

PPRI GLOSSARY

ABBREV	Preferred term	Synonym term	Definition/Description
	Access		The ability to obtain health care, as determined by factors such as the availability and affordability of goods and services.
	Active Ingredient	(Active) Substance, Compound	The primary chemical substance or compound contained in a pharmaceutical. Some pharmaceuticals contain more than one active ingredient (combination product).
	Acute Care Beds		Hospital beds available for acute care, defined as curative care
	Affordability		Financial accessibility of goods and services.
	Analogous Substitution		Dispensation of a pharmaceutical (often generic) by the pharmacist with a different active ingredient (or combination product) but the same therapeutical effect instead of the product prescribed by the doctor. Cf. also generic substitution .
ATC	Anatomic Therapeutic Chemical classification	Anatomic Therapeutic Chemical Code	In this WHO classification system pharmaceuticals are divided into different groups according to the organ or system on which they act and their chemical, pharmacological and therapeutic properties.
	ATC 4 level	ATC-4 level	Defines a therapeutic group within the anatomic therapeutic chemical classification system.
	ATC 5 level	ATC-5 level	Defines a single active ingredient or a fixed combination of active ingredients within the anatomic therapeutic chemical classification system .
	Benchmarking		A process of measuring another organisation's product or service according to specified standards in order to compare it and improve one's own product or service.
	Bioavailability	Bio-availability	The amount of an active ingredient that is absorbed by the organism and the speed at which this occurs when introduced in a given dosage form.
	Bioequivalence	Bio-equivalence	Two pharmaceuticals are bioequivalent if they are pharmaceutically equivalent (cf. pharmaceutical equivalence) and their bioavailability (rate and extent of availability) - after administration in the same moral dose - is similar to such a degree that their effect can be expected to be essentially the same.
	Brand Name		The trade or marketing name of a pharmaceutical . Also generics may have a brand name.
	Branded Generic		Cf. generic .
	Capitation		A remuneration method for general practitioners applied by some social health insurance / national health service . The remuneration is based on a fixed monthly amount for each enrolled/listed patient.
CIP	Carriage and Insurance Packaging	Carriage and Insurance Paid To	A type of price quotation, indicating the delivery of goods including cargo insurance to the named place of destination at seller's expense. In an export the quotation indicates the place of destination (discharge) after the acronym CIP, for example CIP Athens.
	Central Taxation		Taxes collected through central government.
	Centralised Procedure		Way of approval of pharmaceuticals valid in all Member States. The Centralised Procedure is administered by the European Medicines Agency (EMA) in London. It consists of a single application which, when approved, grants marketing authorisation for all markets within the European Union. This procedure is available to all new, innovative pharmaceuticals, and is obligatory for 1. biotechnology-derived products, 2. new active ingredients for treating AIDS, cancer, diabetes and "neuro-degenerative illnesses" as well as 3. orphan drugs . Under certain conditions the centralised authorisation can be limited for one year. If the pharmaceutical is important for public health (especially therapeutic innovations) the appraisal period can be abbreviated.
	Claw-back	Clawback	A system allowing third party payers to recoup (part of the) discounts/rebates granted in a reimbursement system between various stakeholders, e.g. wholesalers and pharmacists.
	Combination Product		A pharmaceutical that contains more than one active ingredient .
	Community pharmacy		Health care facility dispensing pharmaceuticals (POM and OTC, reimbursable and non-reimbursable pharmaceuticals) to out-patients. Pharmacies are subject to pharmacy legislation (e.g. national legislation regarding establishment and ownership of pharmacies), cf. pharmaceutical retailer
	Compulsory Health Insurance	Obligatory Health Insurance	Health Insurance under an obligatory scheme basing on a legal act, usually with income-related contributions. See also social health insurance and voluntary health insurance .
	Consumption-based Reimbursement		The level of reimbursement depends on the expenses for pharmaceuticals of a patient within a certain period of time (increasing reimbursement with rising consumption).

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	Contribution		Money paid by or on behalf of insured persons to a social health insurance to purchase the coverage of a defined range of services (the benefit package).
CIF	Cost, Insurance and Freight		The cargo insurance and delivery of goods to the named port of destination (discharge) at the seller's expense. Buyer is responsible for the import customs clearance and other costs and risks. In the export quotation, indicate the port of destination (discharge) after the acronym CIF, for example CIF Athens.
	Cost-Containment	Cost Containment	Measures like price freezes taken to reduce expenditure or the rate of growth of expenditure, or the unit cost of services.
