



I Quaderni di Orphanet

Collezione Farmaci Orfani

Gennaio 2018

Elenchi dei prodotti medicinali per le malattie rare in Europa*

**Autorizzazione all'immissione in commercio della Comunità Europea nell'ambito della procedura*

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PARTE 1:

Elenco dei prodotti medicinali orfani in Europa con designazione orfana europea e autorizzazione all'immissione in commercio europea*

Indice

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Metodologia

Questa parte del documento fornisce l'elenco di tutti i prodotti medicinali che hanno ottenuto l'autorizzazione all'immissione in commercio (AIC) europea nella data indicata nel documento. Questi prodotti medicinali possono essere accessibili in alcuni paesi europei, ma non necessariamente in tutti. In effetti, l'accessibilità di un determinato prodotto medicinale orfano in un determinato paese dipende dalla strategia intrapresa dal laboratorio e dalle decisioni prese dalle autorità sanitarie nazionali in materia di rimborsi.

I prodotti medicinali orfani in Europa sono prodotti medicinali che hanno ottenuto una designazione orfana europea (secondo il Regolamento (CE) N° 141/2000) e che, in seguito, hanno ottenuto un'autorizzazione all'immissione in commercio europea ed,

eventualmente, una valutazione positiva sui benefici rilevanti.

Questo elenco dei prodotti medicinali orfani in Europa con designazione orfana e autorizzazione all'immissione in commercio europea è il risultato del raffronto incrociato dell'elenco dei prodotti medicinali che hanno ottenuto una designazione orfana (<http://ec.europa.eu/health/documents/community-register/html/alforphreg.htm>) con l'elenco dei prodotti medicinali che hanno ottenuto un'autorizzazione all'immissione in commercio (<http://ec.europa.eu/health/documents/community-register/html/alfregister.htm>). Entrambi gli elenchi sono disponibili sul sito della Direzione generale Salute e Sicurezza Alimentare (DG SANTE) della Commissione Europea.

*Autorizzazione all'immissione in commercio dell'Unione Europea mediante procedura centralizzata

Una prima classificazione per denominazione commerciale fornisce il nome della sostanza attiva, l'indicazione relativa all'autorizzazione all'immissione in commercio, la data e il titolare dell'AIC.

Questa classificazione è seguita da due allegati che comprendono:


- una tabella con i prodotti medicinali orfani ritirati dal Registro Comunitario dei Prodotti Medicinali Orfani (vedi Allegato 1 -"Prodotti medicinali orfani rimossi o ritirati dal Registro Comunitario Europeo dei Prodotti Medicinali Orfani"; le indicazioni dei farmaci in questione sono esplicitate nel dettaglio nella Parte II, "Elenco dei prodotti medicinali per le malattie rare in Europa con autorizzazione all'immissione in commercio senza designazione orfana europea");

- una tabella con i prodotti medicinali orfani ritirati dal commercio nell'Unione Europea (vedi Allegato 2- Prodotti medicinali ritirati dal commercio nell'Unione Europea). Maggiori informazioni disponibili su www.ema.europa.eu

Tre elenchi aggiuntivi propongono altre classificazioni per:

- Data dell'AIC in ordine decrescente;
- Categoria ATC;
- Titolare dell'AIC.

In ciascun elenco, le denominazione commerciali sono disposte in ordine alfabetico.

Ulteriori informazioni sono disponibili nella sezione "Farmaci orfani" del sito web di Orphanet, www.orpha.net, o sul sito web dell'EMA (Agenzia Europea per i Medicinali) <http://www.ema.europa.eu>. Gli elenchi forniti dall'EMA non comprendono solo i prodotti medicinali orfani ma tutti i prodotti medicinali autorizzati all'immissione in commercio. I prodotti medicinali orfani con designazione orfana europea sono contrassegnati da un logo 

Informazioni ufficiali aggiornate sui prodotti medicinali orfani sono disponibili nel Registro comunitario dei prodotti medicinali orfani per uso umano: <http://ec.europa.eu/health/documents/community-register/html/alforphreg.htm>

Classificazione per denominazione

TRADENAME	ACTIVE SUBSTANCE	MARKETING AUTHORISATION INDICATION	MARKETING AUTHORISATION DATE (DD/MM/YYYY)	MARKETING AUTHORISATION HOLDER
ADCETRIS	brentuximab vedotin	<p>Treatment of adult patients with relapsed or refractory CD30+ Hodgkin lymphoma (HL):</p> <ul style="list-style-type: none"> -following autologous stem cell transplant (ASCT) or -following at least two prior therapies when ASCT or multi-agent chemotherapy is not a treatment option. <p>Treatment of adult patients with CD30+ HL at increased risk of relapse or progression following ASCT</p> <p>Treatment of adult patients with relapsed or refractory systemic anaplastic large cell lymphoma (sALCL).</p> <p>Treatment of adult patients with CD30+ cutaneous T-cell lymphoma (CTCL) after at least 1 prior systemic therapy.</p>	25/10/2012	Takeda Pharma A/S
ADEMPAS	riociguat	<p>Treatment of adult patients with WHO Functional Class (FC) II to III with inoperable Chronic thromboembolic pulmonary hypertension (CTEPH), persistent or recurrent CTEPH after surgical treatment, to improve exercise capacity.</p> <p>As monotherapy or in combination with endothelin receptor antagonists, for the treatment of adult patients with pulmonary arterial hypertension (PAH) with WHO Functional Class (FC) II to III to improve exercise capacity.</p> <p>Efficacy has been shown in a PAH population including etiologies of idiopathic or heritable PAH or PAH associated with connective tissue disease.</p>	27/03/2014	Bayer Pharma AG
ALPROLIX	eftrenonacog alfa	Treatment and prophylaxis of bleeding in patients with haemophilia B (congenital factor IX deficiency). ALPROLIX can be used for all age groups.	12/05/2016	Biogen Idec Ltd
ARZERRA	ofatumumab	<p>In combination with chlorambucil or bendamustine, for the treatment of patients with chronic lymphocytic leukaemia (CLL) who have not received prior therapy and who are not eligible for fludarabine-based therapy.</p> <p>In combination with fludarabine and cyclophosphamide for the treatment of adult patients with relapsed CLL.</p> <p>Treatment of CLL in patients who are refractory to fludarabine and alemtuzumab.</p>	19/04/2010	Novartis Europharm Ltd
BAVENCIO	avelumab	As monotherapy for the treatment of adult patients with metastatic Merkel cell carcinoma (MCC) .	18/09/2017	Merck Serono Europe Limited

TRADENAME	ACTIVE SUBSTANCE	MARKETING AUTHORISATION INDICATION	MARKETING AUTHORISATION DATE (DD/MM/YYYY)	MARKETING AUTHORISATION HOLDER
BESPONSA	inotuzumab ozogamicin	As monotherapy for the treatment of adults with relapsed or refractory CD22- positive B cell precursor acute lymphoblastic leukaemia (ALL) . Adult patients with Philadelphia chromosome positive (Ph+) relapsed or refractory B cell precursor ALL should have failed treatment with at least 1 tyrosine kinase inhibitor (TKI).	29/06/2017	Pfizer Limited
BLINCYTO	blinatumomab	Treatment of adults with Philadelphia chromosome negative relapsed or refractory B-precursor acute lymphoblastic leukaemia (ALL) .	23/11/2015	Amgen Europe B.V.
BOSULIF	bosutinib	Treatment of adult patients with chronic phase (CP), accelerated phase (AP), and blast phase (BP) Philadelphia chromosome positive chronic myelogenous leukaemia (Ph+ CML) previously treated with one or more tyrosine kinase inhibitor(s) and for whom imatinib, nilotinib and dasatinib are not considered appropriate treatment options.	27/03/2013	Pfizer Ltd
BRINEURA	cerliponase alfa	Treatment of neuronal ceroid lipofuscinosis type 2 (CLN2) disease, also known as tripeptidyl peptidase 1 (TPP1) deficiency.	30/05/2017	BioMarin International Limited
BRONCHITOL	mannitol	Treatment of cystic fibrosis (CF) in adults aged 18 years and above as an add-on therapy to best standard of care.	13/04/2012	Pharmaxis Pharmaceuticals Ltd
CARBAGLU	carglumic acid	Treatment of hyperammonaemia due to - isovaleric acidaemia , - methymalonic acidaemia , - propionic acidaemia .	01/06/2011	Orphan Europe S.a.r.l.
CAYSTON	aztreonam	Suppressive therapy of chronic pulmonary infections due to <i>Pseudomonas aeruginosa</i> in patients with cystic fibrosis (CF) aged 6 years and older.	21/09/2009	Gilead Sciences International Ltd
CEPLENE	histamine dihydrochloride	Maintenance therapy for adult patients with acute myeloid leukaemia in first remission concomitantly treated with interleukin-2 (IL-2). The efficacy of Ceplene has not been fully demonstrated in patients older than age 60.	07/10/2008	Noventia Pharma Srl
CERDELGA	eliglustat	Long-term treatment of adult patients with Gaucher disease type 1 (GD1) , who are CYP2D6 poor metabolisers (PMs), intermediate metabolisers (IMs) or extensive metabolisers (EMs)	19/01/2015	Genzyme Europe B.V.
CHENODEOXYCHOLIC ACID LEADIANT (previously CHENODEOXYCHOLIC ACID SIGMA-TAU)	chenodeoxycholic acid	Treatment of inborn errors of primary bile acid synthesis due to sterol 27 hydroxylase deficiency (presenting as cerebrotendinous xanthomatosis (CTX)) in infants, children and adolescents aged 1 month to 18 years and adults.	10/04/2017	Leadiant GmbH
COAGADEX	human coagulation factor X	Treatment and prophylaxis of bleeding episodes and for perioperative management in patients with hereditary factor X deficiency .	16/03/2016	Bio Products Laboratory Ltd

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COMETRIQ	cabozantinib	Treatment of adult patients with progressive, unresectable locally advanced or metastatic medullary thyroid carcinoma . For patients in whom Rearranged during Transfection (RET) mutation status is not known or is negative, a possible lower benefit should be taken into account before individual treatment decision.	21/03/2014	TMC PharmaServices Ltd.
CRESEMBA	isavuconazole	In adults for the treatment of: - invasive aspergillosis - mucormycosis in patients for whom amphotericin B is inappropriate	15/10/2015	Basilea Medical Ltd
CYSTADROPS	mercaptamine hydrochloride	Treatment of corneal cystine crystal deposits in adults and children from 2 years of age with cystinosis .	19/01/2017	Orphan Europe S.A.R.L.
DACOGEN	decitabine	Treatment of adult patients aged 65 years and above with newly diagnosed de novo or secondary acute myeloid leukaemia (AML) , according to the World Health Organization (WHO) classification, who are not candidates for standard induction chemotherapy.	20/09/2012	Janssen-Cilag International N.V.
DARZALEX	daratumumab	As monotherapy for the treatment of adult patients with relapsed and refractory multiple myeloma , whose prior therapy included a proteasome inhibitor and an immunomodulatory agent and who have demonstrated disease progression on the last therapy. In combination with lenalidomide and dexamethasone, or bortezomib and dexamethasone, for the treatment of adult patients with multiple myeloma who have received at least one prior therapy.	20/05/2016	Janssen-Cilag International N.V.
DEFITELIO	defibrotide	Treatment of severe hepatic veno-occlusive disease (VOD) also known as sinusoidal obstructive syndrome (SOS) in haematopoietic stem-cell transplantation (HSCT) therapy. It is indicated in adults and in adolescents, children and infants over 1 month of age.	18/10/2013	Gentium S.p.A.
DELTYBA	delamanib	Used as part of an appropriate combination regimen for pulmonary multi-drug resistant tuberculosis (MDR-TB) in adult patients when an effective treatment regimen cannot otherwise be composed for reasons of resistance or tolerability. Consideration should be given to official guidance on the appropriate use of antibacterial agents.	28/04/2014	Otsuka Novel Products GmbH

TRADENAME	ACTIVE SUBSTANCE	MARKETING AUTHORISATION INDICATION	MARKETING AUTHORISATION DATE (DD/MM/YYYY)	MARKETING AUTHORISATION HOLDER
DINUTUXIMAB BETA APEIRON	dinutuximab beta	Treatment of high-risk neuroblastoma in patients aged 12 months and above, who have previously received induction chemotherapy and achieved at least a partial response, followed by myeloablative therapy and stem cell transplantation, as well as patients with history of relapsed or refractory neuroblastoma, with or without residual disease. Prior to the treatment of relapsed neuroblastoma, any actively progressing disease should be stabilised by other suitable measures. In patients with a history of relapsed/refractory disease and in patients who have not achieved a complete response after first line therapy, Dinutuximab beta Apeiron should be combined with interleukin-2 (IL-2).	08/05/2017	Apeiron Biologics AG
ESBRIET	pirfenidone	In adults for the treatment of mild to moderate Idiopathic Pulmonary Fibrosis (IPF) .	28/02/2011	Roche Registration Ltd
FARYDAK	panobinostat lactate anhydrous	In combination with bortezomib and dexamethasone, for the treatment of adult patients with relapsed and/or refractory multiple myeloma who have received at least two prior regimens including bortezomib and an immunomodulatory agent.	28/08/2015	Novartis Europharm Ltd
FIRAZYR	icatibant acetate	Symptomatic treatment of acute attacks of hereditary angioedema (HAE) in adults, adolescents and children aged 2 years and older, with C1-esterase-inhibitor deficiency.	11/07/2008	Shire Orphan Therapies GmbH
FIRDAPSE (previously ZENAS)	amifampridine	Symptomatic treatment of Lambert-Eaton myasthenic syndrome (LEMS) in adults.	23/12/2009	BioMarin Europe Ltd
GALAFOLD	migalastat	Long-term treatment of adults and adolescents aged 16 years and older with a confirmed diagnosis of Fabry disease (α-galactosidase A deficiency) and who have an amenable mutation.	26/05/2016	Amicus Therapeutics UK Ltd
GAZYVARO	obinutuzumab	In combination with chlorambucil, treatment of adult patients with previously untreated chronic lymphocytic leukaemia (CLL) and with comorbidities making them unsuitable for full-dose fludarabine based therapy. In combination with chemotherapy, followed by Gazyvaro maintenance therapy in patients achieving a response, is indicated for the treatment of patients with previously untreated advanced follicular lymphoma . In combination with bendamustine followed by Gazyvaro maintenance is indicated for the treatment of patients with follicular lymphoma (FL) who did not respond or who progressed during or up to 6 months after treatment with rituximab or a rituximab-containing regimen.	23/07/2014	Roche Registration Ltd

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GLYBERA	alipogene tiparvovec	For adult patients diagnosed with familial lipoprotein lipase deficiency (LPLD) and suffering from severe or multiple pancreatitis attacks despite dietary fat restrictions. The diagnosis of LPLD has to be confirmed by genetic testing. The indication is restricted to patients with detectable levels of LPL protein.	25/10/2012	uniQure biopharma B.V.
GRANUPAS (previously PARA-AMINOSALICYLIC ACID LUCANE)	para-aminosalicylic acid	Indicated for use as part of an appropriate combination regimen for multi-drug resistant tuberculosis in adults and paediatric patients from 28 days of age and older when an effective treatment regimen cannot otherwise be composed for reasons of resistance or tolerability. Consideration should be given to official guidance on the appropriate use of antibacterial agents.	07/04/2014	Lucane Pharma
HETLIOZ	tasimelteon	Treatment of Non-24-Hour Sleep-Wake Disorder (Non-24) in totally blind adults.	03/07/2015	Vanda Pharmaceuticals Ltd
HOLOCLAR	ex vivo expanded autologous human corneal epithelial cells containing stem cells	Treatment of adult patients with moderate to severe limbal stem cell deficiency (defined by the presence of superficial corneal neovascularisation in at least two corneal quadrants, with central corneal involvement, and severely impaired visual acuity), unilateral or bilateral, due to physical or chemical ocular burns. A minimum of 1 - 2 mm ² of undamaged limbus is required for biopsy.	17/02/2015	Chiesi Farmaceutici SpA
ICLUSIG	ponatinib	Indicated in adult patients with chronic phase, accelerated phase, or blast phase chronic myeloid leukaemia (CML) who are resistant to dasatinib or nilotinib; who are intolerant to dasatinib or nilotinib and for whom subsequent treatment with imatinib is not clinically appropriate; or who have the T315I mutation. Indicated in adult patients with Philadelphia chromosome positive acute lymphoblastic leukaemia (Ph+ ALL) who are resistant to dasatinib; who are intolerant to dasatinib and for whom subsequent treatment with imatinib is not clinically appropriate; or who have the T315I mutation.	01/07/2013	ARIAD Pharma Ltd
IDELVION	albutrepenonacog alfa	Treatment and prophylaxis of bleeding in patients with haemophilia B (congenital factor IX deficiency) . IDELVION can be used for all age groups.	11/05/2016	CSL Behring GmbH

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IMBRUVICA	ibrutinib	As a single agent for the treatment of adult patients with relapsed or refractory mantle cell lymphoma (MCL) . As a single agent for the treatment of adult patients with previously untreated chronic lymphocytic leukaemia (CLL) . As a single agent or in combination with bendamustine and rituximab (BR) for the treatment of adult patients with CLL who have received at least one prior therapy. As a single agent for the treatment of adult patients with Waldenström's macroglobulinaemia (WM) who have received at least one prior therapy, or in first line treatment for patients unsuitable for chemo-immunotherapy.	21/10/2014	Janssen-Cilag International N.V.
IMNOVID (previously POMALIDOMI DE CELGENE)	pomalidomide	In combination with dexamethasone, in the treatment of adult patients with relapsed and refractory multiple myeloma who have received at least two prior treatment regimens, including both lenalidomide and bortezomib, and have demonstrated disease progression on the last therapy.	05/08/2013	Celgene Europe Ltd
INOVELON	rufinamide	Adjunctive therapy in the treatment of seizures associated with Lennox-Gastaut syndrome in patients aged 4 years and older.	16/01/2007	Eisai Ltd
JORVEZA	budesonide	Treatment of eosinophilic esophagitis (EoE) in adults (older than 18 years of age).	08/01/2018	Dr. Falk Pharma GmbH
KALYDECO	ivacaftor	Treatment of patients with cystic fibrosis (CF) aged 6 years and older and weighing 25kg or more who have one of the following gating (class III) mutations in the CFTR gene: <i>G551D, G1244E, G1349D, G178R, G551S, S1251N, S1255P, S549N</i> or <i>S549R</i> . Treatment of patients with cystic fibrosis (CF) aged 18 years and older who have an <i>R117H</i> mutation in the CFTR gene	23/07/2012	Vertex Pharmaceuticals (Europe) Ltd
KANUMA	sebelipase alfa	Long-term enzyme replacement therapy (ERT) in patients of all ages with lysosomal acid lipase (LAL) deficiency	28/08/2015	Synageva BioPharma Ltd
KETOCONAZOLE HRA	ketoconazole	Treatment of endogenous Cushing's syndrome in adults and adolescents above the age of 12 years.	19/11/2014	Laboratoire HRA Pharma
KOLBAM (previously CHOLIC ACID FGK)	cholic acid	Treatment of inborn errors in primary bile acid synthesis due to sterol 27-hydroxylase (presenting as cerebrotendinous xanthomatosis, CTX) deficiency, 2- (or α-) methylacyl-CoA racemase (AMACR) deficiency or cholesterol 7α-hydroxylase (CYP7A1) deficiency in infants, children and adolescents aged 1 month to 18 years and adults.	08/04/2014	Retrophin Europe Ltd

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KUVAN	sapropterin dihydrochloride	Treatment of hyperphenylalaninaemia (HPA) in adult and paediatric patients of all ages with phenylketonuria (PKU) who have been shown to be responsive to such treatment. Treatment of hyperphenylalaninaemia (HPA) in adult and paediatric patients with tetrahydrobiopterin (BH4) deficiency who have been shown to be responsive to such treatment.	02/12/2008	Merck Serono Europe Ltd
KYPROLIS	carfilzomib	In combination with either lenalidomide and dexamethasone or dexamethasone alone is indicated for the treatment of adult patients with multiple myeloma who have received at least one prior therapy.	19/11/2015	Amgen Europe B.V.
LARTRUVO	olaratumab	In combination with doxorubicin for the treatment of adult patients with advanced soft tissue sarcoma who are not amenable to curative treatment with surgery or radiotherapy and who have not been previously treated with doxorubicin.	09/11/2016	Eli Lilly Nederland B.V.
LEDAGA	chlormethine	Topical treatment of mycosis fungoides-type cutaneous T-cell lymphoma (MF-type CTCL) in adult patients.	03/03/2017	Actelion Registration Ltd.
LENVIMA	lenvatinib	Treatment of adult patients with progressive, locally advanced or metastatic, differentiated (papillary/follicular/Hürthle cell) thyroid carcinoma (DTC) refractory to radioactive iodine (RAI).	28/05/2015	Eisai Ltd
LUTATHERA	lutetium (177Lu) oxodotreotide	Treatment of unresectable or metastatic, progressive, well differentiated (G1 and G2), somatostatin receptor positive gastroenteropancreatic neuroendocrine tumours (GEP NETs) in adults.	26/09/2017	Advanced Accelerator Applications
LYNPARZA	olaparib	Monotherapy for the maintenance treatment of adult patients with platinum-sensitive relapsed BRCA-mutated (germline and/or somatic) high grade serous epithelial ovarian, fallopian tube, or primary peritoneal cancer who are in response (complete response or partial response) to platinum-based chemotherapy.	16/12/2014	AstraZeneca AB
MEPACT	mifamurtide	In children, adolescents and young adults for the treatment of high-grade resectable non-metastatic osteosarcoma after macroscopically complete surgical resection. It is used in combination with post-operative multi-agent chemotherapy. Safety and efficacy have been assessed in studies of patients 2 to 30 years of age at initial diagnosis.	06/03/2009	Takeda France SAS

