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Page 581

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FIFTY YEARS OF PATENT LAW: THE TOP TEN DEVELOPMENTS

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I.	INTR	ODUCTION	582
II.	TOP TEN DEVELOPMENTS		582
	10.	HATCH-WAXMAN ACT	582
	9.	PATENT LITIGATION'S "EXCESS LUGGAGE" (BEST MODE,	
		Inequitable Conduct, Willful Infringement and At	TORNEY
		Fee Awards)	583
	8.	REMEDIES: INJUNCTIONS AND DAMAGES	586
	7.	Venue	587
	6.	STANDARD OF REVIEW	588
	5.	CLAIM INTERPRETATION AND APPLICATION	590
	4.	OBVIOUSNESS; WRITTEN DESCRIPTION	594
	3.	Creation of Federal Circuit	597
	2.	SECTION 101 INELIGIBILITY: DEATH AND REVIVAL	599
	1.	AIA: Post-Issuance Review	603
	0.	THE PROMISED BONUS: GROWTH IN THE PROFESSION	604

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I. INTRODUCTION

The fifty years of the *AIPLA Quarterly Journal* coincide with my professional engagement in patent law. In the early 1970s, I began the research that led to the 1978 publication of a five-volume treatise. After its publication, the treatise required regular updating to reflect patent law developments. Little did I know that I would still be at it after these many years (and 185 "releases").

Many basic features of the U.S. patent system remained fundamentally the same during the past fifty years. But there have been significant developments. Below I review the top ten in reverse order of significance¹ and add a bonus development.

II. TOP TEN DEVELOPMENTS

10. HATCH-WAXMAN ACT

In 1984, Congress passed the "Hatch-Waxman Act." It could well have been titled "The Patent Lawyers and Litigators Full Employment Act." And it should have been given a Pulitzer Prize for linguistic complexity.

The Act negotiated a compromise between the generic drug industry (which wanted a procedure to obtain quicker and easier Food and Drug Administration ("FDA") approval of generic drugs) and the brand drug industry (which wanted extension of patent term for regulatory delays in approval of new drugs).

Some would contend that the Act fostered disrespect for the patent system on both sides. In § 271(e)(2), it authorized a patent owner to sue a generic that filed an abbreviated new drug application ("ANDA") to obtain FDA approval immediately, i.e., before FDA approval to market the drug.³ That gave generics an

For prior top tens, see Donald Chisum, "Top Ten Intellectual Property Cases of the Federal Circuit 1982–2002, Twentieth Anniversary Judicial Conference of the Court of Appeals for the Federal Circuit (April 8, 2002), *in* 217 F.R.D. at 548 and Donald Chisum, The Year in Review: The Patent and Trademark Decisions of the Court of Appeals for the Federal Circuit, Second Annual Judicial Conference of the Court of Appeals for the Federal Circuit (April 26, 1984), *in* 104 F.R.D. at 207.

See Drug Price Competition and Patent Term Restoration Act of 1984, Pub. L. No. 98-417, 98 Stat. 1585. On Hatch-Waxman, see 5 DONALD CHISUM, CHISUM ON PATENTS § 16.03[1][d] (2022).

³ See 35 U.S.C. § 271(e)(2). For a Supreme Court discussion of Section 271(e)(2), see Eli Lilly & Co. v. Medtronic, Inc., 496 U.S. 661, 678 (1990).

incentive to challenge even strong patents on drugs because they could contest a patent's scope and validity without risking a potentially enormous damage award.⁴ The Act also gave patent owners an incentive to obtain weak, incremental patents on drugs that were vulnerable to a validity challenge. By suing for infringement thereof, patent owners obtained an automatic thirty-month stay of FDA approval of a generic drug.

Could we have both a healthy, patent-supported research drug industry and a cost-saving generic industry without the litigation-inducing Hatch-Waxman Act ANDA suit provision? Likely yes. With conventional patent enforcement, a generic, reasonably confident in its position, could develop, obtain FDA approval, and "test" a product in the market. A patent owner, confident in its position, could sue and make an appropriate showing for a preliminary injunction. One might examine how all this works in countries without the ANDA suit procedure.

