



Questions on Immunization and Vaccination and Short Answers

Bağışıklama ve Aşı ile İlgili Sorular ve Kısa Cevaplar

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Question 1: How should the approach be in situations where the child's vaccination status is unknown or unclear?

A child's vaccination status can be found out by reviewing the history, vaccination card and medical records of the child. Among these, history is known to be insufficiently reliable. Since it is administered annually, only the oral history of flu vaccination can be accepted.

In order to determine the vaccination status of the individuals, information should be received from the vaccination card or vaccination records of the healthcare center. In individuals whose vaccination status cannot be reached on paper or digitally, routine serologic test is not recommended due to reasons such as difficulty in implementation and the failure of the non-standard methods in showing protection. Therefore, all children and adults without proper vaccination card or vaccination record should be started to be vaccinated by accepting them as unvaccinated.

In the event of not having any records of vaccination of the individual, general information on the possibility of the vaccinations being administered more than the recommended dosage or more frequently than recommended:

- It is known that the frequency of side effects decreases in the repeat doses of live vaccines.
- In children aged under 6 years with no records of vaccination status, BCG vaccination does not have to be repeated if there is a BCG scar.

All individuals, whose vaccination status is not known, should be vaccinated anew with the age-appropriate immunization calendar of the unvaccinated person. However, if the decision not to continue with the repeat doses of the vaccine has been made by the ASIE Commission (ASIE: adverse effects after vaccination; see ASIE Monitoring System Notice) due to previous vaccine-related systemic or local serious side effects, then these individuals should be evaluated as incomplete vaccination.

Question 2: What are the protective durations of the vaccinations?

The potentially shortest protective durations of some vaccines are given in Table 1.

Question 3: Can vaccinations be administered from an anatomic site with a tattoo on?

Yes, they can. Vaccinations administered intramuscularly or subcutaneously can be administered through the site with the tattoo.

Question 4: Are there any vaccines to beware of if more than one vaccine will be administered simultaneously?

There is no problem in administering the vaccines on the same day. More than one vaccine can be administered on the same day. Particularly, in children who are late at the vaccination calendar may have to be vaccinated with multiple vaccines on the same day. Hepatitis B, BCG, DaBT-IPA-Hib, CPV, OPV, MMR, varicella, and hepatitis A vaccines can be administered

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on the same day. Administering many vaccinations on the same day does not suppress immunity.

- There are two exceptions where two vaccinations cannot be done on the same day though there is indication for both vaccines:
 - Conjugate pneumococcus and polysaccharide pneumococcus are not administered on the same day in risk groups who need to be vaccinated with both (for instance, chronic pulmonary, chronic heart, chronic kidney diseases, immunosuppressed individuals, those with hyposplenism/asplenism, those with diabetes mellitus, and etc.). Polysaccharide pneumococcus vaccines are not used before the age of 2 years. Polysaccharide pneumococcus vaccine can be administered at least 8 weeks after the completion of age-appropriate conjugate pneumococcus doses in children between the age of 2-18 years. In adults (≥ 19 years), first the conjugate pneumococcus vaccine is administered and then the polysaccharide vaccine is administered at least 8 weeks or 1 year later (according to the presence of risk).
 - Both pneumococcus and meningococcus vaccines are indicated in those with congenital or acquired asplenia/hyposplenism. However, it is known that simultaneous administration of CPV-13 and 4-valent conjugate meningococcus vaccine of the brand Menactra (Men ACWY-D) causes a decrease in response to the 3 serotypes of pneumococcus. Therefore, if meningococcus vac-

cine of the brand Menactra is to be administered in those with congenital or acquired asplenia/hyposplenism, it should be given at least 4 weeks after the completion of the CPV-13 series. There is no need for such duration in conjugate meningococcus vaccines of the Nimenrix or Menveo brand vaccinations in this risk group. They can be administered simultaneously with CPV-13 vaccine or without any time interval.

- Paracetamol (first dose, half an hour before the vaccine administration, two other doses once in every 6 hours, hence in total, 3 doses) is recommended when meningococcus serogroup B, a non-routine vaccine, is administered simultaneously with any of the vaccines due to fact that it causes high fever. Conjugate meningococcus vaccines (Menveo, Nimenrix or Menactra) and meningococcus serogroup B vaccine (Bexsero) can be administered simultaneously or without any time interval.
- Oral polio and rota vaccines can be administered simultaneously with childhood vaccines (including BCG) or without any time interval. If vomiting occurs in the 10 minutes of oral polio administration, then the vaccine is repeated. If vomiting occurs following the administration of rotavirus vaccine, then the vaccine is not repeated. When oral polio and oral rotavirus vaccines are administered simultaneously, oral polio should be preferred to be given first. Oral rotavirus vaccine can be given 5 minutes later.

Table 1

Vaccination	Protectiveness	Evaluation
Pertussis	4-6 years	Immunity acquired by infection is also weak. Booster dose is recommended after the age of 11.
Diphtheria	About 10 years	Booster dose is recommended between 45-65 years.
Tetanus	13-14 years: 96%, > 25 years: 72%	Booster dose is recommended between 45-65 years.
Polio	At least 18 years > 99%	Booster dose is recommended in the case of travelling to risky regions.
HibB	> 9 years (accepted lifelong)	With very good immunological response, it is accepted to be protective lifelong.
Hepatitis B	> 20 years (accepted lifelong)	Shown to be protective for 20 years but considered to be protective lifelong.
Measles	> 96%, lifelong	Herd immunity is vital to protect infants too young to be vaccinated and those that cannot get the MMR vaccine.
Mumps	>10 years 90% protection, immunity decreases in time	Protective duration of the vaccine changes from person to person. It does not show lifelong protection like measles and rubella.
Rubella	> 15-20 years, > 90%	Herd immunity is vital to protect infants too young to be vaccinated and those that cannot get the MMR vaccine.
Pneumococcus	> 4-5 years, for conjugated vaccines	It has been shown that antibody concentrations remain high up to 4-5 years. Herd immunity gained by vaccinating children decreases diseases in the community at every age.
Human papillomavirus	> 5-8 years	Antigen response points that it provides a long-term protection. Herd immunity is observed.
Varicella	1 dose, unknown 2 doses, > 14 years	Herd immunity is vital.
Measles-mumps-rubella	> 15-20 years	Herd immunity is vital to protect infants too young to be vaccinated and those that cannot get the MMR vaccine.

The Immunization Advisory Centre. Efficacy and Effectiveness. Available from: <http://www.immune.org.nz/vaccines/efficiency-effectiveness> Accessed on: 24.05.2019.