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MOZOBIL	plerixafor	In combination with granulocyte-colony stimulating factor G-CSF to enhance mobilisation of haematopoietic stem cells to the peripheral blood for collection and subsequent autologous transplantation in patients with lymphoma and multiple myeloma whose cells mobilise poorly.	31/07/2009	Genzyme Europe B.V.
NATPAR	parathyroid hormone	Indicated as adjunctive treatment of adult patients with chronic hypoparathyroidism who cannot be adequately controlled with standard therapy alone.	24/04/2017	Shire Pharmaceuticals Ireland Ltd
NEXAVAR	sorafenib tosylate	Treatment of hepatocellular carcinoma . Treatment of patients with progressive, locally advanced or metastatic, differentiated (papillary/follicular/Hürthle cell) thyroid carcinoma , refractory to radioactive iodine.	19/07/2006	Bayer Pharma AG
NEXOBRID	concentrate of proteolytic enzymes enriched in bromelain	Removal of eschar in adults with deep partial- and full-thickness thermal burns .	18/12/2012	Mediound Germany GmbH
NINLARO	ixazomib	In combination with lenalidomide and dexamethasone is indicated for the treatment of adult patients with multiple myeloma who have received at least one prior therapy.	21/11/2016	Takeda Pharma A/S
NPLATE	romiplostim	Indicated for adult chronic immune (idiopathic) thrombocytopenic purpura (ITP) patients who are refractory to other treatments (e.g. corticosteroids, immunoglobulins)	04/02/2009	Amgen Europe B.V.
OCALIVA	obeticholic acid	Treatment of primary biliary cholangitis (also known as primary biliary cirrhosis) in combination with ursodeoxycholic acid (UDCA) in adults with an inadequate response to UDCA or as monotherapy in adults unable to tolerate UDCA.	12/12/2016	Intercept Pharma Ltd
OFEV	nintedanib	Treatment in adults of Idiopathic Pulmonary Fibrosis (IPF) .	15/01/2015	Boehringer Ingelheim International GmbH
ONIVYDE	irinotecan hydrochloride trihydrate	Treatment of metastatic adenocarcinoma of the pancreas , in combination with 5-fluorouracil (5-FU) and leucovorin (LV), in adult patients who have progressed following gemcitabine based therapy.	14/10/2016	Baxalta Innovations GmbH
OPSUMIT	macitentan	Used as monotherapy or in combination, for the long-term treatment of pulmonary arterial hypertension (PAH) in adult patients of WHO Functional Class (FC) II to III. Efficacy has been shown in a PAH population including idiopathic and heritable PAH, PAH associated with connective tissue disorders, and PAH associated with corrected simple congenital heart disease.	20/12/2013	Actelion Registration Ltd



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ORPHACOL	cholic acid	Treatment of inborn errors in primary bile acid synthesis due to 3beta-hydroxy-delta5-C27- steroid oxidoreductase deficiency or delta4-3-oxosteroid-5beta-reductase deficiency in infants, children and adolescents aged 1 month to 18 years and adults.	12/09/2013	Laboratoires CTRS
OXERVATE	cenegermin	Treatment of moderate (persistent epithelial defect) or severe (corneal ulcer) neurotrophic keratitis in adults.	06/07/2017	Dompe farmaceutici s.p.a.
PEYONA (previously NYMUSA)	caffeine citrate	Treatment of primary apnea of premature newborns.	02/07/2009	Chiesi Farmaceutici SpA
PLENADREN	hydrocortisone	Treatment of adrenal insufficiency in adults.	03/11/2011	ViroPharmaSPRL
PREVYMIS	letermovir	Prophylaxis of cytomegalovirus (CMV) reactivation and disease in adult CMV-seropositive recipients [R+] of an allogeneic haematopoietic stem cell transplant (HSCT). Consideration should be given to official guidance on the appropriate use of antiviral agents.	08/01/2018	Merck Sharp & Dohme Limited
PROCYSBI	mercaptopamine	Treatment of proven nephropathic cystinosis . Cysteamine reduces cystine accumulation in some cells (e.g. leukocytes, muscle and liver cells) of nephropathic cystinosis patients and, when treatment is started early, it delays the development of renal failure.	06/09/2013	Raptor Pharmaceuticals Europe BV
RAVICTI	glycerol phenylbutyrate	Indicated for use as adjunctive therapy for chronic management of adult and paediatric patients ≥ 2 months of age with urea cycle disorders (UCDs) including: deficiencies of carbamoyl phosphate-synthase-I (CPS) -ornithine carbamoyltransferase (OTC) - argininosuccinate synthetase (ASS), - argininosuccinate lyase (ASL) - arginase I (ARG) - ornithine translocase deficiency hyperornithinaemia -hyperammonaemia homocitrullinuria syndrome (HHH) Who cannot be managed by dietary protein restriction and/or amino acid supplementation alone. RAVICTI must be used with dietary protein restriction and, in some cases, dietary supplements (e.g., essential amino acids, arginine, citrulline, protein-free calorie supplements).	27/11/2015	Horizon Therapeutics Ltd
RAXONE	idebenone	Treatment of visual impairment in adolescent and adult patients with Leber's Hereditary Optic Neuropathy (LHON) .	08/09/2015	Santhera Pharmaceuticals (Deutschland) GmbH
REVESTIVE	teduglutide	Treatment of patients aged 1 year and above with Short Bowel Syndrome . Patients should be stable following a period of intestinal adaptation after surgery.	30/08/2012	NPS Pharma Holdings Ltd

TRADENAME	ACTIVE SUBSTANCE	MARKETING AUTHORISATION INDICATION	MARKETING AUTHORISATION DATE (DD/MM/YYYY)	MARKETING AUTHORISATION HOLDER
REVLIMID	lenalidomide	Treatment of patients with transfusion-dependent anaemia due to low-or intermediate-1-risk myelodysplastic syndromes associated with an isolated deletion 5q cytogenetic abnormality when other therapeutic options are insufficient or inadequate. Treatment of adult patients with relapsed or refractory mantle cell lymphoma .	13/06/2013	Celgene Europe Ltd
RYDAPT	midostaurin	In combination with standard daunorubicin and cytarabine induction and high dose cytarabine consolidation chemotherapy, and for patients in complete response followed by Rydapt single agent maintenance therapy, for adult patients with newly diagnosed acute myeloid leukaemia (AML) who are FLT3 mutation positive. As monotherapy for the treatment of adult patients with aggressive systemic mastocytosis (ASM) , systemic mastocytosis with associated haematological neoplasm (SM AHN) , or mast cell leukaemia (MCL) .	18/09/2017	Novartis Europharm Limited
SCENESSE	afamelanotide	Prevention of phototoxicity in adult patients with erythropoietic protoporphyria (EPP)	22/12/2014	Clinuvel UK Ltd
SIGNIFOR	pasireotide	Treatment of adult patients with Cushing's disease for whom surgery is not an option or for whom surgery has failed. Treatment of adult patients with acromegaly for whom surgery is not an option or has not been curative and who are inadequately controlled on treatment with another somatostatin analogue.	24/04/2012	Novartis Europharm Ltd
SIRTURO	bedaquiline	Used as part of an appropriate combination regimen for pulmonary multidrug-resistant tuberculosis (MDR-TB) in adult patients when an effective treatment regimen cannot otherwise be composed for reasons of resistance or tolerability. Consideration should be given to official guidance on the appropriate use of antibacterial agents.	05/03/2014	Janssen-Cilag International N.V.
SOLIRIS	eculizumab	Treatment of adults and children with : - Paroxysmal nocturnal haemoglobinuria (PNH) . Evidence of clinical benefit is demonstrated in patients with haemolysis with clinical symptom(s) indicative of high disease activity, regardless of transfusion history. - atypical haemolytic uraemic syndrome (aHUS) . - Refractory generalized myasthenia gravis (gMG) in patients who are anti-acetylcholine receptor (AChR) antibody-positive	20/06/2007	Alexion Europe SAS

TRADENAME	ACTIVE SUBSTANCE	MARKETING AUTHORISATION INDICATION	MARKETING AUTHORISATION DATE (DD/MM/YYYY)	MARKETING AUTHORISATION HOLDER
SOMAKIT TOC	edotreotide	After radiolabelling with gallium (⁶⁸ Ga) chloride solution, the solution of gallium (⁶⁸ Ga) edotreotide obtained is indicated for Positron Emission Tomography (PET) imaging of somatostatin receptor overexpression in adult patients with confirmed or suspected well-differentiated gastro-enteropancreatic neuroendocrine tumours (GEP-NET) for localizing primary tumours and their metastases.	08/12/2016	Advanced Accelerator Applications
SPINRAZA	nusinersen sodium	Treatment of 5q Spinal Muscular Atrophy .	30/05/2017	Biogen Idec Ltd
STRENSIQ	asfotase alfa	Long-term enzyme replacement therapy in patients with paediatric-onset hypophosphatasia to treat the bone manifestations of the disease.	28/08/2015	Alexion Europe SAS
STRIMVELIS	autologous CD34+ enriched cell fraction that contains CD34+ cells transduced with retroviral vector that encodes for the human adenosine deaminase (ADA) cDNA sequence from human haematopoietic stem/progenitor (CD34+) cells	Treatment of patients with severe combined immunodeficiency due to adenosine deaminase deficiency (ADA-SCID) , for whom no suitable human leukocyte antigen (HLA)-matched related stem cell donor is available.	26/05/2016	GlaxoSmithKline Trading Services Limited
SYLVANT	siltuximab	Treatment of adult patients with multicentric Castlemans disease (MCD) who are human immunodeficiency virus (HIV) negative and human herpesvirus-8 (HHV-8) negative.	22/05/2014	Janssen-Cilag International N.V.
TASIGNA	nilotinib	Treatment of adult and paediatric patients with newly diagnosed Philadelphia chromosome positive chronic myelogenous leukaemia (CML) in the chronic phase. Adult patients with chronic phase and accelerated phase Philadelphia chromosome positive CML with resistance or intolerance to prior therapy including imatinib. Efficacy data in patients with CML in blast crisis are not available. Paediatric patients with chronic phase Philadelphia chromosome positive CML with resistance or intolerance to prior therapy including imatinib.	19/11/2007	Novartis Europharm Ltd

TRADENAME	ACTIVE SUBSTANCE	MARKETING AUTHORISATION INDICATION	MARKETING AUTHORISATION DATE (DD/MM/YYYY)	MARKETING AUTHORISATION HOLDER
TEPADINA	thiotepa	In combination with other chemotherapy medicinal products: 1) with or without total body irradiation (TBI), as conditioning treatment prior to allogeneic or autologous haematopoietic progenitor cell transplantation (HPCT) in haematological diseases in adult and paediatric patients; 2) when high dose chemotherapy with HPCT support is appropriate for the treatment of solid tumours in adult and paediatric patients.	15/03/2010	Adienne S.r.l.
THALIDOMIDE CELGENE (previously THALIDOMIDE PHARMION)	thalidomide	In combination with melphalan and prednisone as first line treatment of patients with untreated multiple myeloma , aged ≥ 65 years or ineligible for high dose chemotherapy. Thalidomide Celgene is prescribed and dispensed according to the Thalidomide Celgene Pregnancy Prevention Programme	16/04/2008	Celgene Europe Ltd
TOBI PODHALER	tobramycin	Suppressive therapy of chronic pulmonary infection due to <i>Pseudomonas aeruginosa</i> in adults and children aged 6 years and older with cystic fibrosis . Consideration should be given to official guidance on the appropriate use of antibacterial agents.	20/07/2011	Novartis Europharm Ltd
TORISEL	temsirolimus	Treatment of adult patients with relapsed and / or refractory mantle cell lymphoma (MCL) .	19/11/2007	Pfizer Ltd
TRANSLARNA	ataluren	Treatment of Duchenne muscular dystrophy resulting from a nonsense mutation in the dystrophin gene, in ambulatory patients aged 5 years and older. Efficacy has not been demonstrated in non-ambulatory patients. The presence of a nonsense mutation in the dystrophin gene should be determined by genetic testing.	31/07/2014	PTC Therapeutics Ltd
VENCLYXTO	venetoclax	As monotherapy for the treatment of chronic lymphocytic leukaemia (CLL) in the presence of 17p deletion or TP53 Mutation in adult patients who are unsuitable for or have failed a B-cell receptor pathway inhibitor. As monotherapy for the treatment of CLL in the absence of 17p deletion or TP53 mutation in adult patients who have failed both chemoimmunotherapy and a B-cell receptor pathway inhibitor.	05/12/2016	AbbVie Ltd

TRADENAME	ACTIVE SUBSTANCE	MARKETING AUTHORISATION INDICATION	MARKETING AUTHORISATION DATE (DD/MM/YYYY)	MARKETING AUTHORISATION HOLDER
VIDAZA	azacitidine	Treatment of adult patients who are not eligible for haematopoietic stem cell transplantation with: - intermediate-2 and high-risk myelodysplastic syndromes (MDS) according to the International Prognostic Scoring System (IPSS), - chronic myelomonocytic leukaemia (CMML) with 10-29% marrow blasts without myeloproliferative disorder, - acute myeloid leukaemia (AML) with 20-30 % blasts and multi-lineage dysplasia, according to World Health Organisation (WHO) classification. Treatment of adult patients aged 65 years or older who are not eligible for HSCT with AML with >30% marrow blasts according to the WHO classification.	17/12/2008	Celgene Europe Ltd
VIMIZIM	elosulfase alfa	Treatment of mucopolysaccharidosis, type IVA (Morquio A Syndrome, MPS IVA) in patients of all ages.	28/04/2014	BioMarin Europe Ltd
VOLIBRIS	ambrisentan	Treatment of pulmonary arterial hypertension (PAH) in adult patients of WHO Functional Class (FC) II to III, including use in combination treatment. Efficacy has been shown in idiopathic PAH (IPAH) and in PAH associated with connective tissue disease.	21/04/2008	Glaxo GroupLtd
VOTUBIA	everolimus	Treatment of adult patients with renal angiomyolipoma associated with tuberous sclerosis complex (TSC) who are at risk of complications (based on factors such as tumour size or presence of aneurysm, or presence of multiple or bilateral tumours) but who do not require immediate surgery. The evidence is based on analysis of change in sum of angiomyolipoma volume. Treatment of patients with subependymal giant cell astrocytoma (SEGA) associated with tuberous sclerosis complex (TSC) who require therapeutic intervention but are not amenable to surgery. The evidence is based on analysis of change in SEGA volume. Further clinical benefit, such as improvement in disease-related symptoms, has not been demonstrated.	02/09/2011	Novartis Europharm Ltd
VPRIV	velaglucerase alfa	Long-term enzyme replacement therapy (ERT) in patients with type 1 Gaucher disease .	26/08/2010	Shire Pharmaceuticals Ireland Ltd
VYNDAQEL	tafamidis	Treatment of transthyretin amyloidosis in adult patients with stage 1 symptomatic polyneuropathy to delay peripheral neurologic impairment.	16/11/2011	Pfizer Ltd
WAKIX	pitolisant	Treatment in adults of narcolepsy with or without cataplexy .	31/03/2016	Bioprojet Pharma
XALUPRINE (previously MERCAPTOPURINE NOVA)	mercaptopurine	Treatment of acute lymphoblastic leukaemia (ALL) in adults, adolescents and children.	09/03/2012	Nova Laboratories Ltd

TRADENAME	ACTIVE SUBSTANCE	MARKETING AUTHORISATION INDICATION	MARKETING AUTHORISATION DATE (DD/MM/YYYY)	MARKETING AUTHORISATION HOLDER
XERMELO	telotristat	Treatment of carcinoid syndrome diarrhoea in combination with somatostatin analogue (SSA) therapy in adults inadequately controlled by SSA therapy.	18/09/2017	Ipsen Pharma
YONDELIS	trabectedin	In combination with pegylated liposomal doxorubicin (PLD), treatment of patients with relapsed platinum-sensitive ovarian cancer .	28/10/2009	Pharma MarS.A.
ZALMOXIS	allogeneic T cells genetically modified with a retroviral vector encoding for a truncated form of the human low affinity nerve growth factor receptor (Δ LNNGFR) and the herpes simplex I virus thymidine kinase (HSV-TK Mut2)	Indicated as adjunctive treatment in haploidentical haematopoietic stem cell transplantation (HSCT) of adult patients with high-risk haematological malignancies .	18/08/2016	MolMed SpA
ZAVESCA	miglustat	Treatment of progressive neurological manifestations in adult patients and paediatric patients with Niemann-Pick type C disease .	28/01/2009	Actelion Registration Ltd
ZEJULA	niraparib	As monotherapy for the maintenance treatment of adult patients with platinum sensitive relapsed high grade serous epithelial ovarian, fallopian tube, or primary peritoneal cancer who are in response (complete or partial) to platinum based chemotherapy.	16/11/2017	Tesaro UK Limited



Allegato 1: Prodotti medicinali orfani ritirati dal Registro Europeo dei Prodotti Medicinali Orfani

Cf. Parte II "Elenco dei prodotti medicinali per le malattie rare in Europa con autorizzazione all'immissione in commercio senza designazione orfana europea".*

TRADENAME	ACTIVE SUBSTANCE	REGULAR STATUS	MARKETING AUTHORIZATION DATE	ORPHAN DESIGNATION WITHDRAWAL DATE
AFINITOR	everolimus	This product was withdrawn from the Community Register of designated orphan medicinal products on request of the sponsor . It was originally designated an orphan medicine on 5 June 2007.	05/08/2009	08/07/2011
ALDURAZYME	laronidase	This product was withdrawn from the Community register of orphan medicinal products at the end of the 10-year period of market exclusivity . It was originally designated an orphan medicine on 14 February 2001.	12/06/2003	12/06/2013
ATRIANCE	nelarabine	This product was withdrawn from the Community register of orphan medicinal products at the end of the 10-year period of market exclusivity . It was originally designated an orphan medicine on 16 June 2005.	22/08/2007	24/08/2017
BUSILVEX	busulfan	This product was withdrawn from the Community register of orphan medicinal products at the end of the 10-year period of market exclusivity . It was originally designated an orphan medicine on 29 December 2000	11/07/2003	11/07/2013
CARBAGLU	carglumic acid	This product was withdrawn from the Community register of orphan medicinal products at the end of the 10-year period of market exclusivity for the following condition: - Treatment of <i>N-acetylglutamate synthetase (NAGS) deficiency</i> . It was originally designated an orphan medicine for this indication on 18 October 2000.	28/01/2003	28/01/2013
CUPRIOR	trientine	This product was withdrawn from the Community Register of designated orphan medicinal products on request of the sponsor . It was originally designated an orphan medicine on 19 March 2015.	05/09/2017	20/07/2017
CYRAMZA	ramucirumab	This product was withdrawn from the Community Register of designated orphan medicinal products on request of the sponsor . It was originally designated an orphan medicine on 6 July 2012.	23/12/2014	27/01/2016
CYSTADANE	betaine anhydrous	This product was withdrawn from the Community register of orphan medicinal products at the end of the 10-year period of market exclusivity . It was originally designated an orphan medicine on 9 July 2001.	15/02/2007	19/02/2017
DIACOMIT	stiripentol	This product was withdrawn from the Community register of orphan medicinal products at the end of the 10-year period of market exclusivity . It was originally designated an orphan medicine on 5 December 2001.	04/01/2007	09/01/2017

TRADENAME	ACTIVE SUBSTANCE	REGULAR STATUS	MARKETING AUTHORIZATION DATE	ORPHAN DESIGNATION WITHDRAWAL DATE
ELAPRASE	idursulfase	This product was withdrawn from the Community register of orphan medicinal products at the end of the 10-year period of market exclusivity . It was originally designated an orphan medicine on 11 December 2001.	08/01/2007	10/01/2017
ELMIRON	pentosan polysulfate sodium	This product was withdrawn from the Community Register of designated Orphan Medicinal Products on request of the sponsor . It was originally designated an orphan medicine on 15 January 2015.	02/06/2017	11/04/2017
ELOCTA	efmorotocog alfa	This product was withdrawn from the Community Register of designated Orphan Medicinal Products on request of the sponsor . It was originally designated an orphan medicine on 20 September 2010.	23/11/2015	23/11/2015
EMPLICITI	elotuzumab	This product was withdrawn from the Community Register of designated orphan medicinal products by the European Commission at the time of the granting of a marketing authorisation . It was originally designated an orphan medicine on 9 August 2012.	11/05/2016	08/04/2016
EVOLTRA	clofarabine	This product was withdrawn from the Community register of orphan medicinal products at the end of the period of market exclusivity . It was originally designated an orphan medicine on 7 February 2002.	31/05/2006	31/05/2016
EXJADE	deferasirox	This product was withdrawn from the Community register of orphan medicinal products at the end of the period of market exclusivity . It was originally designated an orphan medicine on 13 mars 2002	01/09/2006	01/09/2016
FABRAZYME	agalsidase beta	This product was withdrawn from the Community register of orphan medicinal products at the end of the period of market exclusivity . It was originally designated an orphan medicine on 8 August 2000.	07/08/2001	07/08/2011
GLIOLAN	5-aminolevulinic acid hydrochloride	This product was withdrawn from the Community register of orphan medicinal products at the end of the 10-year period of market exclusivity . It was originally designated an orphan medicine on 13 November 2002.	07/09/2007	12/09/2017