9. PATENT LITIGATION'S "EXCESS LUGGAGE" (BEST MODE,
INEQUITABLE CONDUCT, WILLFUL INFRINGEMENT AND ATTORNEY
FEE AWARDS)

Patent litigation should focus primarily on two basic issues: infringement (claim scope) and validity (patentability). During the fifty-year period, four additional issues distracted from the basics. The issues were excess luggage from the start and later became too uncertain. Cases on the issues added many pages to *Chisum on Patents*. In time, each was either trimmed or simplified by the Supreme Court, the Federal Circuit, or Congress.

First was the statutory "best mode" requirement.⁵ A patent otherwise valid and infringed could be invalidated because it failed to disclose an inventor's subjective preference on some aspect of a claimed invention. Cases parsed the details of the requirement, such whose "contemplation" mattered⁶ and on what

⁴ See GlaxoSmithKline LLC v. Teva Pharms. USA, Inc., 7 F.4th 1320, 1340 (Fed. Cir. 2021). In *Glaxo*, which was a "regular" infringement suit, not a Hatch-Waxman Act ANDA suit, a divided Federal Circuit panel affirmed infringement of a treatment method patent. *Id.* A jury awarded the patent owner \$234,110,000 in damages for lost profits (even though the generic's sales amounted to only \$74,500,000 and there were other generic equivalents available). *Id.*

On best mode, see 3 Chisum on Patents, supra note 2, § 7.05.

⁶ See, e.g., Glaxo Inc. v. Novopharm Ltd., 52 F.3d 1043, 1049 (Fed. Cir. 1995).

date.⁷ A favorite defense tactic was to depose an inventor early. Tell me about your invention? The inventor might boast of various advantageous features. Then ask: where is that in the patent? I can't find it. Voila! Summary judgment of invalidity!

In the 2011 American Invents Act ("AIA"), Congress stepped in. The best mode disclosure requirement remained but not as a requirement for priority to a prior application, an invalidity defense in an infringement suit, or a basis for post grant review. That solution was odd. Can you omit a best mode or not? But the amendment quashed best mode as a litigation complicator. The Federal Circuit issued no precedential decisions on best mode from 2011 through 2022.

Second was inequitable conduct. Inventors and their representatives owe a duty of candor in prosecuting a patent application in the Patent and Trademark Office ("PTO"). That includes disclosing known material prior art. But, like best mode, inequitable conduct became a routine, overly pleaded defense against a patent otherwise apparently valid and infringed. One distractive aspect was that the defense focused attention on attorney conduct during prosecution. The Federal Circuit stepped in with an en banc ruling, *Therasense, Inc. v. Becton, Dickinson & Co.*, which raised the bar on the showings of materiality (it must be "but for") and deceptive intent. Thereafter, the defense continued but usually only on a well-supported, factual basis. The Supreme Court has yet to weigh in on the Federal Circuit's standard for the inequitable conduct defense.

Third was willful infringement.¹² The Patent Act authorized a district court to increase damages up to three times actual damages.¹³ It set no standard

See, e.g., Transco Prods. Inc. v. Performance Contracting, Inc. 38 F.3d 551, 554 (Fed. Cir. 1994).

See Leahy-Smith America Invents Act, Pub. L. No. 112-29, § 15(a), 125 Stat. 284, 328 (2011) (amending 35 U.S.C. § 282(b)(3)(A) to state "the failure to disclose the best mode shall not be a basis on which any claim of a patent may be canceled or held invalid or otherwise unenforceable").

On inequitable conduct, see 4 CHISUM ON PATENTS, *supra* note 2, § 11.03b[4] and 6A CHISUM ON PATENTS, *supra* note 2, § 19.03.

See Therasense, Inc. v. Becton, Dickinson & Co., 649 F.3d 1276, 1292 (Fed. Cir. 2011). For a discussion of *Therasense*, see 6A CHISUM ON PATENTS, supra note 2, §§ 19.03[3][e][v], 19.03[4][g][iii].

See, e.g., Belcher Pharms., LLC v. Hospira, Inc., 11 F.4th 1345, 1351 (Fed. Cir. 2021).

