



"Hot chilli peppers" Health risks of capsaicin(oids)

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Emerging risk "Challenges with extremely hot chips"

- Poisoning cases, including hospitalizations and even some fatalities have been associated with exposure to extremely hot chips
- Approx. 40 cases associated with oral exposure to hot chips were registered by German poisoning centres between 2021 and 2023
 - ⇒ Severity: mostly mild and some moderate clinical symptoms
 - ⇒ Symptoms: include gastrointestinal and circulatory complaints



Hot tortilla chips



Relevant ingredients

- Capsaicinoids
- Occurrence in *Capsicum* species in highly variable amounts (< 0,3 to > 1 % per dried matter)
- Total capsaicinoids

Capsaicin \rightarrow 63-77 % \rightarrow 16,000,000 Scoville

Dihydrocapsaicin \rightarrow 20-32 % \rightarrow 16,000,000 Scoville

Nordihydrocapsaicin \rightarrow 1-8 % \rightarrow 9,100,000 Scoville

Minor congeners



Various *Capsicum* species



Hazards of capsaicinoids following oral ingestion

- Gastrointestinal irritation
- Neurophysiological effects (blood pressure alterations etc.)

Mode of action

- Capsaicinoids: agonistic activity at TRPV1 (heat receptor), located, e.g., at sensory peripheral neurons
 - (a) Signal transduction to the CNS
 - → Pain sensation
 - (b) Release of substance P and calcitonin gene-related-peptide (CGRP)
 - → Activation of the unspecific immune system
 - → Neurogenic inflammation (e.g. vasodilatation)



Risk assessment

- BfR's current risk assessment is based on detrimental effects to the stomach mucosa observed in a human intervention study (Myers et al. 1987)
 - Parietal secretion, Pepsin secretion, Gastric cell exfoliation (reflected by DNA loss into gastric content)
 - \rightarrow NOAEL: 8,3 mg hot chilli pepper/kg bw corresponding to approx. 12-83 µg capsaicin/kg bw
 - \rightarrow LOAEL: 25 mg hot chilli pepper/kg bw corresponding to approx. 36-250 µg capsaicin/kg bw

◆ BfR is currently working on an updated risk assessment for capsaicin(oids)



Challenges for risk assessment

- Activation of TRPV1 leads to a very complex downstream effect cascade affecting certain endpoints
- Effect vs. adversity (relevance of certain endpoints)
- High interindividual (and also intraindividual) variability in susceptibility
 - Tolerance development
- Relevance of matrix effects
- Relevance: systemic dose vs. local (bolus) concentration at gastrointestinal mucosa





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