TRADENAME	ACTIVE SUBSTANCE	REGULAR STATUS	MARKETING AUTHORIZATION DATE	ORPHAN DESIGNATION WITHDRAWAL DATE
GLIVEC	imatinib mesilate	<p>This product was withdrawn from the Community register of orphan medicinal products at the end of the 10-year period of market exclusivity for the following conditions:</p> <ul style="list-style-type: none"> - Treatment of chronic myeloid leukaemia (it was designated an orphan medicine on 14/02/2001). <p>It was withdrawn from the Community register of orphan medicinal products on April 2012 on request of the sponsor for the following conditions:</p> <ul style="list-style-type: none"> - Treatment of malignant gastrointestinal stromal tumours (it was designated an orphan medicine on 20/11/2001) - Treatment of dermatofibrosarcoma protuberans (it was designated an orphan medicine on 26/08/2005); - Treatment of acute lymphoblastic leukaemia (it was designated an orphan medicine on 26/08/2005); - Treatment of chronic eosinophilic leukaemia and the hypereosinophilic syndrome (it was designated an orphan medicine on 28/10/2005) - Treatment of myelodysplastic / myeloproliferative diseases (it was designated an orphan medicine on 23/12/2005) 	<p>12/11/2001</p> <p>27/05/2002</p> <p>18/09/2006</p> <p>18/09/2006</p> <p>01/12/2006</p> <p>01/12/2006</p>	<p>12/11/2011</p> <p>16/04/2012</p>
ILARIS	canakinumab	<p>This product was withdrawn from the Community Register of designated orphan medicinal products on request of the sponsor. It was originally designated an orphan medicine on 20 March 2007.</p>	27/10/2009	01/12/2010
INCRELEX	mecasermin	<p>This product was withdrawn from the Community register of orphan medicinal products at the end of the 10-year period of market exclusivity. It was originally designated an orphan medicine on 22 May 2006.</p>	03/08/2007	07/08/2017
IXIARO	Purified inactivated Japanese encephalitis SA14-4-2 virus vaccine	<p>This product was withdrawn from the Community Register of designated orphan medicinal products on request of the sponsor. It was originally designated an orphan medicine on 26 January 2006.</p>	02/04/2009	12/03/2009
JAKAVI	ruxolitinib	<p>This product was withdrawn from the Community Register of designated orphan medicinal products on request of the sponsor for the following conditions:</p> <ul style="list-style-type: none"> -Treatment of polycythaemia vera (19/02/2014) -Treatment of chronic idiopathic myelofibrosis (07/11/2008) -Treatment of myelofibrosis secondary to polycythaemia vera or essential thrombocythaemia (03/04/2009). 	28/08/2012	20/02/2015

TRADENAME	ACTIVE SUBSTANCE	REGULAR STATUS	MARKETING AUTHORIZATION DATE	ORPHAN DESIGNATION WITHDRAWAL DATE
JINARC	tolvaptan	This product was withdrawn from the Community Register of designated orphan medicinal products on request of the sponsor . It was originally designated an orphan medicine on 5 august 2013	29/05/2015	26/03/2015
LITAK	cladribine	This product was withdrawn from the Community register of orphan medicinal products at the end of the 10-year period of market exclusivity . It was originally designated an orphan medicine on 18 September 2001.	19/04/2004	19/04/2014
LYSODREN	mitotane	This product was withdrawn from the Community register of orphan medicinal products at the end of the 10-year period of market exclusivity . It was originally designated an orphan medicine on 12 June 2002.	30/04/2004	30/04/2014
MYOZYME	alglucosidase alfa	This product was withdrawn from the Community register of orphan medicinal products at the end of the 10-year period of market exclusivity . It was originally designated an orphan medicine on 22 February 2001.	31/03/2006	31/03/2016
NAGLAZYME	galsulfase	This product was withdrawn from the Community register of orphan medicinal products at the end of the 10-year period of market exclusivity . It was originally designated an orphan medicine on 22 February 2001.	26/01/2006	26/01/2016
NEOFORDEX	dexamethasone	This product was withdrawn from the Community Register of designated orphan medicinal products on request of the sponsor . It was originally designated an orphan medicine on 9 june 2010.	16/03/2016	25/01/2016
NEXAVAR	sorafenib tosylate	This product was withdrawn from the Community register of orphan medicinal products at the end of the 10-year period of market exclusivity for the following condition: -Treatment of renal cell carcinoma (it was designated an orphan medicine on 29/07/2004)	19/07/2006	22/07/2016
NOVOTHIRTEEN	catridecacog	This product was withdrawn from the Community Register of designated orphan medicinal products on request of the sponsor . It was originally designated an orphan medicine on 12 December 2003.	05/09/2012	01/07/2012
OBIZUR	Recombinant porcine factor VIII (B-domain-deleted)	This product was withdrawn from the Community Register of designated orphan medicinal products on request of the sponsor . It was originally designated an orphan medicine on 20 September 2010.	13/11/2015	23/10/2015
ORFADIN	nitisinone	This product was withdrawn from the Community register of orphan medicinal products at the end of the 10-year period of market exclusivity . It was originally designated an orphan medicine on 29 December 2000.	24/02/2005	24/02/2015

TRADENAME	ACTIVE SUBSTANCE	REGULAR STATUS	MARKETING AUTHORIZATION DATE	ORPHAN DESIGNATION WITHDRAWAL DATE
ORKAMBI	Lumacaftor / ivacaftor	This product was withdrawn from the Community Register of designated orphan medicinal products on request of the sponsor . It was originally designated an orphan medicine on 22 August 2014.	24/11/2015	12/10/2015
PEDEA	ibuprofen	This product was withdrawn from the Community register of orphan medicinal products at the end of the 10-year period of market exclusivity . It was originally designated an orphan medicine on 14 February 2001.	02/08/2004	02/08/2014
PRIALT	ziconotide	This product was withdrawn from the Community register of orphan medicinal products at the end of the 10-year period of market exclusivity . It was originally designated an orphan medicine on 9 July 2001.	24/02/2005	24/02/2015
QUINSAIR	levofloxacin	This product was withdrawn from the Community Register of designated orphan medicinal products on request of the sponsor . It was originally designated an orphan medicine on 23 September 2008.	30/03/2015	01/02/2015
REPLAGAL	agalsidase alfa	This product was withdrawn from the Community register of orphan medicinal products at the end of the 10-year period of market exclusivity . It was originally designated an orphan medicine on 8 August 2000.	07/08/2001	07/08/2011
REVATIO	Sildenafil citrate	This product was withdrawn from the Community register of orphan medicinal products at the end of the 10-year period of market exclusivity . It was originally designated an orphan medicine on 17 December 2003.	04/11/2005	04/11/2015
REVLIMID	lenalidomide	This product was withdrawn from the Community register of orphan medicinal products at the end of the 10-year period of market exclusivity for the following condition: - Treatment of multiple myeloma . It was originally designated an orphan medicine for this indication on 12 December 2003	14/06/2007	19/06/2017
REVOLADE	eltrombopag	This product was withdrawn from the Community Register of designated orphan medicinal products on request of the sponsor . It was originally designated an orphan medicine on 3 August 2007.	15/03/2010	01/01/2012
SAVENE	dexrazoxane	This product was withdrawn from the Community register of orphan medicinal products at the end of the 10-year period of market exclusivity . It was originally designated an orphan medicine on 19 september 2001	02/08/2006	02/08/2016

NEW

TRADENAME	ACTIVE SUBSTANCE	REGULAR STATUS	MARKETING AUTHORIZATION DATE	ORPHAN DESIGNATION WITHDRAWAL DATE
SIKLOS	hydroxycarbamide	Prevention of recurrent painful vaso-occlusive crises including acute chest syndrome in adults, adolescents and children older than 2 years suffering from symptomatic sickle cell syndrome . This product was withdrawn from the Community register of orphan medicinal products at the end of the 10-year period of market exclusivity . It was originally designated an orphan medicine on 9 July 2003.	29/06/2007	Addmedica
SOMAVERT	pegvisomant	This product was withdrawn from the Community register of orphan medicinal products at the end of the 10-year period of market exclusivity . It was originally designated an orphan medicine on 14 February 2001.	15/11/2002	15/11/2012
SPECTRILA	asparaginase	This product was withdrawn from the Community Register of designated orphan medicinal products on request of the sponsor . It was originally designated an orphan medicine on 26 January 2005.	18/01/2016	18/01/2016
SPRYCEL	dasatinib	This product was withdrawn from the Community register of orphan medicinal products at the end of the 10-year period of market exclusivity . It was originally designated an orphan medicine on 23 December 2005	20/11/2006	22/11/2016
SUTENT	sunitinib malate	This product was withdrawn from the Community Register of designated orphan medicinal products on request of the sponsor . It was originally designated an orphan medicine on 10 March 2005.	15/01/2007	23/07/2008
TORISEL	temsirolimus	First-line treatment of adult patients with advanced renal cell carcinoma (RCC) who have at least three of six prognostic risk factors. This product was withdrawn from the Community register of orphan medicinal products at the end of the 10-year period of market exclusivity . It was originally designated an orphan medicine on 6 April 2006.	19/11/2007	Pfizer Ltd

NEW

TRADENAME	ACTIVE SUBSTANCE	REGULAR STATUS	MARKETING AUTHORIZATION DATE	ORPHAN DESIGNATION WITHDRAWAL DATE
TRACLEER	bosentan monohydrate	<p>This product is no longer an orphan medicine.</p> <p>It was withdrawn from the Community register of orphan medicinal products on request of the sponsor for the following condition:</p> <p>-Treatment of systemic sclerosis (it was designated an orphan medicine on 17/03/2003)</p> <p>It was withdrawn from the Community register of orphan medicinal products at the end of the 10-year period of market exclusivity for the following condition:</p> <p>- Treatment of pulmonary arterial hypertension and chronic thromboembolic pulmonary hypertension (it was designated an orphan medicine on 14/02/2001)</p>	<p>11/06/2007</p> <p>17/05/2002</p>	<p>04/04/2014</p> <p>17/05/2012</p>
TRISENOX	arsenic trioxide	<p>This product was withdrawn from the Community register of orphan medicinal products at the end of the 10-year period of market exclusivity. It was originally designated an orphan medicine on 18 October 2000.</p>	07/03/2002	07/03/2012
UPTRAVI	selexipag	<p>This product was withdrawn from the Community Register of designated orphan medicinal products on request of the sponsor at the time of the granting of a marketing authorization. It was originally designated an orphan medicine on 26 August 2005.</p>	12/05/2016	22/02/2016
VENTAVIS	iloprost	<p>This product was withdrawn from the Community register of orphan medicinal products at the end of the 10-year period of market exclusivity. It was originally designated an orphan medicine on 29 December 2000.</p>	18/09/2003	18/09/2013
WILZIN	zinc acetate dihydrate	<p>This product was withdrawn from the Community register of orphan medicinal products at the end of the 10-year period of market exclusivity. It was originally designated an orphan medicine on 31 July 2001.</p>	18/10/2004	18/10/2014
XAGRID	anagrelide hydrochloride	<p>This product was withdrawn from the Community register of orphan medicinal products at the end of the 10-year period of market exclusivity+ 2 years for an agreed paediatric investigation plan (PIP). It was originally designated an orphan medicine on 29 December 2000.</p>	16/11/2004	18/11/2016
XYREM	sodium oxybate	<p>This product was withdrawn from the Community Register of designated orphan medicinal products on request of the sponsor. It was originally designated an orphan medicine on 3 February 2003.</p>	18/10/2005	11/01/2010

TRADENAME	ACTIVE SUBSTANCE	REGULAR STATUS	MARKETING AUTHORIZATION DATE	ORPHAN DESIGNATION WITHDRAWAL DATE
YONDELIS	trabectedin	This product was withdrawn from the Community register of orphan medicinal products at the end of the 10-year period of market exclusivity for the following condition: - Treatment of soft tissue sarcoma . It was originally designated an orphan medicine for this indication on 30 May 2001.	17/09/2007	21/09/2017
ZAVESCA	miglustat	This product was withdrawn from the Community register of orphan medicinal products at the end of the 10-year period of market exclusivity for the following condition: - Treatment of type 1 Gaucher disease . It was originally designated an orphan medicine for this indication on 18 October 2000.	21/11/2002	21/11/2012

Allegato 2: Prodotti medicinali orfani ritirati dal commercio nell'Unione Europea

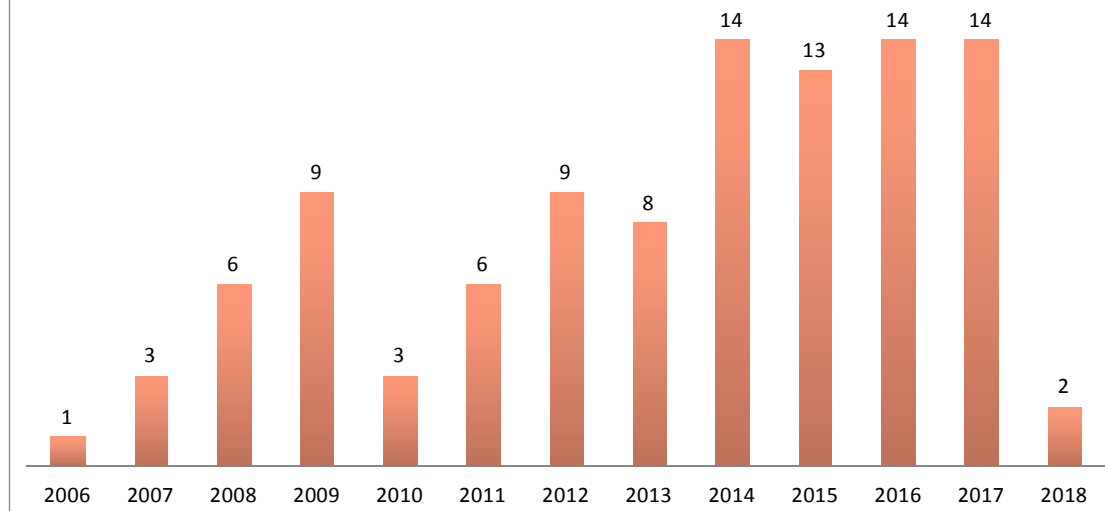
Ulteriori informazioni sul sito www.ema.europa.eu

TRADENAME	ACTIVE SUBSTANCE	MARKETING AUTHORISATION (MA) INDICATION	MA DATE / MA HOLDER	MA WITHDRAWN DATE
BIOGRASTIM	filgrastim	In patients, children or adults, with severe congenital, cyclic, or idiopathic neutropenia with an absolute neutrophil count (ANC) of $\leq 0.5 \times 10^9/l$, and a history of severe or recurrent infections, long term administration of Biograstim was indicated to increase neutrophil counts and to reduce the incidence and duration of infection-related events.	15/09/2008 AbZ-Pharma GmbH	25/09/2015
NUDEXTA	dextro methorphan hydrobromide / quinidine sulfate	Symptomatic treatment of pseudobulbar affect (PBA) in adults. Efficacy has been studied in patients with underlying Amyotrophic Lateral Sclerosis .	24/06/2013 Jenson Pharmaceutical Services Ltd	04/03/2016
ONSENAL	celecoxib	Reduction of the number of adenomatous intestinal polyps in familial adenomatous polyposis (FAP) , as an adjunct to surgery and further endoscopic surveillance.	17/10/2003 Pfizer Ltd	28/03/2011
PHOTOBARR	porfimer sodium (for use with photodynamic therapy)	Ablation of high-grade dysplasia (HGD) in patients with Barrett's oesophagus .	25/03/2004 Pinnacle Biologics B.V.	20/04/2012
RILONACEPT REGENERON (previously ARCALYST)	rilonacept	Treatment of Cryopyrin-Associated Periodic Syndromes (CAPS) with severe symptoms, including Familial Cold Autoinflammatory Syndrome (FCAS) and Muckle-Wells Syndrome (MWS) , in adults and children aged 12 years and older.	23/10/2009 Regeneron UK Ltd	24/10/2012
THELIN	sitaxentan sodium	Treatment of patients with pulmonary arterial hypertension classified as WHO functional class III, to improve exercise capacity. Efficacy has been shown in primary pulmonary hypertension and in pulmonary hypertension associated with connective tissue disease.	10/08/2006 Pfizer Ltd	06/01/2011
UNITUXIN	dinutuximab	Treatment of high-risk neuroblastoma in patients aged 12 months to 17 years, who have previously received induction chemotherapy and achieved at least a partial response, followed by myeloablative therapy and autologous stem cell transplantation (ASCT). It is administered in combination with granulocyte-macrophage colony-stimulating factor (GM-CSF), interleukin-2 (IL-2), and isotretinoin.	14/08/2015 United Therapeutics Europe Ltd	20/03/2017

Classificazione per data dell'AIC in ordine decrescente

2018	VENCLYXTO	VIMIZIM	VPRIV
JORVEZA	WAKIX	2013	2009
PREVYMIS	ZALMOXIS	BOSULIF	CAYSTON
2017	2015	DEFITELIO	FIRDAPSE
BAVENCIO	BLINCYTO	ICLUSIG	MEPACT
BESPONSA	CERDELGA	IMNOVID	MOZOBIL
BRINEURA	CRESEMBA	OPSUMIT	NPLATE
CHENODEOXYCHOLIC ACID LEADIANT	FARYDAK	ORPHACOL	PEYONA
CYSTADROPS	HETLIOZ	PROCYSBI	TORISEL
DINUTUXIMAB BETA APEIRON	HOLOCLAR	REVLIMID	YONDELIS
LEDAGA	KANUMA	2012	ZAVESCA
LUTATHERA	KYPROLIS	ADCETRIS	2008
NATPAR	LENVIMA	BRONCHITOL	CEPLENE
OXERVATE	OFEV	DACOGEN	FIRAZYR
RYDAPT	RAVICTI	GLYBERA	KUVAN
SPINRAZA	RAXONE	KALYDECO	THALIDOMIDE CELGENE
XERMELO	STRENSIQ	NEXOBRID	VIDAZA
ZEJULA	2014	REVESTIVE	VOLIBRIS
2016	ADEMPAS	SIGNIFOR	2007
ALPROLIX	COMETRIQ	XALUPRINE	INOVELON
COAGADEX	DELTYBA	2011	SOLIRIS
DARZALEX	GAZYVARO	CARBAGLU	TASIGNA
GALAFOLD	GRANUPAS	ESBRIET	2006
IDELVION	IMBRUVICA	PLENADREN	NEXAVAR
LARTRUVO	KETOCONAZOLE HRA	TOBI PODHALER	
NINLARO	KOLBAM	VOTUBIA	
OCALIVA	LYNPARZA	VYNDAQEL	
ONIVYDE	SCENESSE	2010	
SOMAKIT TOC	SIRTURO	ARZERRA	
STRIMVELIS	SYLVANT	TEPADINA	
	TRANSLARNA		

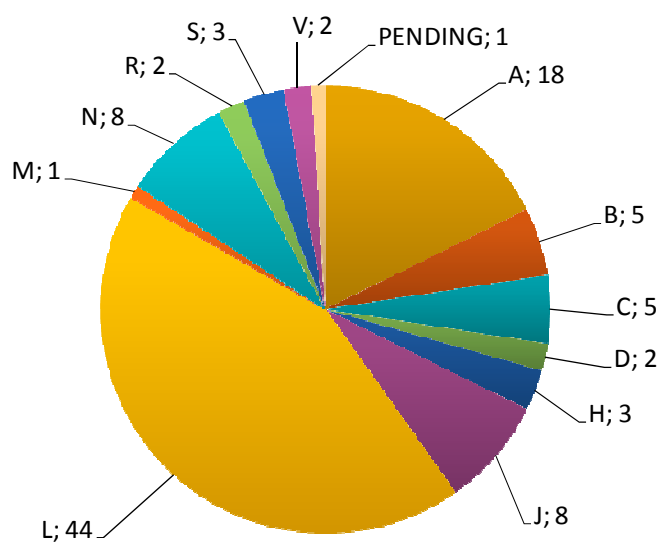
Numero dei farmaci orfani in Europa con designazione orfana europea e autorizzazione all'immissione in commercio europea per data dell'AIC



Classificazione per categoria ATC

A- ALIMENTARY TRACT AND METABOLISM	GLYBERA	DACOGEN	VOTUBIA
BRINEURA	OPSUMIT	DARZALEX	XALUPRINE
CARBAGLU	VOLIBRIS	DINUTUXIMAB BETA APEIRON	YONDELIS
CERDELGA	D- DERMATOLOGICALS	ESBRIET	ZALMOXIS
CHENODEOXYCHOLIC ACID LEADIANT	NEXOBRID	FARYDAK	ZEJULA
GALAFOLD	SCENESSE	GAZYVARO	M- MUSCULO-SKELETAL SYSTEM
JORVEZA	H- SYSTEMIC HORMONAL PREPARATIONS, EXCL. SEX HORMONES AND INSULINS	ICLUSIG	TRANSLARNA
KANUMA	NATPAR	IMBRUVICA	N- NERVOUS SYSTEM
KOLBAM	PLENADREN	IMNOVID	FIRDAPSE
KUVAN	SIGNIFOR	KYPROLIS	HETLIOZ
OICALIVA	J- GENERAL ANTIINFECTIVES FOR SYSTEMIC USE	LARTRUVO	INOVELON
ORPHACOL	CAYSTON	LEDAGA	PEYONA
PROCYSBI	CRESEMBA	LENVIMA	RAXONE
RAVICTI	DELTYBA	LYNPARZA	SPINRAZA
REVESTIVE	GRANUPAS	MEPACT	VYNDAQEL
STRENSIQ	KETOCONAZOLE	MOZOBIL	WAKIX
VIMIZIM	PREVYMIS	NEXAVAR	R- RESPIRATORY
VPRIV	SIRTURO	NINLARO	BRONCHITOL
ZAVESCA	TOBI PODHALER	OFEV	KALYDECO
B- BLOOD AND BLOOD FORMING	L- ANTINEOPLASTIC AND IMMUNOMODULATING	ONIVYDE	S- SENSORY ORGANS
ALPROLIX	ADCETRIS	REVLIMID	CYSTADROPS
COAGADEX	ARZERRA	RYDAPT	HOLOCLAR
DEFITELIO	BAVENCIO	SOLIRIS	OXERVATE
IDELVION	BESPONSA	STRIMVELIS	V- VARIOUS
NPLATE	BLINCYTO	SYLVANT	LUTATHERA
C-CARDIOVASCULAR SYSTEM	BOSULIF	TASIGNA	SOMAKIT TOC
ADEMPAS	CEPLENE	TEPADINA	PENDING
FIRAZYR	COMETRIQ	THALIDOMIDE CELGENE	XERMELO
		TORISEL	
		VENCLYXTO	
		VIDAZA	