	Cost-Effectiveness Analysis	Cost Effectiveness Analysis	Compares the cost per unit of outcome of alternative therapies with the aim of identifying the most efficient therapy. Determines the cost incurred to obtain an increase in health benefit.
	Cost-plus Pricing	Cost Plus Pricing	Pricing procedure which takes besides the production cost of a pharmaceutical other cost like promotional expenses and especially a profit margin for fixation of the price into account. This share is usually expressed as a percentage of the cost.
	Cross-border trade		The act of importing pharmaceuticals into one country from another for the purpose of personal consumption in the import country.
	Decentralised Procedure		The Decentralised Procedure came into operation in late 2005. It is applicable in cases where a marketing authorisation does not yet exist in any of the EU Member States. Identical dossiers be submitted to all Member States where a marketing authorisation is sought. A Reference Member State, selected by the applicant, will prepare draft assessment documents within 120 days and send them to the Concerned Member States. They, in turn, will either approve the assessment or the application will continue into arbitration procedures. The new Decentralised Procedure will involve concerned Member States at an earlier stage of the evaluation than under the MRP in an effort to minimise disagreements and to facilitate the application for marketing authorisation in as many markets as possible.
	Deductible		Out-of-pocket payment in the form of a fixed amount which must be paid for a service or of total cost incurred over a defined period by a covered person beforehand a social health insurance / national health service , then all or a percentage of the rest of the cost is covered.
DDD	Defined Daily Dose		Technical unit developed in the early 1970ies used to measure the consumption of pharmaceuticals in a comparable way. The DDD is the assumed average maintenance dose per day for a pharmaceutical used for its main indication in adults.
	De-listing	Delisting	Exclusion of a pharmaceutical from a pharmaceutical list (e.g. positive list), often resulting in exclusion from reimbursement .
DRG	Diagnosis Related Group		A way of categorising patients according to diagnosis and intensity of resources required, usually for the period of one hospital stay.
	Direct Payments		Payments for goods and services which are not covered by a social health insurance / national health service or a voluntary health insurance (including self-medication).
	Discount		A price reduction granted to specified purchasers of a pharmaceutical .
	Disease-specific		Eligibility for reimbursement is linked to the underlying disease which shall be treated.
	Dispensing fee		Payment of the pharmacist for the service of dispensing a pharmaceutical .
	Distance Selling		Dispensing of pharmaceuticals via internet or posting services.
	Distributors		Distributors rather sell products under a licence obtained from original manufacturers but most likely do not produce it by themselves.
	Effectiveness		The extent to which a specific intervention, when used under ordinary circumstances, does what it is intended to do. Clinical trials that assess effectiveness are sometimes called management trials. Cf. also pharmacoeconomic evaluation and cost-effectiveness analysis .
	Efficacy		The extent to which an intervention produces a beneficial result under ideal conditions. Clinical trials that assess efficacy are sometimes called explanatory trials and are restricted to participants who fully co-operate. Cf. also pharmacoeconomic evaluation and cost-effectiveness analysis .
	Efficiency		Efficiency measures whether health care resources are being used to maximise value for money.

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	Eligibility Scheme(s)		There are, in general, 4 types of eligibility schemes: * patient-specific reimbursement * disease-specific reimbursement , * population-group-specific reimbursement , * consumption-based reimbursement
	Essential Drug Policy		Essential pharmaceuticals satisfy the primary care needs of a population and are selected with due regard to disease prevalence, evidence on efficacy and safety, and comparative cost-effectiveness . An essential drug policy shall guarantee that such pharmaceuticals are available within the context of the respective health care systems at all times in adequate amounts, in the appropriate dosage forms, with assured quality and at a price the individual and the community can afford.
	Ex-factory Price	Manufacturer Price, Procurement Price, List Price, Wholesale Purchasing Price; Ex factory Price	The manufacturer's posted price, in some countries also referred to as list price. This price does not include any discounts or other incentives offered by manufacturers.
	External Price Referencing	Cross Country Referencing, International Price Comparison, International Price benchmarking	The practice of comparing pharmaceutical prices across countries. There are various methods applied and different country baskets relevant.
	Fee-for Service	Fee for Service	Payments to a provider (for example a general practitioner) for each act or service rendered.
	Fixed Co-payment	Co-payment, copayment	A out-of-pocket payment in the form of a fixed amount (like for example a prescription fee) to be paid for a service, a pharmaceutical or a medical device. See also deductible and percentage co-payment .
