D YOUNG®CO INTELLECTUAL PROPERTY

European patent prosecution & litigation: UK

Antony Craggs & Garreth Duncan

Speakers



Antony Craggs arc@dyoung.com Partner, Solicitor London, UK



Garreth Duncan gad@dyoung.com Partner, Patent Attorney Southampton, UK

Any questions? Email us directly or contact registrations@dyoung.com.

Slides and a recording of this webinar will be emailed to you later this week.

Agenda

Agenda

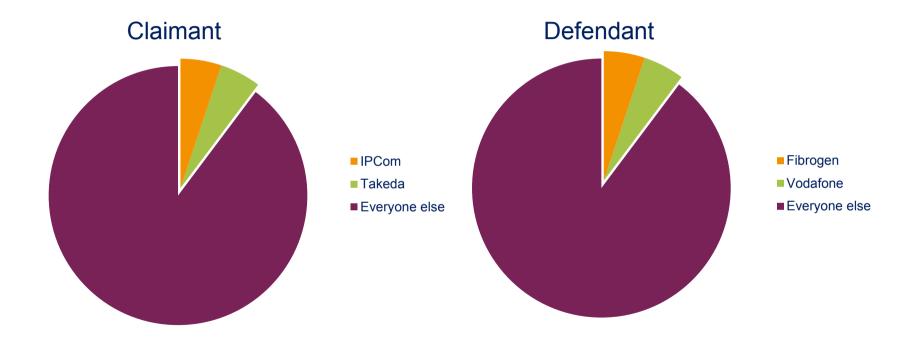
Annual stocktake

Doctrine of equivalents

SEPs & FRAND

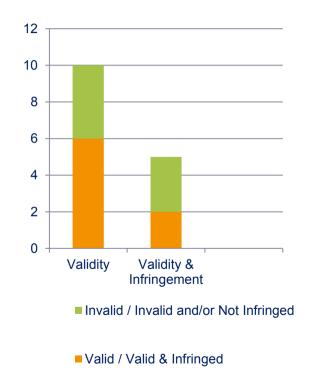
Plausibility and infringement of Swiss Form Claims

Stocktake: 2018



Stocktake: 2018

- In 2018, approximately 13 came to trial.
- Of the 13 cases, 10 trials addressed validity.
 Of these, 6 found one or more patents to be valid.
- Of the 13 cases, 9 trials addressed validity and infringement, with 6 finding at least one patent to be valid and infringed.
- The remaining cases were directed to SPCs or licences.



Stocktake

Patents (2010-2018)		
	103	
Valid		
	39	37.86%
Invalid		
	64	62.14%
Infringed		
	56	54.37%
Non-infringed		
	35	33.98%
Valid and infringed		
	30	29.13%

Stocktake

Patents (2010-2018)						
Judge	Arnold J		Birss J		Carr J	
Patents	30		28		8	
Valid	8	26.67%	16	57.14%	6	75.00%
Invalid	22	73.33%	12	42.86%	2	25.00%
Infringed	16	53.33%	16	57.14%	6	75.00%
Non-infringed	12	40.00%	8	28.57%	0	0.00%
Valid and infringed	8	26.67%	11	39.29%	6	75.00%

Protocol on the Interpretation of Article 69 EPC

Article 1

General principles

Article 69 should not be interpreted as meaning that the extent of the protection conferred by a European patent is to be understood as that defined by the strict, literal meaning of the wording used in the claims, the description and drawings being employed only for the purpose of resolving an ambiguity found in the claims. Nor should it be taken to mean that the claims serve only as a guideline and that the actual protection conferred may extend to what, from a consideration of the description and drawings by a person skilled in the art, the patent proprietor has contemplated. On the contrary, it is to be interpreted as defining a position between these extremes which combines a fair protection for the patent proprietor with a reasonable degree of legal certainty for third parties.

Article 2

Equivalents

For the purpose of determining the extent of protection conferred by a European patent, due account shall be taken of any element which is equivalent to an element specified in the claims.

- What is the doctrine of equivalents?
 - i) Does the variant infringe any of the claims as a matter of <u>normal</u> construction; and, if not
 - ii) Does the variant nonetheless infringe because it varies from the invention in a way or ways which is or are immaterial?
 - If the answer to either issue is "yes", there is an infringement;
 otherwise, there is not.