Numero dei farmaci orfani in Europa con designazione orfana europea e autorizzazione all'immissione in commercio europea per categoria ATC



Classificazione per titolare dell'AIC

ABBVIE LTD	OFEV	PHARMA	PTC THERAPEUTICS LTD
VENCLYXTO	CELGENE EUROPE LTD	KETOCONAZOLE HRA	TRANSLARNA
ACTELION	IMNOVID	LABORATOIRES CTRS	RAPTOR
REGISTRATION LTD	REVLIMID	ORPHACOL	PHARMACEUTICALS
LEDAGA	THALIDOMIDE	LEADIANT GmbH	EUROPE B.V.
OPSUMIT	CELGENE	CHENODEOXYCHOLIC	PROCYSBI
ZAVESCA	VIDAZA	ACID LEADIANT	RETROPHIN EUROPE LTD
ADIENNE SRL	CHIESI FARMACEUTICI SPA	LUCANE PHARMA	KOLBAM
TEPADINA	HOLOCLAR	GRANUPAS	ROCHE REGISTRATION LTD
ADVANCED	PEYONA	MEDIWOUND GERMANY	ESBRIET
ACCELERATOR	CLINUVEL UK LIMITED	GMBH	GAZYVARO
APPLICATIONS	SCENESSE	NEXOBRID	SANTHERA
LUTATHERA	CSL BEHRING GMBH	MERCK SERONO EUROPE	PHARMACEUTICALS
SOMAKIT TOC	IDELVION	LTD	(DEUTSCHLAND) GMBH
ALEXION EUROPE SAS	DOMPE FARMACEUTICI	BAVENCIO	RAXONE
SOLIRIS	S.P.A.	MERCK SHARP & DOHME	SHIRE ORPHAN THERAPIES
STRENSIQ	OXERVATE	LIMITED	GMBH
AMGEN EUROPE B.V.	DR. FALK PHARMA GMBH	PREVYMIS	FIRAZYR
BLINCYTO	JORVEZA	KUVAN	SHIRE PHARMACEUTICALS
KYPROLIS	ELI LILLY B.V.	MolMed SpA	IRELAND LTD
NPLATE	LARTRUVO	ZALMOXIS	NATPAR
AMICUS THERAPEUTICS	ESAI LTD	NOVA LABORATORIES LTD	VPRIV
UK LTD	INOVELON	XALUPRINE	SYNAGEVA BIOPHARMA
GALAFOLD	LENVIMA	NOVARTIS EUROPHARM	LTD
APEIRON BIOLOGICS AG	GENTIUM SPA	ARZERRA	KANUMA
DINUTUXIMAB BETA	DEFITELIO	FARYDAK	TAKEDA FRANCE SAS
APEIRON	GENZYME EUROPE B.V.	RYDAPT	MEPACT
ARIAD PHARMA LTD	CERDELGA	SIGNIFOR	TAKEDA PHARMA A/S.
ICLUSIG	MOZOBIL	TASIGNA	ADCETRIS
ASTRA ZENECA AB	GILEAD SCIENCES	TOBI PODHALER	NINLARO
LYNPARZA	INTERNATIONAL LTD	VOTUBIA	TESARO UK LIMITED
BASILEA MEDICAL LTD	CAYSTON	NOVENTIA PHARMA SRL	ZEJULA
CRESEMBA	GLAXO GROUP LTD	CEPLENE	TMC PHARMA SERVICES
BAXALTA INNOVATIONS	VOLIBRIS	NPS PHARMA HOLDINGS	LTD.
ONIVYDE	GLAXOSMITHKLINE	LIMITED	COMETRIQ
BAYER PHARMA AG	TRADING SERVICES	REVESTIVE	UNIQUIRE BIOPHARMA B.V.
ADEMPAS	LIMITED	ORPHAN EUROPE SARL	GLYBERA
NEXAVAR	STRIMVELIS	CARBAGLU	VANDA
BIO PRODUCTS	HORIZON THERAPEUTICS	CYSTADROPS	PHARMACEUTICALS LTD
LABORATORY LTD	Ltd	OTSUKA NOVEL	HETLIOZ
COAGADEX	RAVICTI	PRODUCTS GMBH	VERTEX
BOGEN IDEC LTD	INTERCEPT PHARMA	DELTYBA	PHARMACEUTICALS
ALPROLIX	OCALIVA	PFIZER LTD	(EUROPE) LTD
SPINRAZA	IPSEN PHARMA	BESPONSA	KALYDECO
BIOMARIN EUROPE LTD	XERMELO	BOSULIF	VIOPHARMA SPRL
BRINEURA	JANSSEN-CILAG	TORISEL	PLENADREN
FIRDAPSE	INTERNATIONAL NV	VYNDAQEL	
VIMIZIM	DACOGEN	PHARMA MAR S.A.	
BIOPROJET PHARMA	DARZALEX	YONDELIS	
WAKIX	IMBRUVICA	PHARMAXIS	
BOEHRINGER INGELHEIM	SIRTURO	PHARMACEUTICALS LTD	
INTERNATIONAL GMBH	SYLVANT	BRONCHITOL	
	LABORATOIRE HRA		

PARTE 2:

Elenco dei prodotti medicinali per le malattie rare in Europa con autorizzazione all'immissione in commercio* senza designazione orfana europea

Indice

Elenco dei prodotti medicinali per le malattie rare in Europa con autorizzazione all'immissione in commercio* senza designazione orfana europea	33
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Metodologia

Questa parte del documento fornisce l'elenco di tutti i prodotti medicinali per le malattie rare che hanno ottenuto l'autorizzazione all'immissione in commercio (AIC) europea per una o più indicazioni terapeutiche per una malattia rara, ma che non hanno ottenuto una designazione orfana europea o per i quali la designazione è stata ritirata.

Questi prodotti medicinali possono avere ottenuto (o non ottenuto) una designazione orfana in un'altra area geografica del mondo e sono disponibili nell'elenco fornito dalla DG SANTE relativo ai prodotti medicinali che hanno ottenuto l'autorizzazione all'immissione in commercio: <http://ec.europa.eu/health/documents/communityregister/html/alfregister.htm>

Una prima classificazione per denominazione commerciale fornisce il nome della sostanza

attiva, l'indicazione rara dell'autorizzazione all'immissione in commercio, la data e il titolare dell'AIC.

Tre elenchi aggiuntivi propongono altre classificazioni per:

- Data dell'AIC in ordine decrescente;
- Categoria ATC;
- Titolare dell'AIC.

In ciascun elenco, le denominazioni commerciali sono disposte in ordine alfabetico.

Ulteriori informazioni sono disponibili nella sezione "Farmaci orfani" del sito web di Orphanet, www.orpha.net, o sul sito web dell'EMA (Agenzia Europea per i Medicinali) <http://www.ema.europa.eu>.

* Autorizzazione all'immissione in commercio dell'Unione Europea mediante procedura centralizzata

Classificazione per denominazione commerciale

Tradename	Active Substance	Marketing Authorisation Indication	Marketing Authorisation Date (Dd/Mm/Yyyy)	Marketing Authorisation Holder
ABRAXANE	paclitaxel	In combination with gemcitabine is indicated for the first-line treatment of adult patients with metastatic adenocarcinoma of the pancreas .	11/01/2008	Celgene Europe Ltd
ADCIRCA	tadalafil	In adults for the treatment of pulmonary arterial hypertension (PAH) classified as WHO functional class II and III, to improve exercise capacity . Efficacy has been shown in idiopathic PAH (IPAH) and in PAH related to collagen vascular disease.	01/10/2008	Eli Lilly Nederland B.V.
ADVATE	octocog alpha	Treatment and prophylaxis of bleeding in patients with haemophilia A (congenital factor VIII deficiency) . It is indicated in all age groups.	02/03/2004	Baxter AG
ADYNOVI	rurioctocog alfa pegol	Treatment and prophylaxis of bleeding in patients 12 years and above with haemophilia A (congenital factor VIII deficiency) .	08/01/2018	Baxalta Innovations GmbH
AFINITOR	everolimus	Treatment of unresectable or metastatic, well- or moderately-differentiated neuroendocrine tumours of pancreatic origin in adults with progressive disease. Treatment of unresectable or metastatic, well-differentiated (Grade 1 or Grade 2) non-functional neuroendocrine tumours of gastrointestinal or lung origin in adults with progressive disease Treatment of patients with advanced renal cell carcinoma , whose disease has progressed on or after treatment with VEGF-targeted therapy.	03/08/2009	Novartis Europharm Ltd
AFSTYLA	lonoctocog alfa	Treatment and prophylaxis of bleeding in patients with haemophilia A (congenital factor VIII deficiency) .	04/01/2017	CSL Behring GmbH
ALDURAZYME	laronidase	Long-term enzyme replacement therapy in patients with a confirmed diagnosis of Mucopolysaccharidosis I (MPS I; a [alpha]-L-iduronidase deficiency) to treat the non-neurological manifestations of the disease.	10/06/2003	Genzyme Europe B.V.
ALIMTA	pemetrexed	Treatment of chemotherapy naïve patients with unresectable malignant pleural mesothelioma .	20/09/2004	Eli Lilly Nederland B.V.
AMGEVITA	Adalimumab	In combination with methotrexate is indicated for the treatment of active polyarticular juvenile idiopathic arthritis , in patients from the age of 2 years who have had an inadequate response to one or more disease-modifying anti-rheumatic drugs (DMARDs). AMGEVITA can be given as monotherapy in case of intolerance to methotrexate or when continued treatment with methotrexate is inappropriate. Adalimumab has not been studied in patients aged less than 2 years. Treatment of active enthesitis-related arthritis in patients, 6 years of age and older, who have had an inadequate response to, or who are intolerant of, conventional therapy. Treatment of non-infectious intermediate, posterior and panuveitis in adult patients who have had an inadequate response to corticosteroids, in patients in need of corticosteroid-sparing, or in whom corticosteroid treatment is inappropriate.	22/03/2017	Amgen Europe B.V.

Tradename	Active Substance	Marketing Authorisation Indication	Marketing Authorisation Date (Dd/Mm/Yyyy)	Marketing Authorisation Holder
AMMONAPS	sodium phenylbutyrate	Adjunctive therapy in the chronic management of urea cycle disorders , involving deficiencies of carbamyl phosphate synthetase, ornithine transcarbamylase, or argininosuccinate synthetase . It is indicated in all patients with <i>neonatal-onset</i> presentation (complete enzyme deficiencies, presenting within the first 28 days of life). It is also indicated in patients with <i>late-onset</i> disease (partial enzyme deficiencies, presenting after the first month of life) who have a history of hyperammonaemic encephalopathy.	08/12/1999	Swedish Orphan Biovitrum AB
ARMISARTE (previously PEMETREXED ACTAVIS)	pemetrexed	In combination with cisplatin is indicated for the treatment of chemotherapy naïve patients with unresectable malignant pleural mesothelioma .	18/01/2016	Actavis Group PTC ehf
ATRIANCE	nelarabine	Treatment of patients with T-cell acute lymphoblastic leukaemia (T-ALL) and T-cell lymphoblastic lymphoma (T-LBL) whose disease has not responded to or has relapsed following treatment with at least two chemotherapy regimens. Due to the small patient populations in these disease settings, the information to support these indications is based on limited data.	22/08/2007	Novartis Europharm Ltd
ATRYN	antithrombin alpha	Prophylaxis of venous thromboembolism in surgery of adult patients with congenital antithrombin deficiency . It is normally given in association with heparin or low molecular weight heparin.	28/07/2006	GTC Biotherapeutics UK Ltd
AVASTIN	bevacizumab	In combination with interferon alfa-2a it is indicated for first line treatment of adult patients with advanced and/or metastatic renal cell cancer . In combination with carboplatin and paclitaxel, it is indicated for the front-line treatment of adult patients with advanced (International Federation of Gynecology and Obstetrics (FIGO) stages III B, III C and IV) epithelial ovarian, fallopian tube, or primary peritoneal cancer . In combination with carboplatin and gemcitabine or in combination with carboplatin and paclitaxel, is indicated for treatment of adult patients with first recurrence of platinum-sensitive epithelial ovarian, fallopian tube or primary peritoneal cancer who have not received prior therapy with bevacizumab or other VEGF inhibitors or VEGF receptor-targeted agents. In combination with paclitaxel, topotecan, or pegylated liposomal doxorubicin it is indicated for the treatment of adult patients with platinum-resistant recurrent epithelial ovarian, fallopian tube, or primary peritoneal cancer who received no more than two prior chemotherapy regimens and who have not received prior therapy with bevacizumab or other VEGF inhibitors or VEGF receptor-targeted agents.	12/01/2005	Roche Registration Ltd
BEMFOLA	follitropin alfa	In adult men: stimulation of spermatogenesis in men who have congenital or acquired hypogonadotropic hypogonadism with concomitant human chorionic gonadotropin (hCG) therapy.	27/03/2014	Finox BiotechAG

Tradename	Active Substance	Marketing Authorisation Indication	Marketing Authorisation Date (Dd/Mm/Yyyy)	Marketing Authorisation Holder
BENEFIX	nonacog alpha	Treatment and prophylaxis of bleeding in patients with haemophilia B (congenital factor IX deficiency) .	27/08/1997	Pfizer Ltd
BLITZIMA	rituximab	Treatment of previously untreated patients with stage III-IV follicular lymphoma in combination with chemotherapy. As maintenance therapy indicated for the treatment of follicular lymphoma patients responding to induction therapy. As monotherapy indicated for the treatment of patients with stage III-IV follicular lymphoma who are chemo-resistant or are in their second or subsequent relapse after chemotherapy. Treatment of patients with CD20 positive diffuse large B cell non-Hodgkin's lymphoma in combination with CHOP (cyclophosphamide, doxorubicin, vincristine, prednisolone) chemotherapy. In combination with chemotherapy for the treatment of patients with previously untreated and relapsed/refractory CLL. Only limited data are available on efficacy and safety for patients previously treated with monoclonal antibodies including Blitzima or patients refractory to previous Blitzima plus chemotherapy.	13/07/2017	Celltrion Healthcare Hungary Kft.
BORTEZOMIB ACCORD	bortezomib	As monotherapy or in combination with pegylated liposomal doxorubicin or dexamethasone for the treatment of adult patients with progressive multiple myeloma who have received at least 1 prior therapy and who have already undergone or are unsuitable for haematopoietic stem cell transplantation. In combination with melphalan and prednisone for the treatment of adult patients with previously untreated multiple myeloma who are not eligible for high-dose chemotherapy with haematopoietic stem cell transplantation. In combination with dexamethasone, or with dexamethasone and thalidomide, for the induction treatment of adult patients with previously untreated multiple myeloma who are eligible for high-dose chemotherapy with haematopoietic stem cell transplantation. In combination with rituximab, cyclophosphamide, doxorubicin and prednisone for the treatment of adult patients with previously untreated mantle cell lymphoma who are unsuitable for haematopoietic stem cell transplantation.	20/07/2015	Accord Healthcare Ltd

Tradename	Active Substance	Marketing Authorisation Indication	Marketing Authorisation Date (Dd/Mm/Yyyy)	Marketing Authorisation Holder
BORTEZOMIB HOSPIRA	bortezomib	As monotherapy or in combination with pegylated liposomal doxorubicin or dexamethasone is indicated for the treatment of adult patients with progressive multiple myeloma who have received at least 1 prior therapy and who have already undergone or are unsuitable for haematopoietic stem cell transplantation. In combination with melphalan and prednisone is indicated for the treatment of adult patients with previously untreated multiple myeloma who are not eligible for high-dose chemotherapy with haematopoietic stem cell transplantation. In combination with dexamethasone, or with dexamethasone and thalidomide, is indicated for the induction treatment of adult patients with previously untreated multiple myeloma who are eligible for high-dose chemotherapy with haematopoietic stem cell transplantation. In combination with rituximab, cyclophosphamide, doxorubicin and prednisone is indicated for the treatment of adult patients with previously untreated mantle cell lymphoma who are unsuitable for haematopoietic stem cell transplantation.	22/07/2016	Hospira UK Limited
BORTEZOMIB SUN	bortezomib	As monotherapy or in combination with pegylated liposomal doxorubicin or dexamethasone is indicated for the treatment of adult patients with progressive multiple myeloma who have received at least 1 prior therapy and who have already undergone or are unsuitable for haematopoietic stem cell transplantation. In combination with melphalan and prednisone is indicated for the treatment of adult patients with previously untreated multiple myeloma who are not eligible for high-dose chemotherapy with haematopoietic stem cell transplantation. In combination with dexamethasone, or with dexamethasone and thalidomide, is indicated for the induction treatment of adult patients with previously untreated multiple myeloma who are eligible for high-dose chemotherapy with haematopoietic stem cell transplantation. In combination with rituximab, cyclophosphamide, doxorubicin and prednisone is indicated for the treatment of adult patients with previously untreated mantle cell lymphoma who are unsuitable for haematopoietic stem cell transplantation.	22/07/2016	SUN Pharmaceutical Industries (Europe) B.V.
BUCCOLAM	midazolam	Treatment of prolonged, acute, convulsive seizures in infants, toddlers, children and adolescents (from 3 months to < 18 years). Buccolam must only be used by parents/carers where the patient has been diagnosed to have epilepsy . For infants between 3-6 months of age treatment should be in a hospital setting where monitoring is possible and resuscitation equipment is available.	05/09/2011	ViroPharma SPRL

Tradenome	Active Substance	Marketing Authorisation Indication	Marketing Authorisation Date (Dd/Mm/Yyyy)	Marketing Authorisation Holder
BUSILVEX	busulfan	Followed by cyclophosphamide (BuCy2), conditioning treatment prior to conventional haematopoietic progenitor cell transplantation in adult patients when the combination is considered the best available option. Following fludarabine (FB), conditioning treatment prior to haematopoietic progenitor cell transplantation in adult patients who are candidates for a reduced-intensity conditioning (RIC) regimen. Followed by cyclophosphamide (BuCy4) or melphalan (BuMel), conditioning treatment prior to conventional haematopoietic progenitor cell transplantation in paediatric patients.	09/07/2003	Pierre Fabre Médicament
CABOMETYX	cabozantinib	Treatment of advanced renal cell carcinoma (RCC) in adults following prior vascular endothelial growth factor (VEGF)-targeted therapy.	09/09/2016	Ipsen Pharma
CAELYX	doxorubicin hydrochloride (pegylated liposomal)	Treatment of advanced ovarian cancer in women who have failed a first-line platinum-based chemotherapy regimen. In combination with bortezomib for the treatment of progressive multiple myeloma in patients who have received at least one prior therapy and who have already undergone or are unsuitable for bone marrow transplant. Treatment of AIDS-related Kaposi's sarcoma (KS) in patients with low CD4 counts (< 200 CD4 lymphocytes/mm ³) and extensive mucocutaneous or visceral disease. Used as first-line systemic chemotherapy, or as second line chemotherapy in AIDS-KS patients with disease that has progressed with, or in patients intolerant to, prior combination systemic chemotherapy comprising at least two of the following agents: a vinca alkaloid, bleomycin and standard doxorubicin (or other anthracycline).	21/06/1996	Janssen-Cilag International N.V.
CANCIDAS (previously CASPOFUNGIN MSD)	casprofungin	Treatment of invasive candidiasis in adult or paediatric patients. Treatment of invasive aspergillosis in adult or paediatric patients who are refractory to or intolerant of amphotericin B, lipid formulations of amphotericin B and/or itraconazole. Empirical therapy for presumed fungal infections (such as Candida or Aspergillus) in febrile, neutropaenic adult or paediatric patients.	24/10/2001	Merck Sharp & Dohme Ltd
CAPRELSA	vandetanib	Treatment of aggressive and symptomatic medullary thyroid cancer (MTC) in patients with unresectable locally advanced or metastatic disease. Caprelsa is indicated in adults, children and adolescents aged 5 years and older. For patients in whom Rearranged during Transfection (RET) mutation is not known or is negative, a possible lower benefit should be taken into account before individual treatment decision.	17/02/2012	AstraZenecaAB
CARBAGLU	carglumic acid	Treatment of hyperammonaemia due to N-acetylglutamate synthase (NAGS) primary deficiency	28/01/2003	Orphan Europe S.A.R.L.

