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# Heptavac P Plus

Awtorizzat

- *Clostridium novyi*, strain 754, toxoid
- *Clostridium tetani*, strain S1123/91, toxoid
- *Clostridium septicum*, strain 505, toxoid
- *Clostridium perfringens*, type D, strain 603, epsilon toxoid
- *Clostridium perfringens*, type C, beta toxoid
- *Bibersteinia trehalosi*, serotype T15, strain S1105/84, Inactivated
- *Bibersteinia trehalosi*, serotype T10, strain S1075/81, Inactivated
- *Bibersteinia trehalosi*, serotype T4, strain S1085/81, Inactivated
- *Bibersteinia trehalosi*, serotype T3, strain S1109/84, Inactivated
- *Mannheimia haemolytica*, serotype A9, strain S994/77, Inactivated
- *Mannheimia haemolytica*, serotype A7, strain S1078/81, Inactivated
- *Mannheimia haemolytica*, serotype A6, strain S1084/81, Inactivated
- *Mannheimia haemolytica*, serotype A2, strain S1126/92, Inactivated
- *Mannheimia haemolytica*, serotype A1, strain S1006/77, Inactivated
- *Clostridium chauvoei*, strain 1048, cells and equivalent toxoid
- *Clostridium chauvoei*, strain 658, cells and equivalent toxoid
- *Clostridium chauvoei*, strain 657, cells and equivalent toxoid
- *Clostridium chauvoei*, strain 656, cells and equivalent toxoid
- *Clostridium chauvoei*, strain 655, cells and equivalent toxoid

## Informazzjoni dwar il-prodott

### Isem tal-mediċina:

Heptavac P Plus

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### Sustanza attiva:

Disponibbli biss fi [Ingliz](#)

Disponibbli biss fi [Ingliz](#)

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### Speċi li fuqhom ser jintuża l-prodott:

Disponibbli biss fi [Bulgaru](#) [Spanjol](#) [Ċek](#) [Daniz](#) [Ġermaniz](#) [Estonjan](#) [Grieg](#) [Ingliz](#) [Franciz](#) [Taljan](#) [Latvjan](#) [Litwan](#) [Ungeriz](#) [Olandiz](#) [Rumen](#) [Finlandiz](#) [Żvediz](#) [Iżlandiz](#) [Norwegian](#)

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### Metodu ta' amministrazzjoni:

Użu għal taħt il-ġilda

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## Dettalji tal-prodott

### Sustanza attiva / Qawwa:

Disponibbli biss fi Ingliz  
3.50 international unit(s) / 1.00 millilitre(s)

Disponibbli biss fi Ingliz  
2.50 international unit(s) / 1.00 millilitre(s)

Disponibbli biss fi Ingliz  
2.50 international unit(s) / 1.00 millilitre(s)

Disponibbli biss fi Ingliz  
5.00 international unit(s) / 1.00 millilitre(s)

Disponibbli biss fi Ingliz  
10.00 international unit(s) / 1.00 millilitre(s)

Disponibbli biss fi Ingliz  
5000000.00 cells / 0.01 millilitre(s)

Disponibbli biss fi Ingliz  
5000000.00 cells / 0.01 millilitre(s)

Disponibbli biss fi Ingliz  
5000000.00 cells / 0.01 millilitre(s)

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Disponibbli biss fi Ingliz  
5000000.00 cells / 0.01 millilitre(s)

Disponibbli biss fi Ingliz  
0.50 Protective Dose / 1.00 millilitre(s)

Disponibbli biss fi Ingliz  
0.50 Protective Dose / 1.00 millilitre(s)

Disponibbli biss fi Ingliz  
0.50 Protective Dose / 1.00 millilitre(s)

Disponibbli biss fi Ingliz  
0.50 Protective Dose / 1.00 millilitre(s)

Disponibbli biss fi Ingliz

0.50 Protective Dose / 1.00 millilitre(s)

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**Forma farmaċewtika:**

Suspensjoni għall-injezzjoni

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**Perjodu ta' rtirar skont ir-rota tal-għoti:**

**Użu għal taħt il-ġilda:**

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**Sheep**

- Meat and offal. 0 day

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**Kodiċi veterinarju anatomiku terapewtiku kimiku (ATCvet):**

QI04AB05

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**Status legali tal-provvista:**

Disponibbli biss fi [Ċek](#) [Estonjan](#) [Ingliz](#) [Franciż](#) [Taljan](#) [Latvjan](#) [Litwan](#) [Portugiż](#) [Rumen](#) [Sloven](#) [Finlandiż](#) [Żvediż](#) [Norwegian](#)

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**Status tal-awtorizzazzjoni:**

Valid

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**Awtorizzat fi:**

Disponibbli biss fi [Spanjol](#) [Ċek](#) [Germaniż](#) [Estonjan](#) [Ingliz](#) [Franciż](#) [Taljan](#) [Olandiż](#) [Portugiż](#) [Slovakk](#) [Żvediż](#) [Iżlandiż](#) [Norwegian](#)

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**Disponibbli fi:**

Ireland

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**Deskrizzjoni tal-pakkett:**

Disponibbli biss fi [Ingliz](#)

Disponibbli biss fi [Ingliz](#)

Disponibbli biss fi [Ingliz](#)

Disponibbli biss fi [Ingliz](#)

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**Tagħrif addizzjonali**

**Tip ta' intitolament:**

Disponibbli biss fi [Ingliz](#) [Franciz](#) [Kroat](#) [Taljan](#) [Latvjan](#) [Finlandiz](#) [Zvediz](#) [Izlandiz](#) [Norwegian](#)

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**Bazi ġuridika tal-awtorizzazzjoni tal-prodott:**

Disponibbli biss fi [Ingliz](#) [Taljan](#)

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**Id-detentur tal-awtorizzazzjoni għat-tqegħid fis-suq:**

Intervet (Ireland) Limited

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**Data tal-awtorizzazzjoni għall-kummerċjalizzazzjoni:**

21/03/2003

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**Siti ta' manifattura b'rilaxx tal-lott:**

Intervet International B.V.

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**Awtorità responsabbli:**

Health Products Regulatory Authority

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**Numru tal-awtorizzazzjoni:**

VPA10996/146/001

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**Data tal-bidla fl-istatus tal-awtorizzazzjoni:**

21/03/2003

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To consult adverse reactions on veterinary medicinal products please go to [www.adrreports.eu/vet](http://www.adrreports.eu/vet)

## Dokumenti

Sommarju tal-karatteristiċi tal-prodott

Dan id-dokument ma jeżistix f 'din il-lingwa (Malti). Tista' ssibha b 'lingwa oħra hawn taħt.