	Framework Agreement		This is an agreement between representatives of stakeholders, which serves as a base for individual agreements (this is for example the case in France). It is not limited to price-volumes control.
	Free Pricing		Pricing system, where pharmaceutical prices may be freely set.
G10	G 10 Medicines Group	G-10 Medicines Group	The G 10 Medicines Group (consisting of then Commissioner Liikanen and then Commissioner Byrne, a number of European health and industry ministers, as well as leaders of the pharmaceutical and health insurance industry and patient representatives) was set up in 2001 with the aim to discuss the major issues relevant to the right balance of health objectives and industry competitiveness in Europe. In May 2002 a report was published containing recommendations on how to improve the competitiveness of industry while meeting public and social health objectives. The results were followed-up in the so-called "Post G 10"-process and are discussed in the EU Pharmaceutical Forum (2006-2007). Still a lot of the recommendations are waiting for implementation.
	Gatekeeper	Gate-keeper	A health care person, most likely a general practitioner (but also paediatrician or gynaecologist), who is responsible for overseeing and coordinating all the medical needs of a patient. The gatekeeper has to authorise any referral of the patient to secondary care (specialist) or tertiary care (hospital). Referral exemptions are possible in emergencies or in some systems if patients accept higher out-of pocket payments .
GP	General Practitioner	Family Doctor, Primary Care Physician	A general doctor, who is the first point of contact with the health services for all non-emergency cases. See also gatekeeper .
	Generic		Bioequivalent of a branded original pharmaceutical , whose patent on the active ingredient has expired (also called off-patent or multi-source pharmaceutical). By law, a generic product must contain an identical amount of the same active ingredient(s) as the branded product. There are branded generics and unbranded generics on the market. Branded generics also have a specific trade name, whereas unbranded generics use the international non-proprietary name and the manufacturer's name.
	Generic Substitution	Original Substitution	Practice of substituting a pharmaceutical , whether marketed under a trade name or generic name (branded or unbranded generic), by a pharmaceutical, often a cheaper one, containing the same active ingredient(s). Generic substitution may be performed by prescribers (doctors) and in some countries also by dispensers (pharmacists).

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GDP	Gross Domestic Product		GDP is the value of goods and services provided in a country by residents and non-residents without regard to their allocation among domestic and foreign claims. This corresponds to the total sum of expenditure (consumption and investment) of the private and government agents of the economy during the reference year.
HTA	Health Technology Assessment		HTA is a multidisciplinary process that summarises information about the medical, social, economic and ethical issues related to the use of a health technology in a systematic, transparent, unbiased, robust manner. Its aim is to inform the formulation of safe, effective, health policies that are patient focused and seek to achieve best value.
	Hospital Price		The price of a pharmaceutical in hospital use.
HOM	Hospital-Only Medicines	Hospital Only Medicine	Type of classification; pharmaceuticals that may be only administered in hospitals (inpatient care and outpatient care).
	Human Pharmaceutical		1. Any active ingredient or combination of active ingredients presented as having properties for treating or preventing disease in human beings. 2. Any active ingredient or combination of active ingredients which may be used in human beings with a view to making a medical diagnosis or to restoring, correcting or modifying physiological functions.
	Inpatient Care	In-patient Care	Medical procedures that require a stationary hospital stay (perhaps of only one day).
	Internal Price Referencing	Therapeutic Referencing	A method to compare prices of pharmaceuticals in a country with the price of identical pharmaceuticals (ATC 5 level) or similar products (ATC 4 level) or even with therapeutical equivalent treatment (not necessarily a pharmaceutical) in a country. Often performed in the course of a reference price system .
INN	International Non-proprietary Name	Generic Name, International Nonproprietary Name	A pharmaceutical is normally identified by either its chemical or "generic" name, which often is referred to as INN or its brand name, which is the trade or marketing name.
	Magistral Formula		Magistral Formula is any pharmaceutical prepared in a pharmacy in accordance with a prescription for an individual patient.
	Manufacturer	Pharmaceutical Industry	Pharmaceutical companies who produce pharmaceuticals and very often also search for and develop new chemical entities, i.e. active ingredients .