Tradename	Active Substance	Marketing Authorisation Indication	Marketing Authorisation Date (Dd/Mm/Yyyy)	Marketing Authorisation Holder
CEPROTIN	human protein c	In purpura fulminans and coumarin-induced skin necrosis in patients with severe congenital protein C deficiency . Short-term prophylaxis in patients with severe congenital protein C deficiency : if surgery or invasive therapy is imminent, while initiating coumarin therapy, when coumarin therapy alone is not sufficient, when coumarin therapy is not feasible.	16/07/2001	Baxter AG
CEREZYME	imiglucerase	Long-term enzyme replacement therapy in patients with a confirmed diagnosis of non-neuronopathic (Type 1) or chronic neuronopathic (Type 3) Gaucher disease and who exhibit clinically significant. The non-neurological manifestations of Gaucher disease include one or more of the following conditions: -Anaemia after exclusion of other causes, such as iron deficiency -Thrombocytopenia -Bone disease after exclusion of other causes such as Vitamin D deficiency -Hepatomegaly or splenomegaly	17/11/1997	Genzyme Europe B.V.
CINRYZE	C1 inhibitor(human)	Treatment and pre-procedure prevention of angioedema attacks in adults and adolescents with hereditary angioedema (HAE) . Routine prevention of angioedema attacks in adults and adolescents with severe and recurrent attacks of hereditary angioedema (HAE), who are intolerant to or insufficiently protected by oral prevention treatments, or patients who are inadequately managed with repeated acute treatment.	15/06/2011	ViroPharma SPRL
COLOBREATHE	colistimethate sodium	Management of chronic pulmonary infections due to <i>Pseudomonas aeruginosa</i> in patients with cystic fibrosis (CF) aged 6 years and older. Consideration should be given to official guidance on the appropriate use of antibacterial agents.	13/02/2012	Forest Laboratories UK Ltd
CUPRIOR	trientine	Treatment of Wilson's disease in adults, adolescents and children ≥ 5 years intolerant to D-penicillamine therapy.	05/09/2017	GMP-Orphan SA
CYLTEZO	adalimumab	In combination with methotrexate is indicated for the treatment of active polyarticular juvenile idiopathic arthritis , in patients from the age of 2 years who have had an inadequate response to one or more disease-modifying anti-rheumatic drugs (DMARDs). Adalimumab can be given as monotherapy in case of intolerance to methotrexate or when continued treatment with methotrexate is inappropriate. Adalimumab has not been studied in patients aged less than 2 years. Treatment of active enthesitis-related arthritis in patients, 6 years of age and older, who have had an inadequate response to, or who are intolerant of, conventional therapy. Treatment of non-infectious intermediate, posterior and panuveitis in adult patients who have had an inadequate response to corticosteroids, in patients in need of corticosteroid-sparing, or in whom corticosteroid treatment is inappropriate.	10/11/2017	Boehringer Ingelheim International GmbH

Tradename	Active Substance	Marketing Authorisation Indication	Marketing Authorisation Date (Dd/Mm/Yyyy)	Marketing Authorisation Holder
CYSTADANE	betaine anhydrous	Adjunctive treatment of homocystinuria , involving deficiencies or defects in cystathionine beta- synthase (CBS), 5,10-methylene-tetrahydrofolate reductase (MTHFR), cobalamin cofactor metabolism (cbl) . Cystadane should be used as supplement to other therapies such as vitamin B6 (pyridoxine), vitamin B12 (cobalamin), folate and a specific diet.	15/02/2007	Orphan Europe S.a.r.l.
CYSTAGON	mercaptamine bitartrate	Treatment of proven nephropathic cystinosis . Cysteamine reduces cystine accumulation in some cells (e.g. leukocytes, muscle and liver cells) of nephropathic cystinosis patients and, when treatment is started early, it delays the development of renal failure.	23/06/1997	Orphan Europe S.A.R.L
DEPOCYTE	cytarabine	Intrathecal treatment of lymphomatous meningitis . In the majority of patients such treatment will be part of symptomatic palliation of the disease.	11/07/2001	Pacira Ltd
DIACOMIT	stiripentol	Used in conjunction with clobazam and valproate as adjunctive therapy of refractory generalized tonic-clonic seizures in patients with severe myoclonic epilepsy in infancy (SMEI, Dravet's syndrome) whose seizures are not adequately controlled with clobazam and valproate.	04/01/2007	Biocodex
DUKORAL	vibrio cholerae and recombinant cholera toxinb-subunit	Indicated for active immunisation against disease caused by Vibrio cholerae serogroup O1 in adults and children from 2 years of age who will be visiting endemic/epidemic areas. The use of Dukoral should be determined on the basis of official recommendations taking into consideration the variability of epidemiology and the risk of contracting disease in different geographical areas and travelling conditions. Dukoral should not replace standard protective measures. In the event of diarrhoea measures of rehydration should be instituted.	28/04/2004	Crucell Sweden AB
ELAPRASE	idursulfase	Long-term treatment of patients with Hunter syndrome (Mucopolysaccharidosis II, MPS II) . Heterozygous females were not studied in the clinical trials.	08/01/2007	Shire Human Genetic Therapies AB
ELMIRON	pentosan polysulfate sodium	Treatment of bladder pain syndrome characterized by either glomerulations or Hunner's lesions in adults with moderate to severe pain, urgency and frequency of micturition.	02/06/2017	bene-Arzneimittel GmbH
ELOCTA	efmoroctocog alfa	Treatment and prophylaxis of bleeding in patients with haemophilia A (congenital factor VIII deficiency). ELOCTA can be used for all age groups.	19/11/2015	Biogen Idec Ltd
EMPLICITI	elotuzumab	In combination with lenalidomide and dexamethasone for the treatment of multiple myeloma in adult patients who have received at least one prior therapy.	11/05/2016	Bristol-Myers Squibb

Tradename	Active Substance	Marketing Authorisation Indication	Marketing Authorisation Date (Dd/Mm/Yyyy)	Marketing Authorisation Holder
ENBREL	etanercept	Treatment of polyarthritis (rheumatoid-factorpositive or -negative) and extended oligoarthritis in children and adolescents from the age of 2 years who have had an inadequate response to, or who have proved intolerant of, methotrexate. Treatment of psoriatic arthritis in adolescents from the age of 12 years who have had an inadequate response to, or who have proved intolerant of, methotrexate. Treatment of enthesitis-related arthritis in adolescents from the age of 12 years who have had an inadequate response to, or who have proved intolerant of conventional therapy. Enbrel has not been studied in children aged less than 2 years.	03/02/2000	Pfizer Ltd
ERBITUX	cetuximab	Treatment of patients with squamous cell cancer of the head and neck : - in combination with radiation therapy for locally advanced disease, - in combination with platinum-based chemotherapy for recurrent and/or metastatic disease.	29/06/2004	Merck KGaA
ERELZI	etanercept	Treatment of polyarthritis (rheumatoid factor positive or negative) and extended oligoarthritis in children and adolescents from the age of 2 years who have had an inadequate response to, or who have proved intolerant of, methotrexate. Treatment of psoriatic arthritis in adolescents from the age of 12 years who have had an inadequate response to, or who have proved intolerant of, methotrexate. Treatment of enthesitis-related arthritis in adolescents from the age of 12 years who have had an inadequate response to, or who have proved intolerant of, conventional therapy. Etanercept has not been studied in children aged less than 2 years.	23/06/2017	Sandoz GmbH
ERIVEDGE	vismodegib	Treatment of adult patients with symptomatic metastatic basal cell carcinoma Treatment of adult patients with locally advanced basal cell carcinoma inappropriate for surgery or radiotherapy	12/07/2013	Roche Registration Ltd
EURARTESIM	piperaquine tetraphosphate/dihydroartemisinin	Treatment of uncomplicated Plasmodium falciparum malaria in adults, children and infants 6 months and over and weighing 5 kg or more. Consideration should be given to official guidance on the appropriate use of antimalarial agents.	27/10/2011	Sigma-Tau Industrie Far- maceutiche Riunite S.p.A
EVOLTRA	clofarabine	Treatment of acute lymphoblastic leukaemia (ALL) in paediatric patients who have relapsed or are refractory after receiving at least two prior regimens and where there is no other treatment option anticipated to result in a durable response. Safety and efficacy have been assessed in studies of patients ≤ 21 years old at initial diagnosis.	29/05/2006	Genzyme Europe B.V.

Tradenname	Active Substance	Marketing Authorisation Indication	Marketing Authorisation Date (Dd/Mm/Yyyy)	Marketing Authorisation Holder
EXJADE	deferasirox	Treatment of chronic iron overload due to frequent blood transfusions (≥ 7 ml/kg/month of packed red blood cells) in patients with beta thalassaemia major aged 6 years and older. Treatment of chronic iron overload due to blood transfusions when deferoxamine therapy is contraindicated or inadequate in the following patient groups: -in paediatric patients with beta thalassaemia major with iron overload due to frequent blood transfusions (≥ 7 ml/kg/month of packed red blood cells) aged 2 to 5 years, -in adult and paediatric patients with beta thalassaemia major with iron overload due to infrequent blood transfusions (< 7 ml/kg/month of packed red blood cells) aged 2 years and older, -in adult and paediatric patients with other anaemias aged 2 years and older. Treatment of chronic iron overload requiring chelation therapy when deferoxamine therapy is contraindicated or inadequate in patients with non-transfusion-dependent thalassaemia syndromes aged 10 years and older.	01/09/2006	Novartis Europharm Limited
FABRAZYME	agalsidase beta	Long-term enzyme replacement therapy in patients with a confirmed diagnosis of Fabry disease (alphagalactosidase A deficiency) .	03/08/2001	Genzyme Europe B.V.
FERRIPROX	deferiprone	Treatment of iron overload in patients with thalassaemia major when deferoxamine therapy is contraindicated or inadequate.	25/08/1999	Apotex Europe B.V.
FILGRASTIM HEXAL	filgrastim	In patients, children or adults, with severe congenital, cyclic, or idiopathic neutropenia with an absolute neutrophil count (ANC) of $\leq 0.5 \times 10^9/l$, and a history of severe or recurrent infections. Long term administration of filgrastim is indicated to increase neutrophil counts and to reduce the incidence and duration of infection-related events.	06/02/2009	Hexal AG
FLEBOGAMMA DIF	human normal immunoglobulin	Replacement therapy in adults, and children and adolescents (2-18 years) in: - Primary immunodeficiency (PID) syndromes with impaired antibody production. - Hypogammaglobulinaemia and recurrent bacterial infections in patients with chronic lymphocytic leukaemia, in whom prophylactic antibiotics have failed. - Hypogammaglobulinaemia and recurrent bacterial infections in plateau phase multiple myeloma patients who have failed to respond to pneumococcal immunisation. - Hypogammaglobulinaemia in patients after allogeneic haematopoietic stem cell transplantation (HSCT). Immunomodulation in adults, and children and adolescents (2-18 years) in - Primary immune thrombocytopenia (ITP) , in patients at high risk of bleeding or prior to surgery to correct the platelet count. - Guillain-Barré syndrome - Kawasaki disease .	23/07/2007	Instituto Grifols S.A.

NEW

Tradename	Active Substance	Marketing Authorisation Indication	Marketing Authorisation Date (Dd/Mm/Yyyy)	Marketing Authorisation Holder
FOTIVDA	tivozanib hydrochloride monohydrate	First line treatment of adult patients with advanced renal cell carcinoma (RCC) and for adult patients who are VEGFR and mTOR pathway inhibitor-naïve following disease progression after one prior treatment with cytokine therapy for advanced RCC.	24/08/2017	EUSA Pharma (UK) Limited
GLIOLAN	5-aminolevulinic acid hydrochloride	In adult patients for visualisation of malignant tissue during surgery for malignant glioma (World Health Organization grade III and IV).	07/09/2007	Medac GmbH
GLIVEC	imatinib mesilate	Treatment of adult and paediatric patients with newly diagnosed Philadelphia chromosome (bcr-abl) positive (Ph+) chronic myeloid leukaemia (CML) for whom bone marrow transplantation is not considered as the first line of treatment. Treatment of adult and paediatric patients with Ph+ CML in chronic phase after failure of interferon-alpha therapy, or in accelerated phase or blast crisis. Treatment of adult and paediatric patients with newly diagnosed Philadelphia chromosome positive acute lymphoblastic leukaemia (Ph+ ALL) integrated with chemotherapy. Treatment of adult patients with relapsed or refractory Ph+ ALL as monotherapy. Treatment of adult patients with myelodysplastic / myeloproliferative diseases (MDS/MPD) associated with platelet-derived growth factor receptor (PDGFR) gene rearrangements. Treatment of adult patients with advanced hypereosinophilic syndrome (HES) and/or chronic eosinophilic leukaemia (CEL) with FIP1L1-PDGFR α rearrangement. The effect of Glivec on the outcome of bone marrow transplantation has not been determined. Treatment of adult patients with Kit (CD 117) positive unresectable and/or metastatic malignant gastrointestinal stromal tumours (GIST) . Adjuvant treatment of adult patients who are at significant risk of relapse following resection of Kit (CD117)-positive GIST . Patients who have a low or very low risk of recurrence should not receive adjuvant treatment. Treatment of adult patients with unresectable dermatofibrosarcoma protuberans (DFSP) and adult patients with recurrent and/or metastatic DFSP who are not eligible for surgery.	07/11/2001	Novartis Europharm Ltd
GONAL-F	follitropin alpha	Stimulation of spermatogenesis in men who have congenital or acquired hypogonadotropic hypogonadism with concomitant human Chorionic Gonadotrophin (hCG) therapy.	20/10/1995	Merck Serono Europe Ltd
GRASTOFIL	filgrastim	In adult or children patients with severe congenital, cyclic, or idiopathic neutropenia with an absolute neutrophil count (ANC) of $\leq 0.5 \times 10^9/L$, and a history of severe or recurrent infections, long term administration of Grastofil is indicated to increase neutrophil counts and to reduce the incidence and duration of infection-related events.	18/10/2013	Apotex Europe B.V.
HALAVEN	eribulin	Treatment of adult patients with unresectable liposarcoma who have received prior anthracycline containing therapy (unless unsuitable) for advanced or metastatic disease.	17/03/2011	Eisai Europe Ltd

NEW

Tradename	Active Substance	Marketing Authorisation Indication	Marketing Authorisation Date (Dd/Mm/Yyyy)	Marketing Authorisation Holder
HELIXATE NEXGEN	octocog alpha	Treatment and prophylaxis of bleeding in patients with haemophilia A (congenital factor VIII deficiency) . This preparation does not contain von Willebrand factor and is therefore not indicated in von Willebrand's disease.	04/08/2000	Bayer Pharma AG
HERCEPTIN	trastuzumab	In combination with capecitabine or 5-fluorouracil and cisplatin, treatment of patients with HER2-positive metastatic adenocarcinoma of the stomach or gastroesophageal junction who have not received prior anticancer treatment for their metastatic disease. Herceptin should only be used in patients with metastatic gastric cancer whose tumours have HER2 overexpression as defined by IHC2+ and a confirmatory SISH or FISH result, or by an IHC3+ result. Accurate and validated assay methods should be used.	28/08/2000	Roche Registration Ltd
HIZENTRA	human normal immunoglobulin (scig)	Replacement therapy in adults and children in primary immunodeficiency syndromes such as: - congenital agammaglobulinaemia and hypogammaglobulinaemia , - common variable immunodeficiency , - severe combined immunodeficiency , - IgG subclass deficiencies with recurrent infections. Replacement therapy in myeloma or chronic lymphocytic leukaemia with severe secondary hypogammaglobulinaemia and recurrent infections.	14/04/2011	CSL Behring GmbH
HUMIRA	adalimumab	In combination with methotrexate is indicated for the treatment of active polyarticular juvenile idiopathic arthritis , in children and adolescents aged 2 to 17 years who have had an inadequate response to one or more disease-modifying antirheumatic drugs (DMARDs). As monotherapy in case of intolerance to methotrexate or when continued treatment with methotrexate is inappropriate. It has not been studied in children aged less than 2 years. Treatment of active enthesitis-related arthritis in patients, 6 years of age and older, who have had an inadequate response to, or who are intolerant of, conventional therapy. Treatment of non-infectious intermediate, posterior and panuveitis in adult patients who have had an inadequate response to corticosteroids, in patients in need of corticosteroid-sparing, or in whom corticosteroid treatment is inappropriate. Treatment of paediatric chronic non-infectious anterior uveitis in patients from 2 years of age who have had an inadequate response to or are intolerant to conventional therapy, or in whom conventional therapy is inappropriate.	08/09/2003	Abbvie Ltd.
HYCAMTIN	topotecan	As monotherapy, treatment of: - patients with metastatic carcinoma of the ovary after failure of first-line or subsequent therapy. - patients with relapsed small cell lung cancer (SCLC) for whom retreatment with the first-line regimen is not considered appropriate.	12/11/1996	Novartis Europharm Ltd

Tradename	Active Substance	Marketing Authorisation Indication	Marketing Authorisation Date (Dd/Mm/Yyyy)	Marketing Authorisation Holder
HYQVIA	human normal immunoglobulin	Replacement therapy in adults (≥ 18 years) in primary immunodeficiency syndromes such as: - congenital agammaglobulinaemia and hypogammaglobulinaemia - common variable immunodeficiency - severe combined immunodeficiency - IgG subclass deficiencies with recurrent infections. Replacement therapy in adults (≥ 18 years) in myeloma or chronic lymphocytic leukaemia with severe secondary hypogammaglobulinaemia and recurrent infections.	16/05/2013	Baxalta Innovations GmbH
IBLIAS	octocog alfa	Treatment and prophylaxis of bleeding in patients with haemophilia A (congenital factor VIII deficiency) . Iblias can be used for all age groups.	18/02/2016	Bayer Pharma AG
ILARIS	canakinumab	Treatment of Cryopyrin-Associated Periodic Syndromes (CAPS) in adults, adolescents and children aged 2 years and older with body weight of 7,5 kg or above, including: - Muckle-Wells Syndrome (MWS) , - Neonatal-Onset Multisystem Inflammatory Disease (NOMID) / Chronic Infantile Neurological, Cutaneous, Articular Syndrome (CINCA) , - Severe forms of Familial Cold Autoinflammatory Syndrome (FCAS) / Familial Cold Urticaria (FCU) presenting with signs and symptoms beyond cold- induced urticarial skin rash. Treatment of active Still's disease including Adult-Onset Still's Disease (AOSD) and Systemic Juvenile Idiopathic Arthritis (SJIA) in patients aged 2 years and older who have responded inadequately to previous therapy with non-steroidal anti-inflammatory drugs (NSAIDs) and systemic corticosteroids. Ilaris can be given as monotherapy or in combination with methotrexate.	23/10/2009	Novartis Europharm Ltd

NEW

Tradename	Active Substance	Marketing Authorisation Indication	Marketing Authorisation Date (Dd/Mm/Yyyy)	Marketing Authorisation Holder
IMATINIB TEVA B.V.	imatinib	<p>Treatment of paediatric patients with newly diagnosed Philadelphia chromosome (bcr-abl) positive (Ph+) chronic myeloid leukaemia (CML) for whom bone marrow transplantation is not considered as the first line of treatment.</p> <p>Treatment of paediatric patients with Ph+ CML in chronic phase after failure of interferon-alpha therapy, or in accelerated phase or blast crisis.</p> <p>Treatment of adult patients with Ph+ CML in blast crisis.</p> <p>Treatment of adult and paediatric patients with newly diagnosed Philadelphia chromosome positive acute lymphoblastic leukaemia (Ph+ ALL) integrated with chemotherapy.</p> <p>Treatment of adult patients with relapsed or refractory Ph+ ALL as monotherapy.</p> <p>Treatment of adult patients with myelodysplastic/myeloproliferative diseases (MDS/MPD) associated with platelet-derived growth factor receptor (PDGFR) gene re-arrangements.</p> <p>Treatment of adult patients with advanced hypereosinophilic syndrome (HES) and/or chronic eosinophilic leukaemia (CEL) with FIP1L1-PDGFRα rearrangement.</p> <p>The effect of imatinib on the outcome of bone marrow transplantation has not been determined.</p> <p>Treatment of adult patients with Kit (CD 117) positive unresectable and/or metastatic malignant gastrointestinal stromal tumours (GIST).</p> <p>Adjuvant treatment of adult patients who are at significant risk of relapse following resection of Kit (CD117)-positive GIST. Patients who have a low or very low risk of recurrence should not receive adjuvant treatment.</p> <p>Treatment of adult patients with unresectable dermatofibrosarcoma protuberans (DFSP) and adult patients with recurrent and/or metastatic DFSP who are not eligible for surgery.</p>	15/11/2017	Teva B.V.
IMRALDI	adalimumab	<p>In combination with methotrexate indicated for the treatment of active polyarticular juvenile idiopathic arthritis, in patients from the age of 2 years who have had an inadequate response to one or more disease-modifying anti-rheumatic drugs (DMARDs). Imraldi can be given as monotherapy in case of intolerance to methotrexate or when continued treatment with methotrexate is inappropriate. Adalimumab has not been studied in patients aged less than 2 years.</p> <p>Treatment of active enthesitis-related arthritis in patients, 6 years of age and older, who have had an inadequate response to, or who are intolerant of, conventional therapy.</p> <p>Treatment of non-infectious intermediate, posterior and panuveitis in adult patients who have had an inadequate response to corticosteroids, in patients in need of corticosteroid-sparing, or in whom corticosteroid treatment is inappropriate.</p>	24/08/2017	Samsung Bioepis UK Limited (SBUK)

