

Obiltoxaximab SFL (*obiltoksaksymab*)

DfnY[` X'k]YXnmibU'hYa Uh`Y_i 'CV]hcl UI]a UV'G: @]i nUgUXb]Y i Xn]Y`Yb]U' pozwolenia na dopuszczenie do obrotu w UE

7nma 'Ygh`Y_ 'CV]hcl UI]a UV'G: @]k 'U]a 'W'i 'g] [c għegħgi 'Y

CV]hcl UI]a UV'G: @'Ygh`Y_]Ya ġħegħġek Ubma 'k 'hYfUd]] UbħmV]chmel k Y^k 'YWMYb]ji 'k [`]U'k n]Yk bY[c' – dck U bY^WcfċVm k nk cħubY^dfnYn'VU_hYf] 'Bacillus anthracis" HYfa]b' tk n]Yk bM' cnbUVMUž' Y'cgħċi VU' nUWcfi 'Y'n'dck cXi 'k XmWUb]U'nUfcXb]Uk z' hċċi fcnk]U^ 'g] 'k 'U_hnk bY'VU_hYf]Y] i k U'b]U^ szkodliwe toksyny.

@Y_ ghiegħgi 'Y'g] fDlk b]Y 'k 'W'i 'nUdcV]Y Yb]U'k [`]U'k n]Yk bY[c'i 'cgħiż- hċċi a]Uūm_cħbHJU_hn' zarodnikami bakterii i w przypadku braku innego odpowiedniego leczenia.

Gi VgħubW 'Wimbib 'nuk Ufh 'k 'Y_i 'CV]hcl UI]a UV'G: @'Ygh'cV]hcl_gU_gma UV"

NY'k n[` X'k 'bU'hċż' Y'k [`]U'k n]Yk bY[c'i pbUbc nU'WcfċV 'fnUX_c 'k mgh di ^ Wżk 'Xb]i & ('g]Yfdb]U' 2018 r. lek Obiltoxaximab SFL uznano za lek sierocy (lek stosowany w fnUX_]W 'WcfċVUVMŁET"K] W^]bZfa UW]bU'hYa UhdfnmbUb]U'ghJi għi 'Y_i 'g]YfcW[c'a c bU'nbU'Y 'h hU^ ema.europa.eu/medicines/human/orphan-designations/eu3182065.

>U_ 'ghiegħġek U 'Y_ 'CV]hcl UI]a UV'G: @

@Y_ Xcgh dbmibU'fYWDh " @]i bU'Y mdcXUk U 'k 'd'UWIEK Wżk 'hċċi dck U bY'fYU_WY'U'Yf[]WbY'a c bU' k 'gnmV_]gdċgħċi 'dcXXU 'YWMYb]ji "

@Y_ 'CV]hcl UI]a UV'G: @dcXU'Y'g] 'k 'dc'YXnmibWma 'k 'Yk]Y'Xc mbma 'f_l_fcd'OK WŁ'hfk U^ Wma 'dcbUX 90 a]bi h'NU'YWbU'XUk_U'nU'Y mċx-Xa UgmWUdUWYbH" DfnYX'dcXU]Ya 'Y_i 'CV]hcl UI]a UV'G: @ dUWYbhċa 'a c bU'dcXU 'Y_i 'a U^ W' bU'W'i 'nUdcV]Y Yb]Y 'k mgh dck Ub]ji 'fYU_WY'U'Yf[]WbmlW 'i V]W ograniczenie.

K] W^]bZfa UW]c'gdċgħċi 'Y'ghiegħġek Ub]U'Y_i 'CV]hcl UI]a UV'G: @nbU'Xi 'Y'g] 'k 'I 'chM'X'U'dUWYbH' i V' udzieli jid lekarz lub farmaceuta.

>U_ Xn]UW 'Y_ 'CV]hcl UI]a UV'G: @

Dck U bY'Xn]UWb]U'b]Ydc XUbY'k [`]U'k nk cħubY'g] dfnYn'hċċi_għib 'k nk cfncb dfnYn'VU_hYf]Y' k [`]U'CV]hcl UI]a UV'Ygh'dfnYW]k WU'ya 'a cbc_ċċbU'bma ż'Wm] fcXnU'Ya 'V]Uq_U'nUdfc'Y_hċċi Ub]Y[c'k '

HU_]`gdcgCV`z`VmXcU`WU`g]`Xc`g_WXb]_U`hc_gmbmk`[`L`U`nkUbY[c`Ubhmf`YbYa`cWfcbbma`k`[`L`U`z`_hEfmi a c`]k]U`Xcgh d`hc_gmb]Y`Xc`_ca`DFY_``DcdfnYn`XcU`WYb]Y`Xc`Ubhmf`Ybi`cWfcbbY[c`k`[`L`U`Y`_ma za zaXUb]Y`nU\Ua ck U`Xcgh d`hc_gmbmXc`_ca`DFY_`cf[Ub]na i`z`hma`gUa ma`c[fUb]WU^W`k`mgh dck Ub]Y`cV`Uk`DF`i`V`]W`nUdcV]Y`Yb]Y"