- What is the doctrine of equivalents?
 - i) Notwithstanding that it is not within the <u>literal</u> meaning of the relevant claim(s) of the patent, does the variant achieve substantially the same result in substantially the same way as the invention, ie the inventive concept revealed by the patent?
 - ii) Would it be obvious to the person skilled in the art, reading the patent at the priority date, but knowing that the variant achieves substantially the same result as the invention, that it does so in substantially the same way as the invention?

- What is the doctrine of equivalents?
 - o iii) Would such a reader of the patent have concluded that the patentee nonetheless intended that strict compliance with the literal meaning of the relevant claim(s) of the patent was an essential requirement of the invention?
- To establish infringement by doctrine of equivalents:
 - Questions i) and ii): "yes"
 - Question iii): "no"

- What issues remained to be determined?
 - Purposive construction
 - Impact on validity
 - Use of prosecution history
 - Limits of doctrine of equivalents

- Purposive construction
 - Generics v. Teva
 - Fisher & Paykel v. ResMed
 - Illumina v Premaitha
 - Liqwd v. L'Oréal
 - Icescape v. Ice-World

- Impact on validity
 - Generics v. Teva
- Use of prosecution history
 - Icescape v. IceWorld
- Limits of doctrine of equivalents
 - Icescape v. IceWorld

- What is a Standard Essential Patent (SEP)?
- What is FRAND?
- What were the issues?
 - Does the court have jurisdiction to determine essentiality?
 - Does the court have jurisdiction to determine FRAND?

- [Cont.]
 - What is FRAND?
 - Geography
 - Portfolio
 - SEPs and non-SEPs?
 - Length?
 - o Rate?

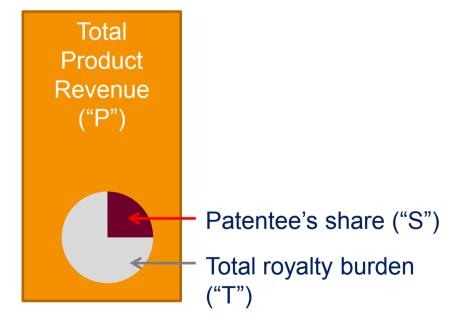
- Unwired Planet v. Huawei:
 - 2012: Licence between Ericsson and Huawei expires
 - 2012: Ericsson disposes of a portion of its patent portfolio to Unwired Planet
 - 2013-2014: Unwired Planet seeks to procure licence from Huawei

- Unwired Planet v. Huawei:
 - 2014: Unwired Planet commences patent infringement proceedings against Huawei (and others)
 - 2015-2016: Five technical trials listed (three proceed, two are stayed): two of the six patents are found to be valid, essential and infringed
 - 2016: FRAND trial
 - 2018: FRAND appeal

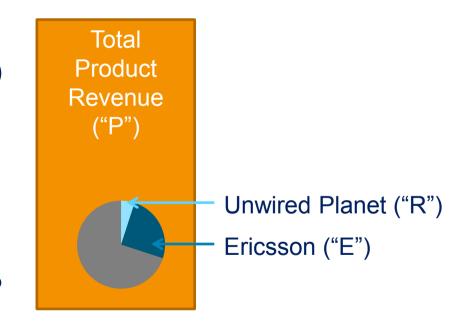
- Unwired Planet v. Huawei:
 - English Court can determine FRAND terms
 - It is FRAND to offer a worldwide portfolio licence
 - It is not FRAND to offer a SEP and non-SEP portfolio licence
 - Court concluded that if the prospective licensee does not take a global FRAND licence it will award a UK injunction.

- FRAND Royalty Rate
 - Approaches:
 - Top Down
 - Comparable Licences

- Top Down:
 - Determine total royalty burden ("T")
 - Determine patentee's share of burden ("S")
 - FRAND Rate: (SxT)xP
 - Example: (25%x10%)xP



- Comparable Licences:
 - Determine Ericsson rate ("E")
 - Determine Unwired Planet portion of Ericsson portfolio ("R")
 - FRAND rate: (RxE)xP
 - Example: (7.69% x 0.8%) x P



- Non-discrimination
 - What does the non-discrimination limb of FRAND mean?