Tradename	Active Substance	Marketing Authorisation Indication	Marketing Authorisation Date (Dd/Mm/Yyyy)	Marketing Authorisation Holder
INCRELEX	mecasermin	<p>For the long-term treatment of growth failure in children and adolescents from 2 to 18 years with severe primary insulin-like growth factor-1 deficiency (Primary IGFD).</p> <p>Severe Primary IGFD is defined by:</p> <ul style="list-style-type: none"> - height standard deviation score \leq -3.0 and - basal IGF-1 levels below the 2.5th percentile for age and gender and - GH sufficiency - exclusion of secondary forms of IGF-1 deficiency, such as malnutrition, hypothyroidism, or chronic treatment with pharmacologic doses of anti-inflammatory steroids. <p>Severe Primary IGFD includes patients with mutations in the GH receptor (GHR), post-GHR signaling pathway, and IGF-1 gene defects; they are not GH deficient, and therefore, they cannot be expected to respond adequately to exogenous GH treatment. It is recommended to confirm the diagnosis by conducting an IGF-1 generation test.</p>	03/08/2007	Ipsen Pharma
INLYTA	axitinib	Treatment of adult patients with advanced renal cell carcinoma (RCC) after failure of prior treatment with sunitinib or a cytokine.	03/09/2012	Pfizer Ltd
INOMAX	nitric oxide	<p>In conjunction with ventilatory support and other appropriate active substances:</p> <ul style="list-style-type: none"> - for the treatment of newborn infants \geq 34 weeks gestation with hypoxic respiratory failure associated with clinical or echo cardiographic evidence of pulmonary hypertension, in order to improve oxygenation and to reduce the need for extracorporeal membrane oxygenation. - as part of the treatment of peri- and post-operative pulmonary hypertension in adults and newborn infants, infants and toddlers, children and adolescents, ages 0-17 years in conjunction to heart surgery, in order to selectively decrease pulmonary arterial pressure and improve right ventricular function and oxygenation. 	01/08/2001	Linde Healthcare AB

Tradename	Active Substance	Marketing Authorisation Indication	Marketing Authorisation Date (Dd/Mm/Yyyy)	Marketing Authorisation Holder
INTRONA	interferon alfa-2b	<p>Treatment of patients with hairy cell leukaemia. As Monotherapy for the treatment of adult patients with Philadelphia chromosome or bcr/abl translocation positive chronic myelogenous leukaemia. Clinical experience indicates that a haematological and cytogenetic major/minor response is obtainable in the majority of patients treated. A major cytogenetic response is defined by < 34 % Ph+ leukaemic cells in the bone marrow, whereas a minor response is > 34 %, but < 90 % Ph+ cells in the marrow.</p> <p>In combination with interferon alfa-2b and cytarabine (Ara-C) during the first 12 months of treatment it has been demonstrated to significantly increase the rate of major cytogenetic responses and to significantly prolong the overall survival at three years when compared to interferon alfa-2b monotherapy.</p> <p>As maintenance therapy in patients with multiple myeloma who have achieved objective remission (more than 50 % reduction in myeloma protein) following initial induction chemotherapy. Current clinical experience indicates that maintenance therapy with interferon alfa-2b prolongs the plateau phase; however, effects on overall survival have not been conclusively demonstrated.</p> <p>Treatment of high tumour burden follicular lymphoma as adjunct to appropriate combination induction chemotherapy such as a CHOP-like regimen. High tumour burden is defined as having at least one of the following: bulky tumour mass (> 7 cm), involvement of three or more nodal sites (each > 3 cm), systemic symptoms (weight loss > 10 %, pyrexia > 38°C for more than 8 days, or nocturnal sweats), splenomegaly beyond the umbilicus, major organ obstruction or compression syndrome, orbital or epidural involvement, serous effusion, or leukaemia.</p> <p>Treatment of carcinoid tumours with lymph node or liver metastases and with "carcinoid syndrome".</p>	09/03/2000	Merck Sharp & Dohme Ltd
IXIARO	japanese encephalitis vaccine (inactivated, adsorbed)	<p>Active immunisation against Japanese encephalitis in adults, adolescents, children and infants aged 2 months and older.</p> <p>IXIARO should be considered for use in individuals at risk of exposure through travel or in the course of their occupation</p>	31/03/2009	Valneva Austria GmbH
JAKAVI	ruxolitinib	<p>Treatment of disease-related splenomegaly or symptoms in adult patients with primary myelofibrosis (also known as chronic idiopathic myelofibrosis), post-polycythaemia-vera myelofibrosis or post-essential-thrombocythaemia myelofibrosis.</p> <p>Treatment of adult patients with polycythaemia vera who are resistant to or intolerant of hydroxyurea.</p>	23/08/2012	Novartis Europharm Ltd

Tradename	Active Substance	Marketing Authorisation Indication	Marketing Authorisation Date (Dd/Mm/Yyyy)	Marketing Authorisation Holder
KEPPRA	levetiracetam	As monotherapy in the treatment of partial onset seizures with or without secondary generalisation in patients from 16 years of age with newly diagnosed epilepsy . As adjunctive therapy in the treatment of partial onset seizures with or without secondary generalisation in adults, children and infants from 1 month of age with epilepsy; in the treatment of myoclonic seizures in adults and adolescents from 12 years of age with Juvenile Myoclonic Epilepsy Treatment of primary generalised tonic-clonic seizures in adults and adolescents from 12 years of age with Idiopathic Generalised Epilepsy .	29/09/2000	UCB Pharma SA
KEYTRUDA	pembrolizumab	As monotherapy for the treatment of adult patients with relapsed or refractory classical Hodgkin lymphoma (cHL) who have failed autologous stem cell transplant (ASCT) and brentuximab vedotin (BV), or who are transplant-ineligible and have failed BV.	17/07/2015	Merck Sharp & Dohme Limited
KINERET	anakinra	Treatment in adults, adolescents, children and infants aged 8 months and older with a body weight of 10 kg or above of Cryopyrin-Associated Periodic Syndromes (CAPS) , including:- Neonatal-Onset Multisystem Inflammatory Disease (NOMID) / Chronic Infantile Neurological, Cutaneous, Articular Syndrome (CINCA) ,- Muckle-Wells Syndrome (MWS) ,- Familial Cold Autoinflammatory Syndrome (FCAS) .	08/03/2002	Swedish Orphan Biovitrum AB
KIOVIG	human normal immunoglobulin	Replacement therapy in adults, and children and adolescents (0-18 years) in: - Primary immunodeficiency syndromes with impaired antibody production, - Hypogammaglobulinaemia and recurrent bacterial infections in patients with chronic lymphocytic leukaemia , in whom prophylactic antibiotics have failed - Hypogammaglobulinaemia and recurrent bacterial infections in plateau phase multiple myeloma patients who have failed to respond to pneumococcal immunisation, - Hypogammaglobulinaemia in patients after allogeneic haematopoietic stem cell transplantation (HSCT) . - Congenital AIDS and recurrent bacterial infections. Immunomodulation in adults, and children and adolescents (0-18 years) in: - Primary immune thrombocytopenia (ITP) , in patients at high risk of bleeding or prior to surgery to correct the platelet count - Guillain Barré syndrome - Kawasaki disease - Multifocal Motor Neuropathy (MMN) .	19/01/2006	Baxter AG
KISPLYX	lenvatinib	in combination with everolimus for the treatment of adult patients with advanced renal cell carcinoma (RCC) following one prior vascular endothelial growth factor (VEGF)-targeted therapy.	25/08/2016	Eisai Europe Ltd

Tradename	Active Substance	Marketing Authorisation Indication	Marketing Authorisation Date (Dd/Mm/Yyyy)	Marketing Authorisation Holder
KOGENATE BAYER	octocog alpha	Treatment and prophylaxis of bleeding in patients with haemophilia A (congenital factor VIII deficiency) . This preparation does not contain von Willebrand factor and is therefore not indicated in von Willebrand's disease. This product is indicated for adults, adolescents and children of all ages.	04/08/2000	Bayer Pharma AG
KOVALTRY	octocog alfa	Treatment and prophylaxis of bleeding in patients with haemophilia A (congenital factor VIII deficiency) . Kovaltry can be used for all age groups.	18/02/2016	Bayer Pharma AG
LITAK	cladribine	Treatment of hairy cell leukaemia .	14/04/2004	Lipomed GmbH
LOJUXTA	lomitapide	Adjunct to a low-fat diet and other lipid-lowering medicinal products with or without low density lipoprotein (LDL) apheresis in adult patients with homozygous familial hypercholesterolaemia (HoFH) . Genetic confirmation of HoFH should be obtained whenever possible. Other forms of primary hyperlipoproteinemia and secondary causes of hypercholesterolaemia (e.g., nephrotic syndrome, hypothyroidism) must be excluded.	31/07/2013	Aegerion Pharmaceuticals SAS
LYSODREN	mitotane	Symptomatic treatment of advanced (unresectable, metastatic or relapsed) adrenal cortical carcinoma . The effect of Lysodren on non functional adrenal cortical carcinoma is not established.	28/04/2004	Laboratoire HRA Pharma
MABTHERA	rituximab	Non-Hodgkin's lymphoma (NHL) - Treatment of previously untreated patients with stage III-IV follicular lymphoma in combination with chemotherapy. - As maintenance therapy, the treatment of follicular lymphoma patients responding to induction therapy. - In monotherapy, treatment of patients with stage III-IV follicular lymphoma who are chemoresistant or are in their second or subsequent relapse after chemotherapy. - Treatment of patients with CD20 positive diffuse large B cell non- Hodgkin's lymphoma in combination with CHOP (cyclophosphamide, doxorubicin, vincristine, prednisolone) chemotherapy. In combination with chemotherapy, treatment of patients with previously untreated and relapsed/refractory chronic lymphocytic leukaemia . Only limited data are available on efficacy and safety for patients previously treated with monoclonal antibodies including MabThera or patients refractory to previous MabThera plus chemotherapy. Granulomatosis with polyangiitis and Microscopic polyangiitis in combination with glucocorticoids, it is indicated for the induction of remission in adult patients with severe, active Granulomatosis with polyangiitis (Wegener's) (GPA) and Microscopic polyangiitis (MPA).	02/06/1998	Roche Registration Ltd
MIGLUSTAT GEN ORPH	miglustat	Oral treatment of adult patients with mild to moderate type 1 Gaucher disease . Miglustat Gen.Orph may be used only in the treatment of patients for whom enzyme replacement therapy is unsuitable.	10/11/2017	Gen.Orph

Tradename	Active Substance	Marketing Authorisation Indication	Marketing Authorisation Date (Dd/Mm/Yyyy)	Marketing Authorisation Holder
MYOZYME	alglucosidase alpha	Long-term enzyme replacement therapy (ERT) in patients with a confirmed diagnosis of Pompe disease (acid α-glucosidase deficiency) . Myozyme is indicated in adults and paediatric patients of all ages	29/03/2006	Genzyme Europe B.V.
MYSILDECARD	sildenafil	Treatment of adult patients with pulmonary arterial hypertension classified as WHO functional class II and III, to improve exercise capacity. Efficacy has been shown in primary pulmonary hypertension and pulmonary hypertension associated with connective tissue disease. Treatment of paediatric patients aged 1 year to 17 years old with pulmonary arterial hypertension. Efficacy in terms of improvement of exercise capacity or pulmonary haemodynamics has been shown in primary pulmonary hypertension and pulmonary hypertension associated with congenital heart disease.	15/09/2016	MYLAN S.A.S.
NAGLAZYME	galsulfase	Long-term enzyme replacement therapy in patients with a confirmed diagnosis of Mucopolysaccharidosis VI (MPS VI; N-acetylgalactosamine 4-sulfatase deficiency; Maroteaux-Lamy syndrome) A key issue is to treat children aged <5 years suffering from a severe form of the disease, even though children <5 years were not included in the pivotal phase 3 study. Limited data are available in patients < 1 year of age.	24/01/2006	BioMarin Europe Ltd
NEOFORDEX	dexamethasone	Indicated in adults for the treatment of symptomatic multiple myeloma in combination with other medicinal products.	16/03/2016	Laboratoires CTRS
NEXAVAR	sorafenib tosylate	Treatment of patients with advanced renal cell carcinoma who have failed prior interferon-alpha or interleukin-2 based therapy or are considered unsuitable for such therapy.	19/07/2006	Bayer Pharma AG
NITISINONE MDK (previously NITISINONE MENDELIKABS)	nitisinone	Treatment of adult and paediatric (in any age range) patients with confirmed diagnosis of hereditary tyrosinemia type 1 (HT 1) in combination with dietary restriction of tyrosine and phenylalanine.	24/08/2017	MendeliKABS Europe Ltd
NIVESTIM	filgrastim	In patients, children or adults, with severe congenital, cyclic, or idiopathic neutropenia with an absolute neutrophil count (ANC) of $\leq 0.5 \times 10^9/l$, and a history of severe or recurrent infections.	08/06/2010	Hospira UK Ltd
NONAFACT	human coagulation factor IX	Treatment and prophylaxis of bleeding in patients with haemophilia B (congenital factor IX deficiency) .	03/07/2001	Sanquin
NORDIMET	methotrexate	Treatment of polyarthritic forms of severe, active juvenile idiopathic arthritis (JIA) , when the response to nonsteroidal anti-inflammatory drugs (NSAIDs) has been inadequate.	18/08/2016	Nordic Group B.V.
NOVOEIGHT	turoctocog alpha	Treatment and prophylaxis of bleeding in patients with haemophilia A (congenital factor VIII deficiency) . NovoEight can be used for all age groups.	13/11/2013	Novo Nordisk A/S

Tradename	Active Substance	Marketing Authorisation Indication	Marketing Authorisation Date (Dd/Mm/Yyyy)	Marketing Authorisation Holder
NOVOSEVEN	eptacog alpha (activated)	Treatment of bleeding episodes and for the prevention of bleeding in those undergoing surgery or invasive procedures in the following patient groups : -patients with congenital haemophilia with inhibitors to coagulation factors VIII or IX > 5 BU -patients with congenital haemophilia who are expected to have a high anamnestic response to factor VIII or factor IX administration -patients with acquired haemophilia -patients with congenital FVII deficiency ; -patients with Glanzmann's thrombasthenia with antibodies toGP IIb - IIIa and/or HLA, and with past or present refractoriness to platelet transfusions.	23/02/1996	Novo Nordisk A/S
NOVOTHIRTEEN	catridecacog	Long term prophylactic treatment of bleeding in in adult and paediatric patients with congenital factor XIII A-subunit deficiency	03/09/2012	Novo Nordisk A/S
NOXAFIL	posaconazole	Treatment of the fungal infections in adults: - Invasive aspergillosis in patients with disease that is refractory to amphotericin B or itraconazole or in patients who are intolerant of these medicinal products - Fusariosis in patients with disease that is refractory to amphotericin B or in patients who are intolerant of amphotericin B. - Chromoblastomycosis and mycetoma in patients with disease that is refractory to itraconazole or in patients who are intolerant of itraconazole - Coccidioidomycosis in patients with disease that is refractory to amphotericin B, itraconazole or fluconazole or in patients who are intolerant of these medicinal products. Refractoriness is defined as progression of infection or failure to improve after a minimum of 7 days of prior therapeutic doses of effective antifungal therapy. Prophylaxis of invasive fungal infections in : - Patients receiving remission-induction chemotherapy for acute myelogenous leukemia (AML) or myelodysplastic syndromes (MDS) expected to result in prolonged neutropenia and who are at high risk of developing invasive fungal infections - Hematopoietic stem cell transplant (HSCT) recipients who are undergoing high-dose immunosuppressive therapy for graft versus host disease and who are at high risk of developing invasive fungal infections.	25/10/2005	Merck Sharp & Dohme Ltd
OBIZUR	susoctocog alfa	Treatment of bleeding episodes in patients with acquired haemophilia caused by antibodies to Factor VIII.	11/11/2015	Baxalta Innovations GmbH

Tradename	Active Substance	Marketing Authorisation Indication	Marketing Authorisation Date (Dd/Mm/Yyyy)	Marketing Authorisation Holder
OMNITROPE	somatropin	<p>Infants, children and adolescents:</p> <ul style="list-style-type: none"> - Growth disturbance due to insufficient secretion of growth hormone (growth hormone deficiency, GHD). - Growth disturbance associated with Turner syndrome. - Growth disturbance associated with chronic renal insufficiency. - Growth disturbance (current height standard deviation score (SDS) < -2.5 and parental adjusted height SDS < -1) in short children/adolescents born small for gestational age (SGA), with a birth weight and/or length below -2 standard deviation (SD), who failed to show catch-up growth (height velocity (HV) SDS < 0 during the last year) by 4 years of age or later. - Prader-Willi syndrome (PWS), for improvement of growth and body composition. The diagnosis of PWS should be confirmed by appropriate genetic testing. <p>Adults</p> <ul style="list-style-type: none"> - Replacement therapy in adults with pronounced growth hormone deficiency. - <i>Adult onset</i>: Patients who have severe growth hormone deficiency associated with multiple hormone deficiencies as a result of known hypothalamic or pituitary pathology, and who have at least one known deficiency of a pituitary hormone not being prolactin. These patients should undergo an appropriate dynamic test in order to diagnose or exclude a growth hormone deficiency. - <i>Childhood onset</i>: Patients who were growth hormone deficient during childhood as a result of congenital, genetic, acquired, or idiopathic causes. Patients with childhood onset GHD should be re-evaluated for growth hormone secretory capacity after completion of longitudinal growth. In patients with a high likelihood for persistent GHD, i.e. a congenital cause or GHD secondary to a hypothalamic-pituitary disease or insult, an insulin-like growth factor-I (IGF-I) SDS < -2 off growth hormone treatment for at least 4 weeks should be considered sufficient evidence of profound GHD. All other patients will require IGF-I assay and one growth hormone stimulation test. 	12/04/2006	Sandoz GmbH
ONCASPAR	pegaspargase	Indicated as a component of antineoplastic combination therapy in acute lymphoblastic leukaemia (ALL) in paediatric patients from birth to 18 years, and adult patients.	14/01/2016	Baxalta Innovations GmbH
OPDIVO	nivolumab	<p>As monotherapy indicated for the treatment of advanced renal cell carcinoma after prior therapy in adults.</p> <p>As monotherapy for the treatment of adult patients with relapsed or refractory classical Hodgkin lymphoma after autologous stem cell transplant (ASCT) and treatment with brentuximab vedotin.</p> <p>As monotherapy for the treatment of squamous cell cancer of the head and neck in adults progressing on or after platinum-based therapy.</p>	19/06/2015	Bristol-Myers Squibb Pharma EEIG

Tradename	Active Substance	Marketing Authorisation Indication	Marketing Authorisation Date (Dd/Mm/Yyyy)	Marketing Authorisation Holder
ORENCIA	abatacept	In combination with methotrexate, for the treatment of moderate to severe active polyarticular juvenile idiopathic arthritis (JIA) in paediatric patients 6 years of age and older who have had an insufficient response to other DMARDs including at least one TNF inhibitor.	21/05/2007	Bristol-Myers SquibbPharma EEIG
ORFADIN	nitisinone	Treatment of adult and paediatric (in any age range) patients with confirmed diagnosis of hereditary tyrosinemia type 1 (HT-1) in combination with dietary restriction of tyrosine and phenylalanine.	21/02/2005	Swedish Orphan Biovitrum AB
ORKAMBI	lumacaftor / ivacaftor	Treatment of cystic fibrosis (CF) in patients aged 12 years and older who are homozygous for the F508del mutation in the CFTR gene	19/11/2015	Vertex Pharmaceuticals (Europe) Ltd
OVALEAP	follitropin alpha	Indicated for the stimulation of spermatogenesis in adult men who have congenital or acquired hypogonadotropic hypogonadism with concomitant human chorionic gonadotropin (hCG) therapy.	27/09/2013	Teva Pharmaceuticals Europe B.V.
OZURDEX	dexamethasone	For the treatment of adult patients with inflammation of the posterior segment of the eye presenting as non-infectious uveitis .	27/07/2010	Allergan Pharmaceuticals Ireland
PANRETIN	alitretinoin	Topical treatment of cutaneous lesions in patients with AIDS-related Kaposi's sarcoma (KS) : - when lesions are not ulcerated or lymphoedematous, and -treatment of visceral KS is not required, and -when lesions are not responding to systemic antiretroviral therapy, and -radiotherapy or chemotherapy are not appropriate.	11/10/2000	Eisai Ltd
PEDEA	ibuprofen	Treatment of a haemodynamically significant patent ductus arteriosus in preterm newborn infants less than 34 weeks of gestational age.	29/07/2004	Orphan Europe S.A.R.L
PEMETREXED ACCORD	pemetrexed	In combination with cisplatin is indicated for the treatment of chemotherapy naïve patients with unresectable malignant pleural mesothelioma .	18/01/2016	Accord Healthcare Ltd
PEMETREXED FRESENIUS KABI	pemetrexed	In combination with cisplatin is indicated for the treatment of chemotherapy naïve patients with unresectable malignant pleural mesothelioma .	22/07/2016	Fresenius Kabi Oncology Plc
PEMETREXED HOSPIRA	pemetrexed	In combination with cisplatin indicated for the treatment of chemotherapy naïve patients with unresectable malignant pleural mesothelioma .	20/11/2015	Hospira UK Ltd
PEMETREXED LILLY	pemetrexed	In combination with cisplatin for the treatment of chemotherapy naïve patients with unresectable malignant pleural mesothelioma .	14/09/2015	Eli Lilly Netherlands
PEMETREXED MEDAC	pemetrexed	In combination with cisplatin indicated for the treatment of chemotherapy naïve patients with unresectable malignant pleural mesothelioma .	27/11/2015	Medac GmbH
PEMETREXED SANDOZ	pemetrexed	In combination with cisplatin for the treatment of chemotherapy naïve patients with unresectable malignant pleural mesothelioma .	18/09/2015	Sandoz GmbH
PIXUVRI	pixantrone dimaleate	As monotherapy for the treatment of adult patients with multiply relapsed or refractory aggressive Non-Hodgkin B-cell Lymphomas (NHL) . The benefit of pixantrone treatment has not been established in patients when used as fifth line or greater chemotherapy in patients who are refractory to last therapy.	10/05/2012	CTI life Sciences Ltd

