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*throughout the process fda works closely with the company producing the vaccine to evaluate the vaccine s safety and effectiveness all safety concerns must be addressed before fda licenses a vaccine every batch of vaccines is tested for quality and safety once a vaccine is approved it continues to be tested, vaccine safety evaluation practical aspects in assessing* - *although smaller pre licensure safety packages may be submitted in settings of public emergency such as pandemic influenza or with new vaccine formulations of existing vaccines most new vaccines are submitted for license approval with an extensive safety data package containing detailed safety information from pre clinical studies and from the thousands or even tens of thousands of individuals that received the vaccine in clinical trials, do your part for vaccine safety what types of events* - *do your part for vaccine safety safety of vaccines after they are licensed vaers vaers is unable to determine that a vaccine caused or did not cause an adverse event sometimes people who are vaccinated get sick from another cause unrelated to the vaccine, vaccine basics vaccine safety* - *vaccines are very safe the united states long standing vaccine safety system ensures that vaccines are as safe as possible in fact currently the united states has the safest most effective vaccine supply in its history safety monitoring begins with the u s food and drug administration fda the agency ensures the safety effectiveness and availability of vaccines for the united states, vaccine safety do combination vaccines or simultaneous* - *see the do vaccines cause seizures summary for more details febrile seizures occur at a rate of about 1 per 2 500 doses of mmr the risk is increased to approximately 1 per 1 250 doses if the first dose is the combination vaccine mmrv proquad 17 18, who clinical evaluation of vaccines* - *clinical evaluation is a critical step to support the approval of vaccines given the limitations of animal models clinical studies to demonstrate the safety and efficacy of vaccines must be conducted in humans before a vaccine can be released for widespread use, evaluation of the efficacy of vaccines drug guide* - *nonliving vaccine products require a much larger antigenic mass dose and more frequent booster injections than do live vaccines to provide adequate levels of protection this is exemplified by the requirement for multiple boosters of immunization against diphtheria tetanus and pertussis compared with the single administration required for*

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