On willful infringement, see 6A CHISUM ON PATENTS, supra note 2, § 20.03[4][b][v][K].

¹³ See 35 U.S.C. § 284.

but had been construed as proper for willful infringement. Like inequitable conduct, willful infringement came to be charged routinely, in this instance, by the patent owner against an accused infringer. Again, attention focused frequently on attorney conduct: was an attorney's noninfringement or invalidity opinion competent? The Federal Circuit responded with two en banc decisions. Knorr v. Dana Corp. barred adverse inferences from an infringer's failure to offer an exculpatory opinion of counsel.14 In re Seagate Technology LLC abolished the "affirmative duty of care," clarified that a waiver of the attorney-client privilege arising from reliance on an advice-of-counsel defense to a charge of willful infringement did not extend to communications with, and work product of, trial counsel, and established a "two-prong" test for willful infringement. 15 First was an objective prong (acting despite "objectively high likelihood" that the acts infringed valid patent). Second was a subjective prong (known or should have known).¹⁶ Both were provable by clear and convincing evidence. In Halo Electronics, Inc. v. Pulse Electronics, Inc., a unanimous Supreme Court rejected the two-prong test and the high proof standard and simplified the willfulness inquiry.¹⁷ The Court reminded us that its early decisions construed the statutory authority to increase damages as allowing a discretionary increase as punishment for willful infringement.¹⁸ Willful infringement was just that: deliberate acts in disregard of known patent rights. The Federal Circuit's *In re* Seagate threshold allowed a willful infringer to escape enhanced damages by mustering "a reasonable (even though unsuccessful) defense at the infringement trial" even when the infringer did not act based on the defense.¹⁹ Despite rejecting In re Seagate's threshold, the Court did not purport to restore the Federal Circuit's pre-Seagate "affirmative duty of care" standard, which was effectively one of negligence, not willfulness, and which allowed patent owners to routinely assert willful infringement.

See Knorr-Bremse Systeme Fuer Nutzfahrzeuge GMBH v. Dana Corp., 383 F.3d 1337, 1346 (Fed. Cir. 2004).

¹⁵ See In re Seagate Tech. LLC, 497 F.3d 1360, 1371 (Fed. Cir. 2007) (en banc).

¹⁶ See id.

See Halo Elecs., Inc. v. Pulse Elecs., Inc., 579 U.S. 93, 103–04 (2016). For a discussion of Halo, see 6A CHISUM ON PATENTS, supra note 2, § 20.03[4][b][x].

¹⁸ See Halo, 579 U.S. at 103–04.

¹⁹ *Id.* at 105.

Fourth was attorney fee awards.²⁰ The Patent Act authorized a district court to award fees in "exceptional cases" to a prevailing party.²¹ Especially in response to suits by non-practicing patent owners, exonerated accused infringers routinely sought fees. Similarly to its cases on willfulness, the Federal Circuit adopted a per se test with a threshold.²² The cases required either litigation misconduct or a showing of both subjective bad faith and objective baselessness to find a case "exceptional." This test tended to shield non-prevailing patent owners just as *In re* Seagate shielded non-prevailing accused infringers. And, again, in *Octane Fitness, LLC v. ICON Health & Fitness, Inc.*²³ and *Highmark Inc. v. Allcare Health Management System, Inc.*,²⁴ the Supreme Court rejected the Federal Circuit's approach as too rigid. An "exceptional case" was "simply one that stands out from others with respect to the substantive strength of a party's litigating position (considering both the governing law and the facts of the case) or the unreasonable manner in which the case was litigated."²⁵ The Court emphasized that an attorney fee award lay heavily within the discretion of a district court.²⁶

8. Remedies: Injunctions and Damages

The primary remedies for patent infringement are an injunction and damages. Two Supreme Court cases were milestones.

On injunctions, the Court, in *eBay Inc. v. MercExchange LLC*, emphasized that there was no "general rule," unique to patent cases, that a permanent injunction must issue, absent extraordinary circumstances, once a patent is adjudged infringed and not invalid.²⁷ Rather, in determining whether to grant a permanent injunction, a court should apply traditional equitable principles. These

On attorney fee awards, see 6A CHISUM ON PATENTS, *supra* note 2, § 20.03[4][c].

