieci nigdy nie została przetestowana pod kątem bezpieczeństwa”, RFK Jr. pozwał Fauciego.

Po roku milczenia prawnicy Fauciego przyznali, że RFK Jr. przez cały czas miał rację.

„Nie ma dalszego biegu... [Mehr anzeigen](#)



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Kennedy twierdzi, że producentów szczepionek nie można pociągnąć do odpowiedzialności, ich produkty nie są testowane i nie inwestują w marketing, ponieważ rząd wymaga co roku szczepień 78 milionów dzieci w wieku szkolnym. W przemówieniu w Hillsdale College Kennedy powiedział, że jest to doskonały model biznesowy dla producentów. Gdy tylko szczepionka zostanie objęta programem, firma będzie generować roczne przychody w wysokości miliarda dolarów.

Obecnie w USA są zatwierdzone 72 szczepionki dla dzieci i 16 w innych. Ustawa o szkodach poszczepiennych z 1986 r. zwalniała producentów szczepionek z odpowiedzialności. Według Kennedy'ego trzy lata później, w 1989 r., liczba przypadków chorób przewlekłych wśród amerykańskich dzieci gwałtownie wzrosła. Przytoczył takie schorzenia, jak ADHD, zaburzenia snu, problemy z mową, autyzm, zespół Tourette'a i narkolepsja.

Coraz więcej i nowe plany

„Wskaźnik autyzmu wzrósł z jednego przypadku na 10 000 dzieci w moim pokoleniu do jednego na 34 dzieci obecnie”.

Donald Trump planuje przyznać Kennedy'emu główną rolę w dziedzinie zdrowia publicznego, ponieważ według Trumpa wie on „więcej niż ktokolwiek inny” o opiece zdrowotnej.

Na podstawie długoterminowego badania kontrolowanego placebo nie zatwierdzono ani jednej rutynowej szczepionki dla dzieci. Ani jednego.



Aarona Siriego
@AaronSiriSG · Podążać



Na podstawie długoterminowego badania kontrolowanego placebo nie zarejestrowano żadnej rutynowej szczepionki dla dzieci. Ani jednego. Dziękuję @ICANdecide za sfinansowanie starannego stworzenia tego wykresu.



None of the vaccine doses the CDC recommends for routine injection into children were licensed by the FDA based on a long-term placebo-controlled trial