Unwired Planet Share (S) for Handsets				
	Unwired Planet Patent	HPA denominator	Adjusted denominator	S
2G	2	350	154	1.30%
3G	1	1089	479	0.21%
4G	6	1812	800	0.75%
Multimode				
2G/3G				0.57%
2G/3G/4G				0.70%

Unwired Planet Share (S) for Infrastructure				
	Unwired Planet Patent	HPA denominator	Adjusted denominator	S
2G	1	305	134	0.75%
3G	2	886	390	0.51%
4G	7	1554	684	1.02%

o Future?



- Swiss form claims (allowed by EPO EBA in G5/83)
 - "Use of substance X <u>for the manufacture of a medicament</u> for the treatment of indication Y"
 - Not permitted since 2011 by EPO but will be around till 2036!
 - EPC 2000 form "Substance X for use in treating indication Y"
 - not considered by Sup Ct

- Pregabalin (Lyrica®) sold for three indications:
 - Epilepsy
 - Generalised anxiety disorder
 - Neuropathic pain
- Basic compound patent:
 - Disclosed epilepsy and anxiety indications
 - Expired in 2013

- Patent in suit:
 - Claim 1: "Use of [pregabalin] for the preparation of a pharmaceutical composition for treating pain"
 - Claim 3: Limited to treating neuropathic pain
- Action:
 - Generics launched "skinny label" product omitting pain indication
 - Revocation / Counterclaim for infringement

- First instance:
 - Invalid for insufficiency
 - No infringement
- Court of appeal:
 - Invalid for insufficiency
 - (Obiter) infringement

Plausibility and Swiss Form Claims

- "Plausibility" in application as filed
 - EPO apply on inventive step
 - UK Courts apply on insufficiency
- Prevent speculative claims justified solely by post-filing data
- If not "plausible" from application as filed, later filed data can't remedy
- "Plausibility" is assessed across entire scope of claim

Plausibility and Swiss Form Claims

- Swiss form (second medical use) claims
 - Application as filed must contain some disclosure as to how or why known product could be expected to work for new indication
 - Data not essential
 - Credible theory with scientific reasoning sufficient
- Threshold not met for either claim 1 or claim 3 patent revoked

Plausibility – Squeeze on Second Medical use Claims?

- Waiting for clinical trial results before filing rarely an option
 - Disclosure of some trial protocols to regulators and patients mandatory
- Pre-clinical data not always available
- Scientific reasoning a double edged sword?
 - good for sufficiency, bad for inventive step?
- Other drugs in same class?

Infringement and Swiss Form Claims

- No infringement
- Direct infringement
 - Lord Sumption and Lord Reed "as manufactured and packaged"
 - Lord Mance "positive exclusion" may be required
 - Lord Hodge and Lord Briggs "subjective intention"
- Indirect infringement

- What is an Arrow declaration?
- What is a Gillette defence?
- Why is an Arrow declaration useful?

- Glaxo Group v. Vectura
 - Background
 - 2010: Licence for Staniforth (but not Non-Assert Patents)
 - 2016: Staniforth expires (no licence taken for Non-Assert Patents)
 - 2017: Vectura sues in USA
 - 2018: GSK commence revocation proceedings in UK

- Glaxo Group v. Vectura
 - Strike out application
 - Undertaking not to enforce the remaining Non Assert Patents
 - Application unsuccessful
 - No undertaking in relation to 415
 - Patents invalid & not infringed
 - Gillette defence

- Fujifilm v AbbVie:
 - i) justice to the claimant;
 - ii) justice to the defendant;
 - iii) whether the declaration will serve a useful purpose; and
 - iv) whether or not there are any other special reasons why the court should or should not grant the declaration.

- Glaxo Group v. Vectura
 - Vectura may be able to establish infringement with reformulated divisionals
 - Open question as to whether patents with differently formulated claims were obvious or not
 - Vectura had declined to give an undertaking in relation to 415
- Declaration granted

Questions and future webinars



Antony Craggs arc@dyoung.com Partner, Solicitor London, UK



Garreth Duncan gad@dyoung.com Partner, Patent Attorney Southampton, UK

Any questions?
Email us directly or contact us by email at registrations@dyoung.com

Listen again?
Slides and a recording this webinar will be emailed to you later this week. We welcome invitations to present this webinar and

Bookmark us
Keep up to date with all our IP news at
www.dyoung.com/knowledgebank

other IP subjects of interest.