²¹ 35 U.S.C. § 285.

E.g., Brooks Furniture Mfg., Inc. v. Dutailier Int'l, Inc., 393 F.3d 1378, 1381 (Fed. Cir. 2005), abrogated by Octane Fitness, LLC v. ICON Health & Fitness, Inc., 572 U.S. 545 (2014).

Octane Fitness, LLC v. ICON Health & Fitness, Inc., 572 U.S. 545, 554 (2014) ("The Federal Circuit's formulation is overly rigid.").

Highmark Inc. v. Allcare Health Mgmt. Sys., Inc., 572 U.S. 559, 563 (2014) ("Our holding in *Octane* settles this case").

²⁵ Octane Fitness, 572 U.S. at 554.

²⁶ See id. at 557; Highmark, 572 U.S. at 564.

²⁷ eBay Inc. v. MercExchange, LLC, 547 U.S. 388, 393–94 (2006)

included whether the patent owner would suffer irreparable injury, whether "remedies available at law, such as monetary damages, are inadequate to compensate for that injury," "the balance of hardships," and whether "the public interest would not be disserved by a permanent injunction." ²⁸ In *eBay*, after a jury found a patent valid and infringed, a district denied a permanent injunction, relying, inter alia, on the fact the patent owner did not practice the patent. A Federal Circuit panel reversed, applying a "general rule" that a court should issue a permanent injunction against patent infringement absent exceptional circumstances. The Court held that both courts erred.

On damages, the Court, in SCA Hygiene Products Aktiebolag v. First Quality Baby Products, LLC, overruling the Federal Circuit's 1992 en banc decision in A.C. Aukerman Co. v. R.L. Chaides Construction Co.,²⁹ held that the traditional equitable defense of "laches," which barred pre-suit damages if a claimant unreasonably delayed suing to the prejudice of a defendant, could not preclude a patent owner's claims for damages for infringements occurring in the six-year pre-suit period prescribed by § 286. The Court left in place the separate defense of equitable estoppel, which could bar all relief against an infringer based on a patent owner's misleading representation that it would not sue. The policy implications of SCA Hygiene are dubious. For example, a company having good faith questions about a patent's scope might communicate them to the patent owner. The patent owner could choose not to respond (and thus avoid any prospect of a declaratory judgment suit) and wait six years to sue while the company built a business around a technology later found infringing.

7. Venue

The Judicial Code restricts venue in a patent infringement suit to either the state of an accused infringer's residence or a district in which it had both a regular and established place of business *and* committed an act of infringement.³⁰ That contrasted with the general venue statute that permitted, via a definition of "residence," a suit against a corporation in any district in which it was subject to personal jurisdiction. Typically, a corporation distributing a product nationally

²⁸ *Id.* at 391.

A.C. Aukerman Co. v. R.L. Chaides Constr. Co., 960 F.2d 1020 (Fed. Cir. 1992) (en banc), abrogated by SCA Hygiene Products Aktiebolag v. First Quality Baby Prods., LLC, 137 S. Ct. 954, 967 (2017).

³⁰ 28 U.S.C. § 1400(b). On venue in patent suits, see 7 CHISUM ON PATENTS, *supra* note 2, § 21.02[2].

would be subject to jurisdiction in most if not all districts in an infringement suit concerning the product.

In VE Holding Corp. v. Johnson Gas Appliance Co., the Federal Circuit changed the then-accepted understanding that the special venue statute precluded a patent owner from suing in its home base or in another preferred district.³¹ The decision facilitated a trend for patent owners, especially non-practicing entities, to file suits in districts, such as the Eastern District of Texas, in which the court offered a quick path to trial.

In *TC Heartland LLC v. Kraft Foods Group Brands LLC*, the Supreme Court blew the whistle, holding that the expansive definition of a domestic corporation's "residence" in the general venue statute did *not* apply to the exclusive venue provision for patent infringement suits.³²

Thus began a process wherein the Federal Circuit faced new and difficult issues on what constituted a place³³ and where an infringing act occurred.³⁴ Those issues had been irrelevant during the *VE Holding* period (1990 to 2017). In some ways, this starting-from-scratch process resembled what the Federal Circuit did for many patent law issues in the early years after its creation in 1982.