MA	Marketing Authorisation	Licensing	A licence issued by a medicines agency approving a pharmaceutical for market use based on a determination by authorities that the pharmaceutical meets the requirements of quality, safety and efficacy for human use in therapeutic treatment. There are four application procedures possible in the EU: " centralised procedure " or " mutual recognition procedure " (MRP), " decentralised procedure " and " national procedure ". For homeopathic pharmaceuticals and medical devices no authorisation but a registration procedure is necessary.
	Maximum Price		This term is used in a different way in different countries: e.g. in some countries it is the maximum amount which is reimbursed (cf. reference price system), in others it is the maximum share that is refunded by social health insurance / national health services expressed as percentage of the reimbursement basis.
	Medical Device		Any instrument, apparatus, appliance, material or other article, whether used alone or in combination, including the software necessary for its proper application intended by the manufacturer to be used for human beings for the purpose of: - diagnosis, prevention, monitoring, treatment or alleviation of disease, - diagnosis, monitoring, treatment, alleviation of or compensation for an injury or handicap, - investigation, replacement or modification of the anatomy or of a physiological process, - control of conception, and which does not achieve its principal intended action in or on the human body by pharmacological, immunological or metabolic means, but which may be assisted in its function by such means. This term defines a wide spectrum of products, ranging from spectacles and clutches to highly sophisticated implantable devices.
	Me-too Pharmaceutical	Me too Pharmaceutical	A me-too pharmaceutical is one that is approved after a pioneering product and which is defined as comparable or similar and is not clinically superior.

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MRP	Mutual Recognition Procedure		The MRP is the most common marketing authorisation procedure in the EU. It states that the marketing authorisation granted in one EU Member State (the so-called "Reference Member State") being "mutually recognised" as valid in other Member State (the "Concerned Member State") upon request. The legal basis is Directive 2001/83/EC, as amended by Directive 2004/27/EC, and further guidance is given in the Notice to Applicants, which forms Chapter 2 of the Rules Governing Medicinal Products in the EU.
NHS	National Health Service	Beveridge System, Semashko System	The system of social security and health services arising out of the Beveridge report in England and Wales, first published in 1943. A NHS System is financed through central taxation or regional taxation , usually covering all inhabitants/residents. The scope of services rendered is identical for every person covered and most services are offered by public institutions. In some countries people may opt for a complementary VHI (see voluntary health insurance) for services, which are not covered through the NHS.
	National Procedure		Independent national marketing authorisation procedures are still applicable during the initial stage of the mutual recognition procedure in the country that is to act as reference Member State (RMS). They are also applicable in situations in which the mutual recognition procedure is not compulsory, namely: - Bibliographical applications for pharmaceuticals with a well-established medicinal use for which no reference product is available in the EU. - Line extensions of nationally registered pharmaceuticals for which no harmonised product information is available within the EU. Although some changes to dossiers for nationally registered pharmaceuticals (such as a change in the strength, pharmaceutical form or route of administration) require the submission of a new marketing authorisation application, these changes are considered as variations to a nationally issued marketing authorisation .
	Negative List		List of pharmaceuticals which cannot be prescribed at the expense of the social health insurance/ national health service .
	Officinal Formula		Officinal Formula is any pharmaceutical which is prepared in a pharmacy in accordance with the prescriptions of a pharmacopoeia and is intended to be supplied directly to the patients served by the pharmacy in question (To distinguish from magistral formula).
	Off-patent Pharmaceutical	Off-patent Product, Multi-source Pharmaceutical	(Branded) original product , whose patent has expired as opposed to on-patent pharmaceuticals . Multisource products may be marketed either under the approved international non-proprietary name or under a brand (proprietary) name.
	On-patent pharmaceutical	On-patent Product	A pharmaceutical that is protected by a patent or a supplementary product certificate (SPC); i.e. a branded original product .
	Original Product	Originator, Original Preparation	The first version of a pharmaceutical , developed and patented by an originator pharmaceutical company which has exclusive rights to marketing the product in the European Union for 15 years. A original product has a unique trade name for marketing purposes, its so-called brand name.
	Orphan Drug		A pharmaceutical which only has a limited target population or which treats a rare disease thus limiting its commercial and financial potential.
	Out-of Pocket Maximum	Annual Ceiling	The maximum amount (e.g. a certain percentage of income) that an insured person has to pay for all covered healthcare services for a defined period (often a year), cf. Out-of pocket payments .
OPP	Out-of Pocket Payments	Out-Of-Pocket Payments	The amount a person has to pay for all covered healthcare services for a defined period (often a year). It includes: * fixed co-payments , * percentage co-payments and * deductibles . Cf. out-of pocket maximum and prescription fee .