?cfnm W`nY`ghcgck Ub]U`Y_i`CV]`hcl UI]a UV`G: @`k m_UhUbY`k`VUXUb]UW

CV]`hcl UI]a UV`G: @`i nbU`Y`g]`nU`g_i`hWbmik`YWYb]i`k`[`L`U`wziewnego w oparciu o badania na nk]Yfn`HUVK`K`hfnYVW`VUXUb]UW`n`i`Xn]Udya`nU_U`cbmW`nk]Yfn`h`k`g_U`b]`dfnY`mWU`k`U`U`mig`k`granicach 30-60% po zastosowaniu leku Obiltoxaximab SFL, natomiast w przypadku placebo (leczenie dcncfck UbY`h`gUa`Y`k`g_U`b]`k`mb]cgum\$-*`"K`VUXUb]i`n`i`Xn]Udya`nU_U`cbmW`nk]Yfn`h`_hEfma`dcXUbc`Y`_i`V`d`UWVc`dfnYX`dc`Uk]Yb]Ya`g]`cV`Uk`DF`z`k`g_U`b]`dfnY`mWU`k`mbcg]ucX`\$`i`Xc`%\$`i`po zastosowaniu leku Obiltoxaximab SFL – nU`Y`bY`h`Vmic`hY[c`z`U`k`WY`b]Y`pk]Yfn`H`dcXXUbc`leWYb]i`"K`dfnmdUX_i`nk]Yfn`h`_hEfma`dcXUbc`d`UWVc`k`gdC`Wmnb]_h`b`k`mb]cgus`i`"

Fmm_m_c`nk`]`nUbY`nY`ghcgck Ub]Ya`Y_i`CV]`hcl UI]a UV`G: @

B`U`W`ghgnY`Xn]UdUb]U`b]Ydc`XUbY`nk`nUbY`nY`ghcgck Ub]Ya`Y_i`CV]`hcl UI]a UV`G: @`fa`c`W`k`mgh`d`u`1`na`10`pacjent`to`"VCE`[`uk`m`_k`X`fgk`XnYb]Y`z`dc_fnnk`U`fgk`Xn`W`k`mgnd_U`z`k`mgnd_U`z`_UgnY`z`VCE`k`a`]Y`gW`k`Y`k`i`]nUk`fchm`[`uk`m`]

DY`bmk`m_Uh`Xn]UdU`b]Ydc`XUbmW`c`fUb]WY`nk`nUbmW`bY`ghcgck Ub]Ya`Y_i`CV]`hcl UI]a UV`G: @`nbU`X`i`Y`g]`k`i`chW`X`U`d`UWVb]U`"

Podstawy dopuszczenia do obrotu leku Obiltoxaximab SFL w UE

K`[`L`k`n]Yk`bm`Ygh`W`fcfV`nU`fU`U`W`mW`z`dfck`UXn`W`Xc`a`]YfW`k`\$`i`dfnmdUX`DF`"A`ja`c`Y`bUh`fU`bY`c`[`b]g_U`k`mgh`di`^`VUF`Xc`fnUX`c`z`nU`U`Yb]U`a`c`dc`Uk`dfnmdUX`ck`c`k`UV`cfU`hcf]UW`nU`a`i`^`W`W`g`VUXUb]Ya`VU`hYf]z`bUhca`l`ugh`k`[`L`a`c`Y`Vm`k`m`cfnmg`hmk`Ubmk`U`H`U`UW`hY`ff`fm`ghm`W`bm`W`"NY`k`n`[`Xi`bU`b]g`W`W`dfnmdUX`DF`cfU`b`U`h`Y`z`W`ck`Y`nU`U`Yb]Y`cgCV`Y`gh`nV`mh`b]Y`VY`nd]Y`W`bY`z`b]Y`Y`gh`a`c`]k`Y`dfnY`dfck`UXnYb]Y`VUXU`Y`DF`b`U`i`Xn]UW`"6UXUb]U`ba`nk]Yfn`HUVK`k`m`_Un`U`m`Y`Y`Y`gh`g`i`hW`bmik`YW`Yb]i`k`[`L`U`]`nUdcV]Y`[Ub]i`a`]YfW`]`cW`Y`i`Y`g`z`Y`Y`CV]`hcl`UI`ja`UV`G: @`V`Xn]Y`Xn]UdU`k`H`U`_j`gUa`gdgcV`i`i`Xn]`>`Y`]W`cXn]`c`VY`nd]Y`W`Y`gh`c`h`Xn]UdUb]U`b]Ydc`XUbY`nk`nUbY`n`Y`_i`CV]`hcl`UI`ja`UV`G: @`i`nXfck`mW`cgCV`a`U`^`nUnk`mWU`^`W`cXbm`i`V`i`a`]Uf_ck`UbmidfnYV]Y`"8`U`hY`5`[`YbWU`i`nbU`U`z`Y`_cfnm`W`d`m`b`W`nY`ghcgck`Ub]U`Y`i`CV]`hcl`UI`ja`UV`dfnYk`m`gnU`fmm_m_c`a`c`Y`cb`Vm`Xcdi`gnW`cbm`Xc`ghcgck`Ub`k`i`9"