Tradenname	Active Substance	Marketing Authorisation Indication	Marketing Authorisation Date (Dd/Mm/Yyyy)	Marketing Authorisation Holder
PRIVIGEN	human normal immunoglobulin (IVIg)	Replacement therapy in adults, and children and adolescents (0-18 years) in: - Primary immunodeficiency (PID) syndromes with impaired antibody production - Hypogammaglobulinaemia and recurrent bacterial infections in patients with chronic lymphocytic leukaemia , in whom prophylactic antibiotics have failed. - Hypogammaglobulinaemia and recurrent bacterial infections in plateau phase multiple myeloma patients who have failed to respond to pneumococcal immunisation. - Hypogammaglobulinaemia in patients after allogeneic haematopoietic stem cell transplantation (HSCT) . - Congenital AIDS with recurrent bacterial infections. Immunomodulation in adults, and children and adolescents (0-18 years) in: - Primary immune thrombocytopenia (ITP) , in patients at high risk of bleeding or prior to surgery to correct the platelet count. - Guillain-Barré syndrome . - Kawasaki disease . - Chronic inflammatory demyelinating polyneuropathy (CIDP) . Only limited experience is available of use of intravenous immunoglobulins in children with CIDP.	25/04/2008	CSL Behring GmbH
PUREGON	follitropin beta	Indicated in adult males with deficient spermatogenesis due to hypogonadotrophic hypogonadism .	03/05/1996	Merck Sharp & Dohme Ltd
QUINSAIR	levofloxacin	Management of chronic pulmonary infections due to <i>Pseudomonas aeruginosa</i> in adult patients with cystic fibrosis	26/03/2015	Aptalis Pharma SAS
RATIOGRASTIM	filgrastim	In patients, children or adults, with severe congenital, cyclic, or idiopathic neutropenia with an absolute neutrophil count (ANC) of $\leq 0.5 \times 10^9/l$, and a history of severe or recurrent infections.	15/09/2008	Ratiopharm GmbH
REFACTO AF	moroctocog alpha	Treatment and prophylaxis of bleeding in patients with haemophilia A (congenital factor VIII deficiency) in adults and children of all ages, including newborns. ReFacto AF is appropriate for use in adults and children of all ages, including newborns. ReFacto AF does not contain von Willebrand factor, and hence is not indicated in von Willebrand's disease.	13/04/1999	Pfizer Ltd
REFIXIA	nonacog beta pegol	Treatment and prophylaxis of bleeding in patients 12 years and above with haemophilia B (congenital factor IX deficiency) .	02/06/2017	Novo Nordisk A/S
REPATHA	evolocumab	Indicated in adults and adolescents aged 12 years and over with homozygous familial hypercholesterolaemia in combination with other lipid-lowering therapies. The effect of Repatha on cardiovascular morbidity and mortality has not yet been determined.	17/07/2015	Amgen Europe B.V.
REPLAGAL	agalsidase alfa	Long-term enzyme replacement therapy in patients with a confirmed diagnosis of Fabry disease (alpha-galactosidase A deficiency)	03/08/2001	Shire Human Genetic Therapies AB

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RESPREEZA	human alpha1-proteinase inhibitor	For maintenance treatment, to slow the progression of emphysema in adults with documented severe alpha1-proteinase inhibitor deficiency (e.g.genotypes PiZZ, PiZ(null), Pi(null,null), PiSZ). Patients are to be under optimal pharmacologic and non-pharmacologic treatment and show evidence of progressive lung disease (e.g.lower forced expiratory volume per second (FEV1) predicted, impaired walking capacity or increased number of exacerbations) as evaluated by a healthcare professional experienced in the treatment of alpha1-proteinase inhibitor deficiency.	20/08/2015	CSL Behring GmbH
REVATIO	sildenafil citrate	Treatment of adult patients with pulmonary arterial hypertension classified as WHO functional class II and III, to improve exercise capacity. Efficacy has been shown in primary pulmonary hypertension and pulmonary hypertension associated with connective tissue disease. Treatment of paediatric patients aged 1 year to 17 years old with pulmonary arterial hypertension. Efficacy in terms of improvement of exercise capacity or pulmonary haemodynamics has been shown in primary pulmonary hypertension and pulmonary hypertension associated with congenital heart disease.	28/10/2005	Pfizer Ltd
REVLIMID	lenalidomide	As monotherapy for the maintenance treatment of adult patients with newly diagnosed multiple myeloma who have undergone autologous stem cell transplantation. As combination therapy for the treatment of adult patients with previously untreated multiple myeloma who are not eligible for transplant. Treatment in combination with dexamethasone of multiple myeloma in adult patients who have received at least one prior therapy.	14/06/2007	Celgene Europe Ltd
REVOLADE	eltrombopag	Indicated for chronic immune (idiopathic) thrombocytopenic purpura (ITP) patients aged 1 year and above who are refractory to other treatments. Indicated in adult patients with acquired severe aplastic anaemia (SAA) who were either refractory to prior immunosuppressive therapy or heavily pretreated and are unsuitable for haematopoietic stem cell transplantation.	11/03/2010	Novartis Europharm Ltd
RILUTEK	riluzole	To extend life or the time to mechanical ventilation for patients with amyotrophic lateral sclerosis (ALS) . Clinical trials have demonstrated that RILUTEK extends survival for patients with ALS.Survival was defined as patients who were alive, not intubated for mechanical ventilation and tracheotomy-free. There is no evidence that RILUTEK exerts a therapeutic effect on motor function, lung function, fasciculations, muscle strength and motor symptoms. RILUTEK has not been shown to be effective in the late stages of ALS. Safety and efficacy of RILUTEK has only been studied in ALS. Therefore, RILUTEK should not be used in patients with any other form of motor neurone disease.	10/06/1996	Aventis Pharma S.A.

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RITEMVIA	rituximab	<p>Treatment of previously untreated patients with stage III, IV follicular lymphoma in combination with chemotherapy.</p> <p>As maintenance therapy for the treatment of follicular lymphoma patients responding to induction therapy.</p> <p>As monotherapy for the treatment of patients with stage III, IV follicular lymphoma who are chemo resistant or are in their second or subsequent relapse after chemotherapy.</p> <p>Treatment of patients with CD20 positive diffuse large B cell non Hodgkin's lymphoma in combination with CHOP (cyclophosphamide, doxorubicin, vincristine, prednisolone) chemotherapy.</p> <p>In combination with glucocorticoids, is indicated for the induction of remission in adult patients with severe, active granulomatosis with polyangiitis (Wegener's) (GPA) and microscopic polyangiitis (MPA).</p>	13/07/2017	Celltrion Healthcare Hungary Kft
RITUZENA (previously TUXELLA)	rituximab	<p>Treatment of previously untreated patients with stage III IV follicular lymphoma in combination with chemotherapy.</p> <p>As monotherapy indicated for treatment of patients with stage III IV follicular lymphoma who are chemo resistant or are in their second or subsequent relapse after chemotherapy.</p> <p>Treatment of patients with CD20 positive diffuse large B cell non Hodgkin's lymphoma in combination with CHOP (cyclophosphamide, doxorubicin, vincristine, prednisolone) chemotherapy.</p> <p>In combination with chemotherapy for the treatment of patients with previously untreated and relapsed/refractory Chronic lymphocytic leukaemia (CLL). Only limited data are available on efficacy and safety for patients previously treated with monoclonal antibodies including Rituzena or patients refractory to previous Rituzena plus chemotherapy.</p>	13/07/2017	Celltrion Healthcare Hungary Kft

Tradename	Active Substance	Marketing Authorisation Indication	Marketing Authorisation Date (Dd/Mm/Yyyy)	Marketing Authorisation Holder
RIXATHON	rituximab	<p>Treatment of previously untreated patients with stage III-IV follicular lymphoma in combination with chemotherapy.</p> <p>As maintenance therapy for the treatment of follicular lymphoma patients responding to induction therapy.</p> <p>As monotherapy for the treatment of patients with stage III-IV follicular lymphoma who are chemoresistant or are in their second or subsequent relapse after chemotherapy.</p> <p>For the treatment of patients with CD20 positive diffuse large B cell non Hodgkin's lymphoma in combination with CHOP (cyclophosphamide, doxorubicin, vincristine, prednisolone) chemotherapy.</p> <p>In combination with chemotherapy is indicated for the treatment of patients with previously untreated and relapsed/refractory chronic lymphocytic leukaemia. Only limited data are available on efficacy and safety for patients previously treated with monoclonal antibodies including rituximab or patients refractory to previous rituximab plus chemotherapy.</p> <p>In combination with glucocorticoids, is indicated for the induction of remission in adult patients with severe, active granulomatosis with polyangiitis (Wegener's) (GPA) and microscopic polyangiitis (MPA).</p>	15/06/2017	Sandoz GmbH
RIXIMYO	rituximab	<p>Treatment of previously untreated patients with stage III-IV follicular lymphoma in combination with chemotherapy.</p> <p>As maintenance therapy for the treatment of follicular lymphoma patients responding to induction therapy.</p> <p>As monotherapy for the treatment of patients with stage III-IV follicular lymphoma who are chemoresistant or are in their second or subsequent relapse after chemotherapy.</p> <p>For the treatment of patients with CD20 positive diffuse large B cell non Hodgkin's lymphoma in combination with CHOP (cyclophosphamide, doxorubicin, vincristine, prednisolone) chemotherapy.</p> <p>In combination with glucocorticoids, is indicated for the induction of remission in adult patients with severe, active granulomatosis with polyangiitis (Wegener's) (GPA) and microscopic polyangiitis (MPA).</p>	15/06/2017	Sandoz GmbH
RIXUBIS	nonacog gamma	<p>Treatment and prophylaxis of bleeding in patients with haemophilia B (congenital factor IX deficiency).</p> <p>RIXUBIS is indicated in patients of all age groups.</p>	19/12/2014	Baxalta Innovations GmbH

Tradename	Active Substance	Marketing Authorisation Indication	Marketing Authorisation Date (Dd/Mm/Yyyy)	Marketing Authorisation Holder
ROACTEMRA	tocilizumab	Treatment of active systemic juvenile idiopathic arthritis (sJIA) in patients 2 years of age and older, who have responded inadequately to previous therapy with NSAIDs and systemic corticosteroids. RoActemra can be given as monotherapy (in case of intolerance to MTX or where treatment with MTX is inappropriate) or in combination with MTX. In combination with methotrexate (MTX) it is indicated for the treatment of juvenile idiopathic polyarthritis (pJIA; rheumatoid factor positive or negative and extended oligoarthritis) in patients 2 years of age and older, who have responded inadequately to previous therapy with MTX.	16/01/2009	Roche Registration Ltd
RUCONEST	conestat alfa	Treatment of acute angioedema attacks in adults and adolescents with hereditary angioedema (HAE) due to C1 esterase inhibitor deficiency.	28/10/2010	Pharming Group N.V.
SAVENE	dexrazoxane	Treatment of anthracycline extravasation in adults.	28/07/2006	Clinigen Healthcare Ltd
SIKLOS	hydroxycarbamide	Prevention of recurrent painful vaso-occlusive crises including acute chest syndrome in adults, adolescents and children older than 2 years suffering from symptomatic sickle cell syndrome.	29/06/2007	Addmedica
SIMPONI	golimumab	In combination with methotrexate (MTX) for the treatment of polyarticular juvenile idiopathic arthritis in children with a body weight of at least 40 kg, who have responded inadequately to previous therapy with MTX	01/10/2009	Janssen Biologics B.V.
SOMAVERT	pegvisomant	Treatment of adult patients with acromegaly who have had an inadequate response to surgery and/or radiation therapy and in whom an appropriate medical treatment with somatostatin analogues did not normalize IGF-I concentrations or was not tolerated.	13/11/2002	Pfizer Ltd
SPECTRILA	asparaginase	Indicated as a component of antineoplastic combination therapy for the treatment of acute lymphoblastic leukaemia (ALL) in paediatric patients from birth to 18 years and adults.	14/01/2016	Medac Gesellschaft fuer klinische Spezialpraeparate mbH
SPRYCEL	dasatinib	Treatment of adult patients with: - newly diagnosed Philadelphia chromosome positive (Ph+) chronic myelogenous leukaemia (CML) in the chronic phase. - chronic, accelerated or blast phase CML with resistance or intolerance to prior therapy including imatinib mesilate. - Ph+ acute lymphoblastic leukaemia (ALL) and lymphoid blast CML with resistance or intolerance to prior therapy.	20/11/2006	Bristol-Myers SquibbPharma EEIG

Tradename	Active Substance	Marketing Authorisation Indication	Marketing Authorisation Date (Dd/Mm/Yyyy)	Marketing Authorisation Holder
STAYVEER	bosentan monohydrate	Treatment of pulmonary arterial hypertension (PAH) to improve exercise capacity and symptoms in patients with WHO functional class III. Efficacy has been shown in: <ul style="list-style-type: none"> • Primary (idiopathic and heritable) pulmonary arterial hypertension • Pulmonary arterial hypertension secondary to scleroderma without significant interstitial pulmonary disease • Pulmonary arterial hypertension associated with congenital systemic-to-pulmonary shunts and Eisenmenger's physiology. Some improvements have also been shown in patients with pulmonary arterial hypertension WHO functional class II. Indicated to reduce the number of new digital ulcers in patients with systemic sclerosis and ongoing digital ulcer disease	24/06/2013	Marklas Nederland BV
SUTENT	sunitinib	Treatment of unresectable and/or metastatic malignant gastrointestinal stromal tumour (GIST) in adults after failure of imatinib treatment due to resistance or intolerance. Treatment of advanced/metastatic renal cell carcinoma (MRCC) in adults. Treatment of unresectable or metastatic, well-differentiated pancreatic neuroendocrine tumours (pNET) with disease progression in adults. Experience with SUTENT as first-line treatment is limited	19/07/2006	Pfizer Ltd
TADALAFIL GENERICS	tadalafil	Indicated in adults for the treatment of pulmonary arterial hypertension (PAH) classified as WHO functional class II and III, to improve exercise capacity. Efficacy has been shown in idiopathic PAH (IPAH) and in PAH related to collagen vascular disease.	09/01/2017	MYLAN S.A.S
TARCEVA	erlotinib	In combination with gemcitabine, for the treatment of patients with metastatic pancreatic cancer . When prescribing Tarceva, factors associated with prolonged survival should be taken into account. No survival advantage could be shown for patients with locally advanced disease.	19/09/2005	Roche Registration Ltd
TARGRETIN	bexarotene	Treatment of skin manifestations of advanced stage cutaneous T-cell lymphoma (CTCL) patients refractory to at least one systemic treatment.	29/03/2001	Eisai Ltd
TAXESPIRA (previously DOCETAXEL HOSPIRA UK LIMITED)	docetaxel	In combination with cisplatin and 5-fluorouracil for the induction treatment of patients with locally advanced squamous cell carcinoma of the head and neck .	28/08/2015	Hospira UK Ltd
TAXOTERE	docetaxel	In combination with cisplatin and 5-fluorouracil for the induction treatment of patients with locally advanced squamous cell carcinoma of the head and neck .	27/11/1995	Aventis Pharma S.A.
TEMODAL	temozolomide	Treatment of adult patients with newly-diagnosed glioblastoma multiforme concomitantly with radiotherapy (RT) and subsequently as monotherapy treatment. Treatment of children from the age of three years, adolescents and adult patients with malignant glioma , such as glioblastoma multiforme or anaplastic astrocytoma , showing recurrence or progression after standard therapy.	26/01/1999	Merck Sharp & Dohme Ltd

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TEVAGRASTIM	filgrastim	In patients, children or adults, with severe congenital, cyclic, or idiopathic neutropenia with an absolute neutrophil count (ANC) of $\leq 0.5 \times 10^9/l$, and a history of severe or recurrent infections.	15/09/2008	Teva GmbH
TEYSUNO	tegafur/gimeracil/o teracil	In adults for the treatment of advanced gastric cancer when given in combination with cisplatin.	14/03/2011	Nordic Group BV
THYROGEN	thyrotropin alfa	For use with serum thyroglobulin (Tg) testing with or without radioiodine imaging for the detection of thyroid remnants and well-differentiated thyroid cancer in post-thyroidectomy patients maintained on hormone suppression therapy (THST). Low-risk patients with well-differentiated thyroid carcinoma who have undetectable serum Tg levels on THST and no rh (recombinant human) TSH- stimulated increase of Tg levels may be followed-up by assaying rh TSH-stimulated Tg levels. For pre-therapeutic stimulation in combination with a range of 30 mCi (1.1 GBq) to 100 mCi (3.7 GBq) radioiodine for ablation of thyroid tissue remnants in patients who have undergone a near-total or total thyroidectomy for well-differentiated thyroid cancer and who do not have evidence of distant metastatic thyroid cancer.	09/03/2000	Genzyme Europe B.V.
TORISEL	temsirolimus	First-line treatment of adult patients with advanced renal cell carcinoma (RCC) who have at least three of six prognostic risk factors.	19/11/2007	Pfizer Ltd
TRACLEER	bosentan monohydrate	Treatment of pulmonary arterial hypertension (PAH) to improve exercise capacity and symptoms in patients with WHO functional class III. Efficacy has been shown in : - primary (idiopathic and heritable) PAH , - PAH secondary to scleroderma without significant interstitial pulmonary disease, - PAH associated with congenital systemic-to-pulmonary shunts and Eisenmenger's physiology . Some improvements have also been shown in patients with PAH WHO functional class II. To reduce the number of new digital ulcers in patients with systemic sclerosis and ongoing digital ulcer disease.	15/05/2002	Actelion Registration Ltd
TRISENOX	arsenic trioxide	Indicated for induction of remission, and consolidation in adult patients with: • Newly diagnosed low-to-intermediate risk acute promyelocytic leukaemia (APL) (white blood cell count, $\leq 10 \times 10^3/\mu l$) in combination with all-trans-retinoic acid (ATRA) • Relapsed/refractory acute promyelocytic leukaemia (APL) (Previous treatment should have included a retinoid and chemotherapy) characterised by the presence of the t(15;17) translocation and/or the presence of the Pro-Myelocytic Leukaemia/Retinoic-Acid-Receptor-alpha (PML/RAR-alpha) gene. The response rate of other acute myelogenous leukaemia subtypes to arsenic trioxide has not been examined.	05/03/2002	Teva Pharma B.V.

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TRUXIMA	rituximab	<p>Treatment of previously untreated patients with stage III-IV follicular lymphoma in combination with chemotherapy.</p> <p>Truxima maintenance therapy is indicated for the treatment of follicular lymphoma patients responding to induction therapy.</p> <p>Truxima monotherapy is indicated for treatment of patients with stage III-IV follicular lymphoma who are chemo-resistant or are in their second or subsequent relapse after chemotherapy.</p> <p>Treatment of patients with CD20 positive diffuse large B cell non-Hodgkin's lymphoma in combination with CHOP (cyclophosphamide, doxorubicin, vincristine, prednisolone) chemotherapy.</p> <p>In combination with chemotherapy is indicated for the treatment of patients with previously untreated and relapsed/refractory Chronic lymphocytic leukaemia (CLL) . Only limited data are available on efficacy and safety for patients previously treated with monoclonal antibodies including Truxima or patients refractory to previous Truxima plus chemotherapy.</p> <p>In combination with glucocorticoids, is indicated for the induction of remission in adult patients with severe, active granulomatosis with polyangiitis (Wegener's) (GPA) and microscopic polyangiitis (MPA).</p>	17/02/2017	Celltrion Healthcare Hungary Kft.
UCEDANE	carglumic acid	Treatment of hyperammonaemia due to N-acetylglutamate synthase primary deficiency .	23/06/2017	Lucane Pharma
UPTRAVI	selexipag	<p>Long-term treatment of pulmonary arterial hypertension (PAH) in adult patients with WHO functional class (FC) II–III, either as combination therapy in patients insufficiently controlled with an endothelin receptor antagonist (ERA) and/or a phosphodiesterase type 5 (PDE-5) inhibitor, or as monotherapy in patients who are not candidates for these therapies.</p> <p>Efficacy has been shown in a PAH population including idiopathic and heritable PAH, PAH associated with connective tissue disorders, and PAH associated with corrected simple congenital heart disease.</p>	12/05/2016	Actelion Registration Ltd
VEDROP	tocofersolan	Indicated in vitamin E deficiency due to digestive malabsorption in paediatric patients with congenital chronic cholestasis or hereditary chronic cholestasis , from birth (full term newborns) up to 18 years of age.	24/07/2009	Orphan Europe S.A.R.L