6. STANDARD OF REVIEW

In systems for resolving disputes over the facts, the law or both, it is common to provide a review structure, i.e., appeals. The "standard of review" on appeal can be critical. Is it "de novo," i.e., the review starts from scratch, or subject to some form of deference to an initial decider? In the fifty-year span, the Supreme Court addressed review of patent decisions in three areas.

The first area concerned court review of decisions by the PTO and in particular, the PTO's findings of fact in the course of examining patent applications. From its beginning in 1983, the Federal Circuit applied the same

VE Holding Corp. v. Johnson Gas Appliance Co., 917 F.2d 1574, 1575 (Fed. Cir. 1990), abrogated by TC Heartland LLC v. Kraft Foods Grp. Brands LLC, 137 S. Ct. 1514 (2017).

³² *TC Heartland*, 137 S. Ct. at 1517.

See, e.g., Andra Grp., LP v. Victoria's Secret Stores, LLC, 6 F.4th 1283, 1287 (Fed. Cir. 2021); In re Google LLC, 949 F.3d 1338, 1343 (Fed. Cir. 2020); In re Cray Inc., 871 F.3d 1355, 1359–60 (Fed. Cir. 2017).

See, e.g., Celgene Corp. v. Mylan Pharms. Inc., 17 F.4th 1111, 1120–22 (Fed. Cir. 2021); Valeant Pharms. N. Am. LLC v. Mylan Pharms. Inc., 978 F.3d 1374, 1375 (Fed. Cir. 2020).

"clear error" standard used for reviewing district court findings.³⁵ The PTO campaigned for a more deferential "substantial evidence" standard as provided in the Administrative Procedure Act ("APA"). In *In re* Lueders, Judge Giles Rich provided an extensive historical analysis defending the clear error standard.³⁶ In *Dickinson v. Zurko*, the Federal Circuit sitting en banc rejected the PTO's arguments.³⁷ The Supreme Court reversed.³⁸

The APA substantial evidence review standard acquired even greater significance when, in the AIA, Congress expanded post-issuance review by the PTO, including inter partes review. The AIA contained a provision that ostensibly precluded judicial review of a decision by the PTO's director to institute an inter partes review. Three Supreme Court decisions grappled with that provision.³⁹

The second area concerned the allocation of decisional authority between the judge and a jury in a patent infringement suit. The Seventh Amendment to the U.S. Constitution guarantees a right to trial by jury in civil cases. That has long been understood to include patent infringement suits seeking damages. In *Markman v. Westview Instruments, Inc.*, the Supreme Court held that the interpretation of a patent claim was "a matter of law reserved entirely for the court."⁴⁰ For historic reasons, there was no right to have a jury resolve a dispute about the meaning of a claim, even when the patent owner offered expert testimony on the meaning of a "term of art." The decision induced creation a new pre-trial procedure in infringement suits, the "*Markman* hearing" on claim construction.⁴¹

The third area concerned the standard of appellate review of claim construction, whether by a district court, the PTO, or the International Trade Commission. The Supreme Court's *Markman* did not resolve that question; it only held that construction was not for a jury, and did not necessarily exclude appellate

In re Caveney, 761 F.2d 671, 674 (Fed. Cir. 1985) ("reviews PTO findings under the clearly erroneous standard").

³⁶ *In re* Lueders, 111 F.3d 1569, 1574–78 (Fed. Cir. 1997).

³⁷ See In re Zurko, 142 F.3d 1447, 1459 (Fed. Cir. 1998) (en banc), rev'd, 527 U.S. 150 (1999).

³⁸ See Dickinson v. Zurko, 527 U.S. 150, 165 (1999).

See Cuozzo Speed Techs., LLC v. Lee, 579 U.S. 261, 265–66 (2016); see also SAS Inst., Inc. v. Iancu, 138 S. Ct. 1348, 1352 (2018); Thryv, Inc. v. Click-To-Call Techs., LP, 140 S. Ct. 1367, 1370 (2020).

⁴⁰ Markman v. Westview Instruments, Inc., 517 U.S. 370, 372 (1996).