	Outpatient Care	Out-Of-Hospital Care, Out-patient Care	Medical procedures that do not require a hospital stay. They typically occur in the ambulatory of a hospital (~ outpatient clinic) or a doctor's practice. Cf. also primary care and secondary care .
	Outpatient Doctors	Ambulatory Doctors, Ambulatory Care, Out-patient	Health care e.g. in a doctor's office, clinic, or day surgery center (not in-patient, not in hospital). Outpatient care today is also called ambulatory care.

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	Over prescribing	Overprescribing	If a doctor prescribes more pharmaceuticals than comparable doctors (e.g. with similar patient groups or in the same region). The measure of over prescribing is of particular importance if the doctor has been approved a pharmaceutical budget .
OTC	Over-The-Counter Pharmaceutical	Non-prescription medicine, Over-The-Counter medicine	Pharmaceuticals which may be dispensed without a doctor's prescription being submitted and which are in some countries available via self-service in pharmacies a/o other retail outlets (e.g. drug stores). Selected OTC may be reimbursed for certain indications in some countries.
	Parallel Trade	Parallel Distribution	Parallel trades are products imported into one Member State from another (the "export" country) and placed on the market in the destination Member State, outside the manufacturer's or its licensed distributor's formal channels.
	Parapharmaceuticals		Parapharmaceuticals are substances or compounds which do not correspond to the legal definition of a pharmaceutical . They are in any event products which, by virtue of their composition, utilisation or presentation, are compatible with the dignity of the profession of pharmacist.
	Patient-specific Reimbursement		Eligibility for reimbursement depends on the pharmaceutical in question (either a pharmaceutical is considered as reimbursable or as non-reimbursable).
	Pay-back	Payback	A cost-containment measure; a financial mechanism that requires manufacturers to refund a part of their revenue to a payer (i.g. social health insurance / national health service) if sales exceed a previously determined or agreed target-budget.
	Percentage Co-payment	Co-insurance, Percentage Copayment	Cost-sharing in the form of a set proportion of the cost of a service or product. The patient pays a certain fixed proportion of the cost of a service or product, with the social health insurance / national health service paying the remaining proportion.
	Pharmaceutical	Medicine, Drug, Medicinal Product	Any active ingredient or combination product presented for treating or preventing disease in human beings or animals. Any active ingredient or combination product which may be administered to human beings or animals with a view to making a medical diagnosis or to restoring, correcting or modifying physiological functions in human beings or in animals is likewise considered a pharmaceutical.
	Pharmaceutical Budget		Pharmaceutical budgets are a cost-containment measure of third party payers . The maximum amount of money to be spent on pharmaceuticals in a specific region or period of time is fixed ex-ante.
	Pharmaceutical Equivalence		Pharmaceuticals are pharmaceutical equivalents if they contain the same amount of the same active ingredient(s) in the same dosage form; if they meet the same comparable standards; and if they are intended to be administered by the same route. However, pharmaceutical equivalence does not necessarily imply therapeutic equivalence as differences in the excipients and/or the manufacturing process can lead to differences in product performance.
	Pharmaceutical Form		The pharmaceutical- technological form in which an active ingredient is made available. Pharmaceuticals may be administered in solid form (e.g. tablets, powders), in semi-liquid form (e.g. ointments, pastes), in liquid form (e.g. drops, infusions) or in gaseous form (inhalation).
	Pharmaceutical Retailer	Dispensary	Umbrella term for facilities that dispense/sell pharmaceuticals (POM and OTC) to outpatients, e.g. private pharmacies, public pharmacies, POM dispensaries, self-dispensing doctors , hospital pharmacies, pharmacy outlets, medicine chests, drugstores, supermarkets etc. In most countries the dispensation of pharmaceuticals is regulated by law, e.g. stating that supermarkets or pharmacy outlets may only sell a limited range of OTC .
	Pharmacoeconomic Evaluation	Pharmacoeconomics, Pharmacoeconomic Guidelines	The comparative analysis of alternative courses of action in terms of both their costs and consequences.
	Pharmacological Class		Pharmaceuticals that have similar therapeutic effects and similar safety and tolerability, both in nature and extent; cf. therapeutic group
	Pharmacopoeia		Pharmacopoeia (literally, the art of the drug compounder), in its modern technical sense, is a book containing directions for the identification of samples and the preparation of combination products , and published by the authority of a government or a medical or pharmaceutical society.

