Tolerable upper limits for vitamin D:

In this guide

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- 1. <u>Background Statement on vitamin D Exposure Levels in Formula Fed</u> Infants and Children
- 2. <u>Introduction Statement on vitamin D Exposure Levels in Formula Fed</u> Infants and Children
- 3. Limits for vitamin D content in infant and follow-on formulae
- 4. Tolerable upper limits for vitamin D:
- 5. <u>Exposure assessment Statement on vitamin D Exposure Levels in Formula</u> Fed Infants and Children
- 6. <u>Risk characterisation Statement on vitamin D Exposure Levels in Formula</u> Fed Infants and Children
- 7. <u>Summary & conclusions Statement on vitamin D Exposure Levels in Formula</u> Fed Infants and Children
- 8. References Statement on vitamin D Exposure Levels in Formula Fed Infants and Children
- 9. <u>Abbreviations Statement on vitamin D Exposure Levels in Formula Fed</u> Infants and Children
- 10. <u>Annex A Statement on vitamin D Exposure Levels in Formula Fed Infants</u> and Children
- 11. <u>Annex B Statement on vitamin D Exposure Levels in Formula Fed Infants</u> and Children
- 11. In 2012, the European Food Safety Authority (EFSA) Panel on Dietetic Products, Nutrition and Allergies (NDA) established tolerable upper levels (TULs) for vitamin D (EFSA, 2012), based on a risk assessment conducted in 2003 by the Scientific Committee on Food (SCF, 2003). The SCF risk assessment used hypercalcaemia as the adverse effect induced by excessive vitamin D exposure. The TULs established by EFSA in 2012 were as follows:
 - For infants (birth to 1 year of age), the TUL was 25 µg per person, per day.

- For children aged 1 to 4 years, the TUL was 50 μg per person, per day.
- 12. In 2014, the COT published a statement on the adverse effects of high levels of vitamin D, in which the COT agreed with the TULs set by EFSA in 2012 (COT, 2014).
- 13. However, in 2018, based on the overall evidence, the EFSA NDA Panel kept the TUL of 25 μ g/day for infants up to 6 months old, but set a new UL of 35 μ g /day for infants aged 6-12 months (EFSA, 2018). A summary of EFSA's rationale for this is provided in Annex A. The TUL for toddlers above 1 year of age (50 μ g per person, per day) was not changed in EFSA's 2018 assessment following its original establishment in 2003 and confirmation in 2012.
- 14. The COT reviewed the rationale for the revised TUL for 6 <12 month-olds of 35 μ g/person/day established by the EFSA NDA Panel and agreed with the revised TUL for this age group.