On Markman hearings, see 5A CHISUM ON PATENTS, supra note 2, § 18.06[2][a][vii][A].

deference to a trial court's resolution of factual issues pertinent to construction. The Federal Circuit determined that its review was "de novo," i.e., without deference, but individual judges protested that such review violated the general rules requiring deference to trial court findings of fact. The Circuit affirmed its de novo position in en banc decisions in 1998 and 2014. But, yet again, the Supreme Court disagreed. In *Teva Pharmaceuticals USA, Inc. v. Sandoz, Inc.*, it held that, when a construction of a term of art in a patent claim has "evidentiary underpinnings" and a district court resolves an underlying factual dispute, the Federal Circuit on appeal must review the district court's fact finding under the "clear error" standard. ⁴³

The disagreement was less than might be apparent. In *Teva*, the Court agreed that the "ultimate construction" of a patent claim, based on any fact findings, remained a "legal conclusion" reviewable de novo.⁴⁴ And the Court agreed that when a district court reviewed "only evidence intrinsic to the patent (the patent claims and specifications, along with the patent's prosecution history)," its determination was one of law and the Federal Circuit would "review that construction de novo."⁴⁵ Since *Teva*, in most cases, the Federal Circuit has determined that a claim construction was resolvable by reference to the "intrinsic evidence," that resort to extrinsic evidence that would require fact finding was unnecessary, and thus that review was without deference.

5. CLAIM INTERPRETATION AND APPLICATION

A patent's claims define the invention for all purposes in patent law – for infringement, of course, but also for patentability and other issues, such as inventorship. Three landmark decisions, one by the Federal Circuit and two by the Supreme Court, addressed the interpretation and application of claims.

In *Phillips v. AWH Corp.*, the Federal Circuit addressed en banc the basic approach to interpreting a claim, including the relative weight to "intrinsic evidence" (claim language, specification (written description) and prosecution history) and "extrinsic evidence" (including expert testimony).⁴⁶ Leading up to

⁴² See Cybor Corp. v. FAS Techs., Inc., 138 F.3d 1448, 1451 (Fed. Cir. 1998) (en banc); Lighting Ballast Control LLC v. Philips Elecs. N. Am. Corp., 744 F.3d 1272, 1276–77 (Fed. Cir. 2014) (en banc), vacated & remanded, 135 S. Ct. 831 (2015).

⁴³ Teva Pharms. USA, Inc. v. Sandoz, Inc., 574 U.S. 318, 321–22 (2015).

⁴⁴ Id. at 332-33.

⁴⁵ *Id.* at 331.

⁴⁶ Phillips v. AWH Corp., 415 F.3d 1303, 1317 (Fed. Cir. 2005) (en banc).

Phillips, three-judge panel decisions oscillated between two opposing schools.⁴⁷ One emphasized the context of claim language, including particularly the specification and its examples.⁴⁸ The other emphasized ordinary meaning of a claim term, often derived from dictionary definitions, and allowed deviation from the meaning only when a patent clearly redefined the term or the patent owner had unmistakably disavowed ordinary meaning.⁴⁹ In *Phillips*, Judge Bryson, in a thorough opinion, synthesized elements from both schools, emphasizing the specification's importance but warning against reading limitations from the specification's examples into the claims. *Phillips* had a calming effect, but remnants of the two schools occasionally surfaced.⁵⁰

The Supreme Court's decisions addressed the doctrine of equivalents and a significant restraint on its use, prosecution history estoppel.

The doctrine of equivalents has a venerable history in the Court, dating back to the 1853 *Winans v. Denmead* case, in which a Court majority of five justices held a patent claim to a railroad car with a container in the shape of the frustum of a cone (i.e., circular) infringed by an accused infringer's car with an eight-sided container because the latter "substantially," though not literally, embodied "the patentee's mode of operation" and thereby attained "the same kind of result."⁵¹ Four justices dissented. The division was that which is always raised in discussions of the doctrine. Strictly enforcing literal claim scope potentially undermines the ability of competitors to determine what a patent covers but risks undermining the value of patents as incentives for innovation.