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	Pharmacovigilance		Pharmacovigilance is the ongoing surveillance of product safety occurring throughout the product life cycle. The EU legislation definition is "The collection and scientific evaluation of adverse drug reactions (ADR), under normal conditions of use, for regulatory purposes. It includes collection of data on drug consumption as well as misuse and abuse".
	Pharmacy Margin		The gross profit of pharmacies expressed as a percentage of the pharmacy retail price .
	Pharmacy Mark Up	Pharmacy Mark-up	The gross profit of pharmacies expressed as a percentage of the pharmacy purchasing price .
PPP	Pharmacy Purchasing Price	Wholesale Price, Pharmacy Purchase Price; Wholesale Selling price	The price charged by wholesalers to the retailers (usually pharmacies). It includes any wholesale mark up .
PRP (gross)	Pharmacy Retail Price (gross)	Gross Pharmacy Selling Price	The price charged by retail pharmacists to the general public. It includes any pharmacy mark ups or dispensing fees and VAT .
PRP (net)	Pharmacy Retail Price (net)	Net Pharmacy Selling Price	The price charged by retail pharmacists to the general public. It includes any pharmacy mark ups or dispensing fees and does not include VAT (see also pharmacy retail price (gross)).
	POM Dispensary		Pharmaceutical retailers (e.g. pharmacies, self-dispensing doctors, hospital pharmacies) which are allowed to dispense prescription-only medicines (POM) to patients.
	Population-group-specific Reimbursement		Specific population groups (e.g. children, old-age pensioners) are eligible for pharmaceuticals , while others are not.
	Positive List	Formulary	List of pharmaceuticals that may be prescribed more or less without further conditions at the expense of a social health insurance / national health service . Cf. negative list .
	Prescription		Is an order mostly in written form (~ receipt) by a qualified health care professional to a pharmacist or other therapist for a pharmaceutical or treatment to be provided to their patient. One receipt may contain several items. The maximum number of items on a receipt is in many countries regulated.
	Prescription Fee	Prescription Charge	The patient has to pay a fixed fee for each prescription item dispensed on the expense of a third party payer , i.e. a form of fixed co-payment .
POM	Prescription-Only-Medicines		Pharmaceuticals that may be dispensed only on a doctor's prescription.
POM dispensary	Prescription-Only-Medicines dispensary		Umbrella term for facilities that are allowed to sell POM to outpatients , e.g. pharmacies, self-dispensing doctors , hospital pharmacies.
	Price Cap	Price Ceiling	A cost-containment measure which fixes ex-ante the maximum price of pharmaceutical , e.g. taking into consideration inflation rates and production cost. Companies are allowed to choose any price below this threshold and in exchange authorities refrain from further control of company data (profit margins, sales etc.).
	Price Control		Pricing policies where government authorities set the price of a pharmaceutical and/or indirectly influence it (e.g. statutory pricing, price negotiations, public procurement). Contrary to free pricing .
	Price Freeze		A popular cost-containment method. The price of a pharmaceutical is fixed at a given level, mostly for a predetermined period of time. Price freezes are sometimes based on agreements between pharmaceutical industry and authorities but in most cases it is done by law.
	Price Negotiations		A form of pricing procedure , where pharmaceutical prices are discussed / negotiated (e.g. between manufacturer and social health insurance / national health service).
	Price-Volume Agreement		Like a framework agreement , a volume control tool. The price of a pharmaceutical is agreed between public authorities and a manufacturer on the basis of a forecast volume of sales. If the actual sales volume exceeds the forecast, the price of the pharmaceutical is usually reviewed downwards.
	Pricing	Price Setting	The act of setting a price for a pharmaceutical .

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	Pricing Policies		Regulations or procedures used by government authorities to set or limit the amount paid by purchasers or the amount received by sellers (e.g. free pricing, statutory pricing, price negotiation and price control).
	Pricing Procedure		There are several methods of determining the price of a pharmaceutical: internal price referencing, external price referencing, cost-plus pricing and profit control.
	Primary Care		Primary care is provided by health care persons (mainly general practitioners , but also family nurses, gynaecologist or paediatrics) trained for and skilled in comprehensive first contact and continuing care for persons with any undiagnosed sign, symptom, or health concern not limited by problem origin (biological, behavioural, or social), organ system, or diagnosis. Primary care includes health promotion, disease prevention, health maintenance, counselling, patient education, diagnosis and treatment of acute and chronic illnesses in a variety of health care settings. Specialist care and hospital care are also referred to as secondary care and tertiary care .