Type	Doses	Age	Brand	Company	Control	Placebo?	Safety Review After Injection?	Long	Source	Note
HepB	3	Birth 1M 6M	Recombivax HB	M	None	Yes	5 days	Yes	Package insert at § 6.1	Note that to license a vaccine for children, the FDA relies upon the clinical trial conducted with children, not adults, because as the FDA explains, "It's important that the public recognize that, because young children are still growing and developing, it's critical that thorough and robust clinical trials of adequate size are completed to evaluate the safety and the immune response to a ... vaccine in this population. Children are not small adults!"
			Engerix B	G	None	Yes	4 days	Yes	Package insert at § 6.1	
DTaP	15	2M 4M/6M 15M 4Y	Infanrix	G	DTP	Yes	30 days	Yes	Package insert at § 6.1	DTP was also not licensed based on a placebo controlled trial and it increases mortality .
			Daptacel	S	DT or DTP	Yes	Up to 2 months + 1 trial 6 months	Yes	Package insert at § 6.1	
PCV	4	2M 4M 6M 12M	Prevnar 13, PCV-13	P	Prevnar 7	Yes	6 months	Yes	Package insert at § 6.1	Prevnar 2 trial's control was an "[i]nvestigational meningococcal group C conjugate vaccine." In Prevnar 13 trial, "[s]erious adverse events reported following vaccination in infants and toddlers occurred in 8.2% among Prevnar 13 recipients and 7.2% among Prevnar 7 recipients." In Vaneuvance trial, "serious adverse events... were reported by 9.6% of VAXNEUVANCE recipients and by 8.9% of Prevnar 13 recipients" but deemed "safe" because "no notable patterns or numerical imbalances between vaccination groups." Prevnar 20 had similar result split into "serious adverse events" and "newly diagnosed chronic medical conditions."
			Vaneuvance PCV-15	M	Prevnar 13	Yes	6 months	Yes	Package insert at § 6.1	
			Prevnar 20, PCV-20	P	Prevnar 13	Yes	6 months	Yes	Package insert at § 6.1; Clinical review	
IPV	4	2MM 6M 4Y	IPOL	S	None	Yes	3 days	Yes	Package insert at 14-17	IPOL is <i>very different</i> than the polio vaccine created by Jonas Salk in the 1950s (used until 1960s). Hence, trials of Salk's vaccine from the 1950s were not relied upon to license IPOL.
Hib	3 or 4	2M 4M 6M 12M	ActHib	S	HepB	Yes	30 days	Yes	Package insert at § 6.1; Basis of Approval at 8	Within 30 days of injection in the ActHib trial, 3.4% experienced a serious adverse event but "[n]one was assessed by the investigators [Sonali] as related to the study of vaccines."
			Hiberix	G	HB-TITER or other vaccine	Yes	31 days	Yes	Package insert at § 6.1; Clinical review at 20-21	
			Liquid PedvaxHib	M	Lypophilized PedvaxHib	Yes	3 days	Yes	Package insert at 6-8	
RV	2 or 3 or 6M	2M 4M 6M	Rotarix	G	None	Yes	11 days + 1 year for intussusception	Yes	Package insert at § 6.1; Clinical review at 23-24	[T]here were 68 (0.19%) deaths following ROTARIX and 50 (0.15%) deaths following placebo... The most common... cause... was pneumonia... observed in 19 (0.05%) recipients of ROTARIX and 10 (0.03%) placebo recipients." Its clinical review admits "[t]he placebo consisted of all components of Rotarix, but without any RV particles." The package insert for RotaTeq similarly admits its "placebo" contains multiple ingredients as seen to the left.
			RotaTeq	M	None	Yes	42 days + 1 year for intussusception	Yes	Package insert at § 6.1; Clinical reports at 445 etc.	
Covid19	3	6M 7M 10M	Cominarty	P	Placebo	Yes	6 months	Yes	Package insert at § 6.1	Cominarty licensed for only 12+ (Spikevax, Moderna, only 18+). Package controls unblinded and most vaccinated during the trial. All data 16+ is combined but 12-15 data is separate, had 1,131 vaccinated children, and one participant shows how this trial was conducted.
Flu	19	6M Yearly	Various	None	None	Yes	Flu shots change annually without any clinical trial	Yes	CDC 22-23 Flu Shots; FDA Flu Shots	The trials of the original flu shot formulations for children also did not have a placebo control (see pp. 11-14) even though some adult trials did. The one inhaled influenza vaccine had a placebo but, again, it changes every year and is not safety tested in any trial.
MMR	6	12M 4Y	M-M-R-II	M	None	Yes	42 days	Yes	Clinical reports	M-M-R-II trials totaled only 334 children and a third developed gastrointestinal issues and a third respiratory issues. In Proria trial, both vaccine groups had high rate of serious adverse events, emergency room visits, and new chronic diseases (e.g., autoimmune disorders, asthma, type 1 diabetes, celiac, and allergies). See Table 6 of the Supplementary Materials.
			Proria	G	M-M-R-II	Yes	6 months	Yes	Package insert at § 6.1; Sup materials at 12	
VAR	2	12M 4Y	Varivax	M	45 mg of neomycin per milliliter	Yes	70 days	Yes	Package insert at § 6.1; Merck study at 2; Clinical reports	One controlled trial with 956 children, half Varivax and half neomycin, and one trial with 32 vaccinated and another 29 vaccinated 8 weeks later, during which the first group had double the ear infections and 50% more respiratory infections.
HepA	2	12M 18M	Havrix	G	Engerix-B	Yes	6 months	Yes	Package insert at § 6.1	Trials for both occurred at the same time when there was no licensed Hep A vaccine and hence no excuse for not using a placebo control. It is also startling: Engerix B, see above, was the control for Havrix, and an injection of cyto-and-neuro toxic substances, AAHS and thimerosal, were used as a control for Viocta instead of a saline injection.
			Viocta	M	AAHS and Thimerosal	Yes	42 days	Yes	Package insert at § 6.1; Merck study at 454	
Tdap	3	11Y	Adacel	S	Td, for adults	Yes	6 months	Yes	Package insert at § 6.1	Due to reactions, Tdap (Adacel) given at 11Y has 12.5 times less diphtheria toxin (25U v 2U) and 10 times less pertussis toxin (25mcg v 2.5mcg) than DTaP (Infanrix) given to babies.
			Boostrix	G	Diphtheria or Adacel	Yes	6 months	Yes	Package insert at § 6.1	
HPV	2 or 3	9Y or 9.5Y	Gardasil 9	M	Gardasil 4 (see note)	Yes	1 month in five trials, 6 months in one trial, and 4 years in one trial	Yes	Clinical review at 17-19	Gardasil 9 trial gave 306 people placebo after full series of Gardasil 4. In Gardasil 4's trial, controls received aluminum adjuvant, AAHS, except 320 people labeled "saline Placebo" that actually received all vaccine ingredients except antigens and AAHS. Across trials, 2.3% receiving vaccine or aluminum adjuvant (used to induce autoimmunity) had a suspected autoimmune disorder.
			Menactra	S	Menomune	Yes	6 months	Yes	Package insert at § 6.1	Incredibly, the safety section of the package insert for Menomune lists the trial in which it was used as a control for the trial of Menactra. This provides another good example of the safety pyramid scheme in which Menomune is licensed without a placebo-controlled trial and then used as the control to license Menactra. Menactra is then used as the control to license
MenA	2	11Y	Menveo	G	Menactra or	Yes	6 months	Yes	Package insert at § 6.1	