Almost a century later, the Court again applied the doctrine of equivalents in *Graver Tank & Manufacturing Co. v. Linde Air Products Co.*⁵²

See 6 Chisum on Patents, supra note 2, § 18.07.

⁴⁸ E.g., Vitronics Corp. v. Conceptronic, Inc., 90 F.3d 1576, 1582 (Fed. Cir. 1996).

⁴⁹ E.g., Texas Digit. Sys., Inc. v. Telegenix, Inc., 308 F.3d 1193, 1202 (Fed. Cir. 2002).

Compare Thorner v. Sony Comput. Ent. Am. LLC, 669 F.3d 1362 (Fed. Cir. 2012) (holding that terms should be defined according to their ordinary meaning), with Columbia Univ. v. Symantec Corp., 811 F.3d 1359 (Fed. Cir. 2016) (holding that a claim construction should be understood through the entire context of the patent and specification).

⁵¹ Winans v. Denmead, 56 U.S. (15 How.) 330, 344 (1853).

See Graver Tank & Mfg. Co., Inc. v. Linde Air Prods. Co., 339 U.S. 605, 610 (1950).

Thereafter, the regional circuits applied the doctrine, using a "range of equivalents" standard, which accorded greater equivalents to patents on "pioneer" inventions and lesser equivalents to those on mere improvements.⁵³

After 1982, the Federal Circuit paid little attention to the range idea. However, its judges disputed other aspects of the doctrine, in particular, what should be the standard for equivalence, whether it should be limited to instances in which an infringer copied the patented technology (as opposed to developing its technology independently), and what the role of a jury should be. The dispute culminated in an en banc decision with multiple opinions.⁵⁴

In *Warner-Jenkinson Co. v. Hilton Davis Chemical Co.*, the Supreme Court granted review and yet again affirmed the doctrine's viability.⁵⁵ The Court held that "intent plays no role in the application of the doctrine of equivalents."⁵⁶ Equivalency was determined "at the time of infringement, not at the time the patent was issued."⁵⁷ The Court sympathized with the concerns of the Federal Circuit dissenters that the doctrine had "taken on a life of its own, unbounded by the patent claims."⁵⁸ To alleviate those concerns, it adopted the suggestion of dissenting Federal Circuit Judge Helen Nies that equivalency be applied on an element-by-element basis, not "as a whole."⁵⁹

On the role of juries, the Court dropped a highly significant footnote providing "guidance, not a specific mandate" about "the concern over unreviewability due to black-box jury verdicts." Summary judgment of non-infringement should be entered when "no reasonable jury could determine two elements to be equivalent." Additionally, "various legal limitations" on the doctrine of equivalents could be determined by summary judgment or by motions

⁵⁷ *Id.* at 37.

⁵⁸ *Id.* at 28–29.

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See John Zink Co. v. Nat'l Airoil Burner Co., 613 F.2d 547, 556–58 (5th Cir. 1980); Nelson v. Batson, 322 F.2d 132, 136 (9th Cir. 1963).

⁵⁴ See Hilton Davis Chem. Co. v. Warner-Jenkinson Co., 62 F.3d 1512, 1514 (Fed. Cir. 1995), rev'd & remanded, 520 U.S. 17 (1997). For a discussion of Hilton Davis, see 5B CHISUM ON PATENTS, supra note 2, § 18.04a[1][a][iii][G].

⁵⁵ See Warner-Jenkinson Co. v. Hilton Davis Chem. Co., 520 U.S. 17, 40 (1997).

⁵⁶ *Id.* at 36.

⁵⁹ See id. at 29.

⁶⁰ See id. at 39 n.8.

⁶¹ See id.

at trial.⁶² The limitations included "prosecution history estoppel" or "a theory of equivalence" that "would entirely vitiate a particular claim element."⁶³

On prosecution history estoppel, the Court rejected an accused infringer's argument that there should be a "rigid" rule under which a surrender of subject matter, such as by an amendment narrowing a broad claim through adding a limitation in response to a PTO examiner rejection, precluded recapture of any part of the surrendered subject matter. The Court indicated that there was a rebuttable "presumption" of surrender.