	Private Pharmaceutical Expenses	Household Expenses	This term includes all forms of - out-of pocket payments (OPP): * percentage co-payment, * fixed co-payment, * deductibles as well as - direct payments.
	Private Pharmacy		Pharmacies in the ownership of private individual(s) (e.g. pharmacists) or private enterprises. Usually, community pharmacies are private pharmacies.
	Procurement		The act of purchasing a pharmaceutical by a third party payer via tendering , cf. public procurement
	Public Funds		Umbrella term for the State, regional and local government bodies as well as social health insurance / national health service .
	Public Pharmacy		Pharmacies in the ownership of public entities (e.g. local public authorities like communities and universities)
	Public Procurement		Buying pharmaceuticals by the state (e.g. public hospitals) on the basis of a tendering procedure, granting the contract to the best tenderer (pharmaceutical company / importer).
PPPs	Purchasing Power Parities		Purchasing power parities reflect the amount of a national currency that will buy the same basket of goods and services in a given country. When PPPs are taken into account, the differences between poorer countries vis a vis richer countries diminish.
QALY	Quality-adjusted-life year	Quality adjusted life year	A QALY is a measure of the value of health outcomes. Since health is a function of length of life and quality of life, the QALY was developed as an attempt to combine the value of these attributes into a single index number. The change in utility value induced by the treatment is multiplied by the duration of the treatment effect to provide the number of QALYs gained.
	Rebate		A partial refund following a purchase, possible in kind or in cash.
	Reference Price		cf. reference price system .
RPS	Reference Price System		The social health insurance / national health service determines a maximum price (= Reference Price) to be reimbursed for certain pharmaceuticals . On buying a pharmaceutical for which a fixed price / amount (~ the so-called reimbursement price) has been determined, the insured person must pay the difference between the fixed price / amount and the actual pharmacy retail price of the pharmaceutical in question, in addition to any fixed co-payment or percentage co-payment rates. Usually the reference price is the same for all pharmaceuticals in a given ATC 4 level and/or ATC 5 level group.
	Regional Taxation		Taxes collected/generated through single municipalities/communities or provinces/regions.
	Registration		Simplified licensing procedure (instead of marketing authorisation) that is foreseen for herbal pharmaceuticals .
	Reimbursable		Pharmaceuticals which are eligible for reimbursement
	Reimbursement		Reimbursement is the percentage of the reimbursement price (for a service or a pharmaceutical) which the social health insurance / national health service pays. So 100% reimbursement means that the social health insurance/national health service covers 100% of the reimbursement price / amount of a pharmaceutical or service except a possible prescription fee .

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	Reimbursement Category	Reimbursement Group	Pharmaceuticals eligible for reimbursement are often grouped according to selected characteristics, e.g. route of administration (oral, etc.), main indication (oncology, paediatric, etc.), ATC level, classification (hospital-only , etc.). In many countries different reimbursement rates are determined for different reimbursement categories.
	Reimbursement Price	Public Price	This price is the basis for reimbursement of pharmaceuticals in a health care system, i.e. the maximum amount paid for by a third party payer . The reimbursed amount can either be the full reimbursement price (like e.g. Austria) or a percentage share of the reimbursement price (e.g. in Denmark). In a reference price system the reimbursement price is lower than the full price of the pharmaceutical , leaving the patient to pay the difference privately (or through complementary voluntary health insurance).
	Reimbursement Rate		The percentage share of the price of a pharmaceutical or medicinal service, which is reimbursed/subsidised by social health insurance / national health service . The difference to the full price of the pharmaceutical or medicinal service is paid by the patients (out-of-pocket payment).
	Reimbursement Scheme	Key Reimbursement Scheme, General Reimbursement,	The reimbursement system which covers the majority of residents in a country, in some countries also referred to as "general" reimbursement .
	Risk-sharing Agreement		An agreement between public authorities and one manufacturer which links the price of a pharmaceutical to a defined risk. The risk can be a risk of inappropriate use (over prescribing compared to targeted population or prescription of inappropriate dosages) or can be related to the cost-effectiveness claimed by the manufacturer.
	Sales/Pharmacy Tax		A tax - other than VAT - levied by a state or city on the pharmacy retail price of an item, collected by the retailer.