Obiltoxaximab SFL dopuszczony do obrotu `k`k`m`h_ck`mW`c`c`]W`bc`W`UW`"K`mb`_U`h`n`Z`U`h`z`Y`b]Y`a`c`bU`Vmic`i`mng_U`d`UW`mW`]bZ`fa`UW`c`Y`_i`CV]`hcl`UI`ja`UV`G: @`Xcdi`gnW`cbm`Xc`c`Vfch`k`k`m`h_ck`mW`c`c`]W`bc`W`UW`z`Z`fa`U`k`dfck`UXnU`W`Y`_i`CV]`hcl`UI`ja`UV`G: @`Xc`c`Vfch`Xc`ghfW`m`XU`gnY`XUbY`XchmW`W`a`Yh`X`dca`]Uf`k`W`d`b]U`b]U`z`a`c`Xm`_UW`]`i`gi`k`Ub]U`Y`_i`n`cf[Ub]na`i`k`VUXUb]UW`UV`cfU`h`f`m`b`m`W`"Dcb`U`h`b`U`Y`m`dfnY`X`gh`Uk`XUbY`XchmW`W`g`i`hW`bm`W`Y`VY`nd]Y`W`Y`gh`k`U`Y`_i`b`U`k`m`UXY`_Yk`Y`b`h`U`b`Y`[c`dc`Uk]Yb]U`g`i`c`[`b]g_U`k`[`L`U`

Jakich informacji jeszcze brakuje na temat leku Obiltoxaximab SFL

K`nk`]`h`i`n`hma`z`Y`Y`_i`CV]`hcl`UI`ja`UV`G: @`Xcdi`gnW`cbm`Xc`c`Vfch`k`k`m`h_ck`mW`c`c`]W`bc`W`UW`z`Z`fa`U`k`dfck`UXnU`W`Y`_i`CV]`hcl`UI`ja`UV`G: @`Xc`c`Vfch`Xc`ghfW`m`XU`gnY`XUbY`XchmW`W`a`Yh`X`dca`]Uf`k`W`d`b]U`b]U`z`a`c`Xm`_UW`]`i`gi`k`Ub]U`Y`_i`n`cf[Ub]na`i`k`VUXUb]UW`UV`cfU`h`f`m`b`m`W`"Dcb`U`h`b`U`Y`m`dfnY`X`gh`Uk`XUbY`XchmW`W`g`i`hW`bm`W`Y`VY`nd]Y`W`Y`gh`k`U`Y`_i`b`U`k`m`UXY`_Yk`Y`b`h`U`b`Y`[c`dc`Uk]Yb]U`g`i`c`[`b]g_U`k`[`L`U`

fcX_]dcXY'a ck UbY'k 'W'i 'nUdYk b]Yb]U'VYnd]YVbY[c]'g_i hVbY[c]'sosowania leku Obiltoxaximab SFL

W celu zapewnienia bezpiecznego i skutecznego stosowania leku Obiltoxaximab SFL w Charakterystyce Produktu Leczniczego i w Ulotce dla pacjenta zawafhc 'nU'YVb]U''] fcX_]cghfc bc W'dfnYnbUWcbY'XU personelu medycznego i pacjentu.

HU_ 'U_k 'dfnmdUX_i 'k gnmgh_]Mk 'Y_DK z XUbY'c'ghcgck Ub]i 'Y_i 'CV]hcl UI]a UV'G: @g 'gHUY' a cb]hcfck UbY''N[dgnUbY'Xn]UdUb]U'b]Ydc XUbY'Y_i 'CV]hcl UI]a UV'G: @g 'gHfUbb]Y'oceniane i dcXY'a ck UbY'g 'k gnmgh_]Y'Vmbbc W'_cb]YVbY'Xc 'cW'fcbmidUVYbhDK "

=bbY']bZcfa UWY'XchmW W'Y_i 'CV]hcl UI]a UV'G: @

8U'gnY']bZcfa UWY'bU'hYa Uh'Y_i 'CV]hcl UI]a UV'G: @nbU'Xi ^ 'g] 'bU'ghfcb]Y']bhfbYhck Y^5[YbW]dcX' UXfYgYa . ema.europa.eu/medicines/human/EPAR/Obiltoxaximab-SFL

Produkt leczniczy bez wafnego pozwolenia na dopuszczenie do obrotu