Tradename	Active Substance	Marketing Authorisation Indication	Marketing Authorisation Date (Dd/Mm/Yyyy)	Marketing Authorisation Holder
VELCADE	bortezomib	As monotherapy or in combination with pegylated liposomal doxorubicin or dexamethasone is indicated for the treatment of adult patients with progressive multiple myeloma who have received at least 1 prior therapy and who have already undergone or are unsuitable for haematopoietic stem cell transplantation. In combination with melphalan and prednisone is indicated for the treatment of adult patients with previously untreated multiple myeloma who are not eligible for high-dose chemotherapy with haematopoietic stem cell transplantation. In combination with dexamethasone, or with dexamethasone and thalidomide, is indicated for the induction treatment of adult patients with previously untreated multiple myeloma who are eligible for high-dose chemotherapy with haematopoietic stem cell transplantation. In combination with rituximab, cyclophosphamide, doxorubicin and prednisone is indicated for the treatment of adult patients with previously untreated mantle cell lymphoma who are unsuitable for haematopoietic stem cell transplantation.	26/04/2004	Janssen-Cilag International N.V.
VENTAVIS	iloprost	Treatment of patients with primary pulmonary hypertension , classified as NYHA functional class III, to improve exercise capacity and symptoms.	16/09/2003	Bayer Pharma AG
VFEND	voriconazole	In adults and children aged 2 years and above as follows: - treatment of invasive aspergillosis . - treatment of serious fungal infections caused by <i>Scedosporium spp.</i> and <i>Fusarium spp.</i> Vfend should be administered primarily to patients with progressive, possibly life-threatening infections. Prophylaxis of invasive fungal infections in high risk allogeneic hematopoietic stem cell transplant (HSCT) recipients.	19/03/2002	Pfizer Ltd
VONCENTO	human coagulation factor viii/ von willebrand factor	Treatment of haemorrhage or prevention and treatment of surgical bleeding in patients with von Willebrand disease (VWD) , when desmopressin (DDAVP) treatment alone is ineffective or contraindicated. Prophylaxis and treatment of bleeding in patients with haemophilia A (congenital FVIII deficiency) .	12/08/2013	CSL Behring GmbH
VORICONAZOLE HOSPIRA	voriconazole	In adults and children aged 2 years and above as follows: - treatment of invasive aspergillosis . - treatment of serious fungal infections caused by <i>Scedosporium spp.</i> and <i>Fusarium spp.</i> Voriconazole should be administered primarily to patients with progressive, possibly life – threatening infections.	27/05/2015	Hospira UK Ltd

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VOTRIENT	pazopanib	In adults for the first-line treatment of advanced renal cell carcinoma (RCC) and for patients who have received prior cytokine therapy for advanced disease. For the treatment of adult patients with selective subtypes of advanced soft-tissue sarcoma (STS) who have received prior chemotherapy for metastatic disease or who have progressed within 12 months after (neo)-adjuvant therapy. Efficacy and safety have only been established in certain STS histological tumour subtypes.	14/06/2010	Novartis Europharm Ltd
WILZIN	zinc acetate dihydrate	Treatment of Wilson's disease .	13/10/2004	Orphan Europe S.A.R.L
XAGRID	anagrelide hydrochloride	Reduction of elevated platelet counts in at-risk essential-thrombocythaemia (ET) patients who are intolerant to their current therapy or whose elevated platelet counts are not reduced to an acceptable level by their current therapy. An at risk ET is defined by one or more of the following features: - > 60 years of age or - a platelet count > 1000 x 10 ⁹ /l or - a history of thrombo-haemorrhagic events.	16/11/2004	Shire Pharmaceutical Contracts Ltd
XELODA	capecitabine	First-line treatment of advanced gastric cancer in combination with a platinum-based regimen	02/02/2001	Roche Registration Ltd
XYREM	sodium oxybate	Treatment of narcolepsy with cataplexy in adult patients.	13/10/2005	UCB Pharma Ltd
YARGESA	miglustat	For the oral treatment of adult patients with mild to moderate type 1 Gaucher disease . Yargesa may be used only in the treatment of patients for whom enzyme replacement therapy is unsuitable	22/03/2017	JensonR+ Limited
YONDELIS	trabectedin	Treatment of adult patients with advanced soft tissue sarcoma , after failure of anthracyclines and ifosfamide, or who are unsuited to receive these agents. Efficacy data are based mainly on liposarcoma and leiomyosarcoma patients.	17/09/2007	Pharma MarS.A.
ZARZIO	filgrastim	In children and adults with severe congenital, cyclic, or idiopathic neutropenia with an absolute neutrophil count (ANC) of $\leq 0.5 \times 10^9/l$, and a history of severe or recurrent infections, long term administration of filgrastim is indicated to increase neutrophil counts and to reduce the incidence and duration of infection-related events.	06/02/2009	Sandoz GmbH
ZAVESCA	miglustat	Treatment of adult patients with mild to moderate type 1 Gaucher disease . Zavesca may be used only in the treatment of patients for whom enzyme replacement therapy is unsuitable.	21/11/2002	Actelion Registration Ltd
ZEVALIN	ibritumomab tiuxetan	Consolidation therapy after remission induction in previously untreated patients with follicular lymphoma . Treatment of adult patients with rituximab relapsed or refractory CD20+ follicular B-cell non-Hodgkin's lymphoma (NHL) .	16/01/2004	Spectrum Pharmaceuticals B.V.

Tradename	Active Substance	Marketing Authorisation Indication	Marketing Authorisation Date (Dd/Mm/Yyyy)	Marketing Authorisation Holder
ZUTECTRA	human hepatitis b immunoglobulin	Prevention of hepatitis B virus (HBV) re-infection in HBV-DNA negative patients over 6 months after liver transplantation for hepatitis B induced liver failure . Zutectra is indicated in adults only. The concomitant use of adequate virostatic agents should be considered, if appropriate, as standard of hepatitis B re-infection prophylaxis.	30/11/2009	Biotest Pharma GmbH
ZYDELIG	idelalisib	In combination with rituximab, treatment of adult patients with chronic lymphocytic leukaemia (CLL) : - who have received at least one prior therapy, or - as first line treatment in the presence of 17p deletion or TP53 mutation in patients unsuitable for chemo-immunotherapy. As monotherapy, treatment of adult patients with follicular lymphoma (FL) that is refractory to two prior lines of treatment.	18/09/2014	Gilead Sciences International Ltd.

Classificazione per data dell'AIC in ordine decrescente

2018
ADYNOVI
2017
AFSTYLA
AMGEVITA
BLITZIMA
CUPRIOR
CYLTEZO
ELMIRON
ERELZI
FOTIVDA
IMATINIB TEVA B.V.
IMRALDI
MIGLUSTAT GEN ORPH
NITISINONE MDK
REFIXIA
RITEMVIA
RITUZENA
RIXATHON
RIXIMYO
TADALAFIL GENERICS
TRUXIMA
UCEDANE
YARGESA
2016
ARMISARTE
BORTEZOMIB HOSPIRA
BORTEZOMIB SUN
CABOMETYX
EMPLICITI
IBLIAS
KISPLYX
KOVALTRY
MYSILDECARD
NEOFORDEX
NORDIMET
ONCASPAS
PEMETREXED ACCORD
PEMETREXED FRESENIUS KABI
SPECTRILA
UPTRAVI
2015
BORTEZOMIB ACCORD
TAXESPIRA
ELOCTA
KEYTRUDA
OBIZUR
OPDIVO
ORKAMBI
PEMETREXED HOSPIRA
PEMETREXED LILLY
PEMETREXED MEDAC
PEMETREXED SANDOZ

QUINSAIR
REPATHA
RESPREEZA
VORICONAZOLE HOSPIRA
2014
BEMFOLA
RIXUBIS
ZYDELIG
2013
ERIVEDGE
GRASTOFIL
HYQVIA
LOJUXTA
NOVOEIGHT
OVALEAP
STAYVEER
VONCENTO
2012
CAPRELSA
COLOBREATHA
INLYTA
JAKAVI
NOVOTHIRTEEN
PIXUVRI
2011
BUCCOLAM
CINRYZE
EURARTESIM
HALAVEN
HIZENTRA
TEYSUNO
2010
NIVESTIM
OZURDEX
REVOLADE
RUCONEST
VOTRIENT
2009
AFINITOR
FILGRASTIM HEXAL
ILARIS
IXIARO
ROACTEMRA
SIMPONI
VEDROP
ZARZIO
ZUTECTRA
2008
ABRAXANE
ADCIRCA
PRIVIGEN
RATIOGRASTIM
TEVAGRASTIM

2007
ATRIANCE
CYSTADANE
DIACOMIT
ELAPRASE
FLEBOGAMMA DIF
GLIOLAN
INCRELEX
ORENCIA
REVLIMID
SIKLOS
TORISEL
YONDELIS
2006
ATRYN
KIOVIG
EVOLTRA
EXJADE
NEXAVAR
OMNITROPE
MYOZYME
NAGLAZYME
SAVENE
SPRYCEL
SUTENT
2005
AVASTIN
NOXAFIL
ORFADIN
REVATIO
TARCEVA
XYREM
2004
ADVATE
ALIMTA
DUKORAL
ERBITUX
LITAK
LYSODREN
PEDEA
VELCADE
WILZIN
XAGRID
ZEVALIN
2003
ALDURAZYME
BUSILVEX
CARBAGLU
HUMIRA
VENTAVIS
2002
KINERET
SOMAVERT

TRACLEER
TRISENOX
VFEND
ZAVESCA
2001
CANCIDAS
CEPROTIN
DEPOCYTE
FABRAZYME
GLIVEC
INOMAX
NONAFAC
REPLAGAL
TARGRETIN
XELODA
2000
ENBREL
HELIXATE NEXGEN
HERCEPTIN
INTRONA
KEPPRA
KOGENATE BAYER
PANRETIN
THYROGEN
1999
AMMONAPS
FERRIPROX
REFACTO AF
TEMODAL
1998
MABTHERA
1997
BENEFIX
CEREZYME
CYSTAGON
1996
CAELYX
HYCANTIN
NOVOSEVEN
PUREGON
RILUTEK
1995
GONAL-F
TAXOTERE

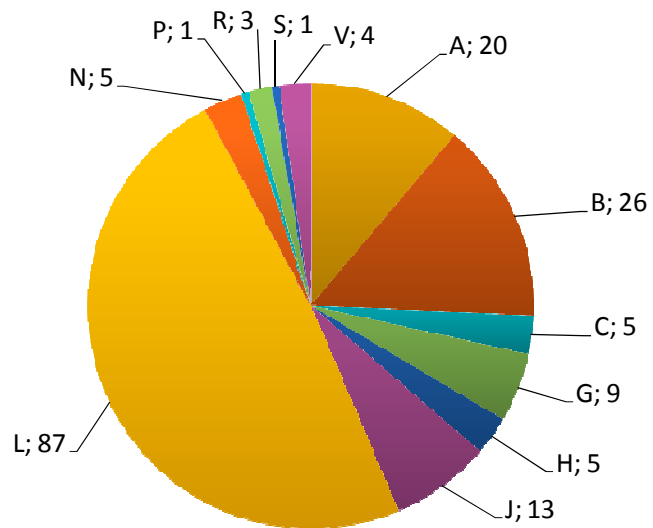
Numero dei farmaci orfani in Europa con autorizzazione all'immissione in commercio europea senza designazione orfana europea per data dell'AIC



Classificazione per categoria ATC

A- ALIMENTARY TRACT AND METABOLISM	LOJUXTA	BLITZIMA	PEMETREXED SANDOZ
ALDURAZYME	PEDEA	BORTEZOMIB ACCORD	PIXUVRI
AMMONAPS	REPATHA	BORTEZOMIB HOSPIRA	RATIOGRASTIM
CARBAGLU	STAYVEER	BORTEZOMIB SUN	REVLIMID
CEREZYME	TRACLEER	BUSILVEX	RITEMVIA
CUPRIOR	G- GENITO URINARY SYSTEM AND SEX HORMONES	CABOMETYX	RITUZENA
CYSTADANE	ADCIRCA	CAELYX	RIXATHON
CYTAGON	BEMFOLA	CAPRELSA	RIXIMYO
ELAPRASE	ELMIRON	CYLTEZO	ROACTEMRA
FABRAZYME	GONAL-F	DEPOCYTE	SIKLOS
MIGLUSTAT GEN ORPH	MYSILDECARD	TAXESPIRA	SIMPONI
MYOZYME	OVALEAP	EMPLICITI	SPECTRILA
NAGLAZYME	PUREGON	ENBREL	SPRYCEL
NITISINONE MDK	REVIATIO	ERBITUX	SUTENT
ORFADIN	TADALAFIL GENERICS	ERELZI	TARCEVA
REPLAGAL	H- SYSTEMIC HORMONAL PREPARATIONS, EXCL. SEX HORMONES AND INSULINS	ERIVEDGE	TARGRETIN
UCEDANE	INCRELEX	EVOLTRA	TAXOTERE
VEDROP	NEOFORDEX	FILGRASTIM HEXAL	TEMODAL
WILZIN	OMNITROPE	FOTIVDA	TEVAGRASTIM
YARGESA	SOMAVERT	GLIOLAN	TEYSUNO
ZAVESCA	THYROGEN	GLIVEC	TORISEL
B- BLOOD AND BLOOD FORMING ORGANS	J- GENERAL ANTIINFECTIVES FOR SYSTEMIC USE	GRASTOFIL	TRISENOX
ADVATE	CANCIDAS	HALAVEN	TRUXIMA
ADYNOVI	DUKORAL	HERCEPTIN	VELCADE
AFSTYLA	FLEBOGAMMA DIF	HUMIRA	VOTRIENT
ATRYN	HIZENTRA	HYCAMTIN	XAGRID
BENEFIX	HYQVIA	ILARIS	XELODA
CEPROTIN	IXIARO	IMATINIB TEVA B.V.	YONDELIS
CINRYZE	KIOVIG	IMRALDI	ZARZIO
ELOCTA	NOXAFIL	INLYTA	ZYDELIG
HELIXATE NEXGEN	PRIVIGEN	INTRONA	N- NERVOUS SYSTEM
IBLIAS	QUINSAIR	JAKAVI	BUCCOLAM
KOGENATE BAYER	VFEND	KEYTRUDA	DIACOMIT
KOVALTRY	VORICONAZOLE HOSPIRA	KINERET	KEPPRA
NONAFAC	ZUTECTRA	KISPLYX	RILUTEK
NOVOEIGHT	L- ANTINEOPLASTIC AND IMMUNOMODULATING AGENTS	LITAK	XYREM
NOVOSEVEN	ABRAXANE	LYSODREN	P- ANTIPARASITIC PRODUCTS, INSECTICIDES AND REPELLENTS
NOVOTHIRTEEN	AFINITOR	MABTHERA	EURARTESIM
OBIZUR	ALIMTA	NEXAVAR	R- RESPIRATORY SYSTEM
REFACTO AF	AMGEVITA	NIVESTIM	COLOBREATHE
REFIXIA	ARMISARTE	NORDIMET	INOMAX
RESPREEZA	ATRIANCE	ONCASPAR	ORKAMBI
REVOLADE	AVASTIN	OPDIVO	S- SENSORY ORGANS
RIXUBIS		ORENCIA	OZURDEX
RUCONEST		PANRETIN	V- VARIOUS
UPTRAVI		PEMETREXED ACCORD	EXJADE
VENTAVIS		PEMETREXED FRESENIUS KABI	FERRIPROX
VONCENTO		PEMETREXED HOSPIRA	SAVENE
C- CARDIOVASCULAR SYSTEM		PEMETREXED LILLY	ZEVALIN
		PEMETREXED MEDAC	

Numero dei farmaci orfani in Europa con autorizzazione all'immissione in commercio europea senza designazione orfana europea precedente per categoria ATC



Classificazione per titolare dell'AIC

ABBVIE LTD	CYLTEZO	GTC BIOTHERAPEUTICS UK LIMITED	AFINITOR
HUMIRA	BRISTOL-MYERS SQUIBB PHARMA EEIG	ATRYN	ATRIANCE
ADDMEDICA	EMPLICITI	HEXAL AG	EXJADE
SIKLOS	OPDIVO	FILGRASTIM HEXAL	GLIVEC
ACCORD HEALTHCARE LTD	ORENCIA	HOSPIRA UK LTD	HYCANTIN
BORTEZOMIB ACCORD	SPRYCEL	BORTEZOMIB HOSPIRA	ILARIS
PEMETREXED ACCORD	CELGENE EUROPE LTD	TAXESPIRA	JAKAVI
ACTAVIS GROUP PTC EHF	ABRAXANE	PEMETREXED HOSPIRA	REVOLADE
ARMISARTE	REVLIMID	NIVESTIM	VOTRIENT
ACTELION REGISTRATION LTD	CELLTRION HEALTHCARE HUNGARY KFT.	VORICONAZOLE HOSPIRA	NOVO NORDISK A/S
TRACLEER	BLITZIMA	INSTITUTO GRIFOLS S.A.	NOVOEIGHT
UPTRAVI	RITEMVIA	FLEBOGAMMA DIF	NOVOSEVEN
ZAVESCA	RITUZENA	IPSEN PHARMA	NOVOTHIRTEEN
AEGERION PHARMACEUTICALS SAS	TRUXIMA	CABOMETYX	REFIXIA
LOJUXTA	CLINIGEN HEALTHCARE LTD	INCRELEX	ORPHAN EUROPE S.A.R.L.
ALLERGAN PHARMACEUTICALS IRELAND	SAVENE	JANSSEN-CILAG INTERNATIONAL NV	CARBAGLU
OZURDEX	CRUCELL SWEDEN AB	CAELYX	CYSTADANE
AMGEN EUROPE BV	DUKORAL	SIMPONI	CYSTAGON
AMGEVITA	CSL BEHRING GMBH	VELCADE	PEDEA
REPATHA	AFSTYLA	JENSON PHARMACEUTICALS SERVICES LIMITED	VEDROP
APOTEX EUROPE B.V.	HIZENTRA	YARGESA	WILZIN
FERRIPROX	PRIVIGEN	LABORATOIRES CTRS	PACIRA LIMITED
GRASTOFIL	RESPREEZA	NEOFORDEX	DEPOCYTE
APTALIS PHARMA SAS	VONCENTO	LABORATOIRE HRA PHARMA	PFIZER LTD
QUINSAIR	CTI LIFE SCIENCES LTD	LYSODREN	BENEFIX
ASTRAZENECA AB	PIXUVRI	LINDE HEALTHCARE AB	ENBREL
CAPRELSA	EISAI EUROPE LTD	INOMAX	INLYTA
AVENTIS PHARMA S.A.	HALAVEN	LIPOMED GMBH	REFACTO AF
RILUTEK	KISPLYX	LITAK	REVATIO
TAXOTERE	PANRETIN	LUCANE PHARMA	SOMAVERT
BAXALTA INNOVATIONS GMBH	TARGRETIN	UCEDANE	SUTENT
ADYNOVI	ELI LILLY NEDERLAND B.V.	MARKLAS NEDERLAND BV	TORISEL
HYQVIA	ADCIRCA	STAYVEER	VFEND
OBIZUR	ALIMTA	MEDAC GMBH	PHARMA MARS.A.
ONCASPAR	PEMETREXED LILLY	GLIOLAN	YONDELIS
RIXUBIS	EUSA Pharma (UK) Limited	PEMETREXED MEDAC	PHARMING GROUP N.V.
BAXTER AG	FOTIVDA	SPECTRILA	RUCONEST
ADVATE	FINOX BIOTECH AG	MENDELKABS EUROPE LTD	PIERRE FABRE MÉDICAMENTS
CEPROTIN	BEMFOLA	NITISINONE MDK	BUSILVEX
KIOVIG	FOREST LABORATORIES UK LTD	MERCK KGAA	RATIOPHARM GMBH
BAYER PHARMA AG	COLOBREATHE	ERBITUX	RATIOGRASTIM
HELIXATE NEXGEN	FRESENIUS KABI ONCOLOGY PLC	MERCK SERONO EUROPE LTD	ROCHE REGISTRATION LTD
IBLIAS	PEMETREXED FRESENIUS	GONAL-F	AVASTIN
KOGENATE BAYER	GEN.ORPH	MERCK SHARP & DOHME LTD	ERIVEDGE
KOVALTRY	MIGLUSTAT GEN ORPH	CANCIDAS	HERCEPTIN
NEXAVAR	GENZYME EUROPE B.V.	INTRONA	MABTHERA
VENTAVIS	ALDURAZYME	KEYTRUDA	ROACTEMRA
BENE- ARZNEIMITTEL GMBH	CEREZYME	NOXAFIL	TARCEVA
ELMIRON	EVOLTRA	PUREGON	XELODA
BIOCODEX	FABRAZYME	TEMODAL	SAMSUNG BIOEPIS UK LIMITED (SBUK)
DIACOMIT	MYOZYME	MYLAN SAS	IMRALDI
 BIOGEN IDEC LTD	THYROGEN	MYSILDECARD	SANDOZ GMBH
ELOCTA	GILEAD SCIENCES INTERNATIONAL LTD	TADALAFIL GENERICS	ERELZI
BIOMARIN EUROPE LTD	ZYDELIG	NORDIC GROUP BV	OMNITROPE
NAGLAZYME	GMP-ORPHAN SA	NORDIMET	PEMETREXED SANDOZ
BIOTEST PHARMA GMBH	CUPRIOR	TEYSUNO	RIXATHON
ZUTECTRA		NOVARTIS EUROPHARM LTD	RIXIMYO
Boehringer Ingelheim International GmbH			ZARZIO
			SANQUIN
			NONAFACT
			SHIRE PHARMACEUTICAL CONTRACTS LTD
			XAGRID

SHIRE HUMAN GENETIC THERAPIES AB	ZEVALIN	TEVA GMBH	VALNEVA AUSTRIA GMBH
ELAPRASE	SUN Pharmaceutical Industries (Europe) B.V.	TEVAGRASTIM	IXIARO
REPLAGAL	BORTEZOMIB SUN	TEVA PHARMA BV	VERTEX PHARMACEUTICALS (EUROPE) LTD
SIGMA-TAU INDUSTRIE FARMACEUTICHE RIUNITE S.P.A	SWEDISH ORPHAN BIOVITRUM INTERNATIONAL AB	IMATINIB TEVA B.V.	ORKAMBI
EURARTESIM	AMMONAPS	OVALEAP	VIROPHARMA SPRL
SPECTRUM PHARMACEUTICALS B.V.	KINERET	TRISENOX	BUCCOLAM
	ORFADIN	UCB PHARMA LTD	CINRYZE
		XYREM	
		UCB PHARMA SA	
		KEPPRA	

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