	Secondary Care	Specialist Care, Ambulant Care	Summarizes all types of specialised medical treatment. In many countries specialist treatment is offered not only in ambulatory care but also in outpatient departments of hospitals (sometimes called 'Polyclinics'). Free access for patients is often only possible with referral from primary care services, otherwise patients are charged with out-of-pocket payments .
SD-doctors	Self-Dispensing Doctor	Dispensing Doctor	Physicians who have been granted the right to dispense pharmaceuticals to their patients.
	Self-Medication	Self Medication	Self-Medication refers to pharmaceuticals purchased without prescription.
	Sickness Fund	Social Insurance Institution	A single social health insurance institution. In some countries there are several sickness funds operating (Austria) or even competing each other (Germany). Some sickness funds are operating on a regional basis whereas others are limited to specific professional groups like farmers or self-employed persons.
SHI	Social Health Insurance	Health Insurance, Bismarck System	Social health insurance is a type of health care provision, often funded through insurance contributions by employers and employees as well as state subsidies. In many countries there are obligatory schemes for (<i>employed</i>) persons whose income does not exceed a certain amount/limit (= insurance obligation) in place. Social health insurance is often organised in different sickness funds - in some countries allowing the patient to select a sickness fund (Germany) whereas in others the membership is determined mandatory, e.g. depending on the type of occupation (e.g. Poland, Austria). In some Social Health Insurance countries persons with higher income as well as self-employed persons may opt for substitutive private health insurance . In addition to social health insurance in some countries voluntary health insurance , covering e.g. out-of pocket payments or allowing for free choice of doctors, is very popular.
	Statutory Pricing		Pricing system, where pharmaceutical prices are set on a regulatory basis (e.g. law, enactment, decree).
	Substitutive Private Health		Cf. voluntary health insurance .
SPC	Supplementary Protection Certificate		SPC gives original products a complementary period of market exclusivity beyond patent expiry to compensate for delays of marketing in the pharmaceutical sector. SPC are available in EU countries but such complementary protection exists in other countries.
	Switch		Reclassification of prescription-only-medicines to over-the-counter pharmaceuticals .
	Tertiary Care		Tertiary care services are provided by specialised hospitals or departments that are often linked to medical schools or teaching hospitals. They treat patients with complex conditions who have usually been referred by other hospitals or specialist doctors.
	Therapeutic Benefit		The effect conveyed on a patient following administration of a pharmaceutical which either restores, corrects or modifies a physiological function(s) for that patient.
	Therapeutic Group		Pharmaceuticals from the same pharmacological class , such as statins. See also ATC 4 level .

PPRI GLOSSARY

ABBREV	Preferred term	Synonym term	Definition/Description
	Therapeutically Interchangeable		An interchangeable pharmaceutical is one which is therapeutically equivalent to a original product , i.e. meaning that the two molecules both deliver a similar therapeutic benefit to patients.
	Third Party Payer		Any organisation, public or private, that pays or insures health care expenses for beneficiaries at the time at which they are patients, e.g. social health insurance or national health service or other public funds .
TD	Transparency Directive		Directive 89/105/EEC (of 21 December 1988) relates to the transparency of measures regulating the pricing of pharmaceuticals for human use and their inclusion in the scope of National social health insurance systems / national health service .
	Unbranded Generic		Cf. generic .
VAT	Value Added Tax		A sales tax levied on the sale of goods and services (compulsory for EU Member States). The VAT rate of pharmaceuticals in the EU is often lower than the standard VAT rate of 15%.
VHI	Voluntary Health Insurance		Health Insurance that is taken up and paid for at the discretion of individuals or employers on behalf of individuals. VHI can be offered by public or quasi-public bodies and by for-profit (commercial) and non-profit private organisations. In the European context, VHI can be classified in three different ways: - Substitutive Private Health Insurance provides cover that would otherwise be available provided by state. In a social health insurance system people who have no insurance obligation (in some countries e.g. self-employed) may opt for substitutive private health Insurance. - Complementary VHI provides cover for services excluded or not fully covered by the state (e.g. dental care), including cover for co-payments imposed by the statutory health care system. - Supplementary VHI provides cover for faster access and increased consumer choice.
	Wholesale		The sale of goods (e.g. pharmaceuticals) to retailers (e.g. pharmacies).
	Wholesale Margin		Gross profit of wholesalers, expressed as a percentage of the pharmacy purchasing price .
	Wholesale Mark up	Wholesale Mark-up	Gross profit of wholesalers, expressed as a percentage of the ex-factory price .