

REACH
RoHS 3
MDR

REACH
Registration, Evaluation, Authorisation and Restriction of Chemicals
Specifically Annex XIV ("Authorisation")

EUROPEAN CHEMICALS AGENCY
DEHP entry added to the candidate list for SVHC due to its **reprotoxic properties** [57c].

EUROPEAN COMMISSION
DEHP **reprotoxic properties** added to REACH. Exemptions granted for medical devices.

EUROPEAN CHEMICALS AGENCY
DEHP entry added to the candidate list for SVHC due to its **endocrine disrupting properties to the environment** [57(f)].

Sunset date for the use of DEHP.
Exemptions granted for medical devices.

EUROPEAN CHEMICALS AGENCY
DEHP entry added to the candidate list for SVHC due to its **endocrine disrupting properties for human health** [57(f)].

EUROPEAN CHEMICALS AGENCY
Eight-week public consultation on draft recommendation.

EUROPEAN CHEMICALS AGENCY
Draft recommendation published, considering whether exemptions are still justified.

EUROPEAN CHEMICALS AGENCY
90-day public consultation on draft recommendation.

MEMBER STATE COMMITTEE
Opinion on draft recommendation adopted.

EUROPEAN CHEMICALS AGENCY
Recommendation published.

EUROPEAN COMMISSION
DEHP **endocrine disrupting properties to human health and the environment** added to REACH.

EUROPEAN COMMISSION
Latest application date for authorisation.

Sunset date after which use of DEHP is banned.

RoHS 3 (EU 2015/863)
Amendment to the Restriction of Hazardous Substances Directive adding DEHP

EUROPEAN COMMISSION
RoHS 3 adopted.

OFFICIAL JOURNAL OF THE EUROPEAN UNION
RoHS 3 published in the official journal.

MEMBER STATES
Regulations and provisions to comply with RoHS 3 adopted by member states.

RoHS 3 enters into force.

RoHS 3 goes into effect for medical devices containing DEHP.

MDR
Medical Device Regulation

EUROPEAN COMMISSION
Proposal for regulation.

COUNCIL OF THE EU
MDR adopted.

EUROPEAN PARLIAMENT
MDR adopted.

OFFICIAL JOURNAL OF THE EUROPEAN UNION
MDR published in the official journal.

MDR enters into force.

Provisions applied.
Including the justification for the presence of CMR and ED substances, such as DEHP.

Deadline for CE marking of all new and legacy medical devices.
Presence of DEHP must be duly justified according to the latest relevant SCHEER guidelines.



Year	Date	Event
2008	28 October	EUROPEAN CHEMICALS AGENCY: DEHP entry added to the candidate list for SVHC due to its reprotoxic properties [57c].
2011	February	EUROPEAN COMMISSION: DEHP reprotoxic properties added to REACH. Exemptions granted for medical devices.
2012	26 September	EUROPEAN COMMISSION: Proposal for regulation.
2014	12 December	EUROPEAN CHEMICALS AGENCY: DEHP entry added to the candidate list for SVHC due to its endocrine disrupting properties to the environment [57(f)].
2015	21 February	Sunset date for the use of DEHP. Exemptions granted for medical devices.
	31 March	EUROPEAN COMMISSION: RoHS 3 adopted.
	4 June	OFFICIAL JOURNAL OF THE EUROPEAN UNION: RoHS 3 published in the official journal.
2016	31 December	MEMBER STATES: Regulations and provisions to comply with RoHS 3 adopted by member states.
2017	7 March	COUNCIL OF THE EU: MDR adopted.
	5 April	EUROPEAN PARLIAMENT: MDR adopted.
	5 May	OFFICIAL JOURNAL OF THE EUROPEAN UNION: MDR published in the official journal.
	25 May	MDR enters into force.
	6 July	EUROPEAN CHEMICALS AGENCY: DEHP entry added to the candidate list for SVHC due to its endocrine disrupting properties for human health [57(f)].
2018	5 June – 6 August	EUROPEAN CHEMICALS AGENCY: Eight-week public consultation on draft recommendation.
	12 December	EUROPEAN CHEMICALS AGENCY: Draft recommendation published, considering whether exemptions are still justified.
	12 December –	
2019	12 March	EUROPEAN CHEMICALS AGENCY: 90-day public consultation on draft recommendation.
	26 June	MEMBER STATE COMMITTEE: Opinion on draft recommendation adopted.
	10 July	EUROPEAN CHEMICALS AGENCY: Recommendation published.
	22 July	RoHS 3 enters into force.
2021	26 May	RoHS 3 goes into effect for medical devices containing DEHP.
	22 July	EUROPEAN COMMISSION: DEHP endocrine disrupting properties to human health and the environment added to REACH.
	21 December	
2023	27 November	EUROPEAN COMMISSION: Latest application date for authorisation.
2024	26 May	Deadline for CE marking of all new and legacy medical devices. Presence of DEHP must be duly justified according to the latest relevant SCHEER guidelines.
2025	27 May	Sunset date after which use of DEHP is banned.