After *Warner-Jenkinson*, the Federal Circuit judges debated the impact of that decision on prosecution history estoppel. In *Festo Corp. v. Shoketsu Kinzoku Kogyo Kabushiki Co.*, the Supreme Court reviewed two of the Federal Circuit's rules on prosecution history estoppel. ⁶⁴

The first rule postulated that an estoppel arises when an applicant by amendment narrows a claim limitation for any reason relating to statutory requirements for obtaining a patent. The Court confirmed that rule. It rejected an argument that estoppel should arise only from amendments made to distinguish prior art and not from amendments made to meet § 112's disclosure and clarity requirements.

The second rule, described as an "absolute" or "complete" bar rule, dictated that a patentee's act of amending a claim limitation during prosecution created an estoppel that bars every equivalent to the amended claim limitation. The Supreme Court rejected the Federal Circuit's absolute bar rule, deeming it an impermissible "new rule" that would unfairly diminish the scope and value of existing patents. But it also recognized the uncertainty caused by a "flexible bar" approach to the estopping effect of claim amendments. Accordingly, the Court held that if a patentee narrows a claim by adding or amending a claim limitation, it should be presumed to have surrendered all equivalents to the amended claim limitation. The patentee may rebut the presumption by showing that:

[t]he equivalent [was] unforeseeable at the time of the application; the rationale underlying the amendment [bore] no more than a tangential relation to the equivalent in question; or there [was] some other reason suggesting that the patentee could not

⁶² See id.

⁶³ See id.

See Festo Corp. v. Shoketsu Kinzoku Kogyo Kabushiki Co., 535 U.S. 722, 730 (2002).

reasonably [have been] expected to have described the insubstantial substitute in question.⁶⁵

After *Festo*, the "no more than tangential relation" rebuttal criteria proved to be most difficult one to apply consistently.⁶⁶

4. OBVIOUSNESS; WRITTEN DESCRIPTION

That a patentable invention should be more than an obvious modification or combination of prior art teachings can hardly be questioned. It has always been the key legal condition for patentability.

Until the 1952 Act, the patent statutes articulated expressly only a requirement that a claimed invention be "new." But, starting with the 1851 *Hotchkiss v. Greenwood* case, the Supreme Court read "invention" and "new" to include an non-obviousness component.⁶⁷ Unfortunately, instead of applying that straightforward proposition on a case-specific basis, courts purported to establish various negative and positive rules on what was and was not an "invention." The Court's application of the "invention" requirement proceeded historically through patent-favorable and patent-hostile periods, the period from about 1930 through 1950 being particularly hostile.⁶⁹

In the 1952 Act, Congress added § 103, expressly stating the would-not-have-been-obvious-to-a-skilled-artisan standard. Some argued that the intent was to "lower" the Court's high standard. In the 1966 *Graham v. John Deere Co.* trilogy, the Court disagreed and suggested that such would have been unconstitutional.⁷⁰ It acknowledged, however, that § 103, "when followed realistically," by both the Patent Office and the courts, was "a more practical test of patentability": "The

See Pharma Tech Sols., Inc. v. Lifescan, Inc., 942 F.3d 1372, 1380–81 (Fed. Cir. 2019); Eli Lilly & Co. v. Hospira, Inc., 933 F.3d 1320, 1333 (Fed. Cir. 2019); Ajinomoto Co., Inc. v. Int'l Trade Comm'n, 932 F.3d 1342, 1354 (Fed. Cir. 2019).

⁶⁵ See id. at 740-41.

⁶⁷ See Hotchkiss v. Greenwood, 52 U.S. 248, 271–72 (1851).

See 2A Chisum on Patents, supra note 2, $\S 5.04[5]$.

⁶⁹ See 2 Chisum on Patents, supra note 2, § 5.02[3].

See Graham v. John Deere Co., 383 U.S. 1, 16–17 (1966); United States v. Adams, 383 U.S. 39, 48 (1966).

emphasis on non-obviousness is one of inquiry, not quality, and, as such, comports with the constitutional strictures."⁷¹

After *Graham*, the regional circuits laced the § 103 condition with sundry non-statutory variants. For example, the Ninth Circuit, sitting en banc and resolving conflicting panel opinions, indicated that the test for a "combination patent" was "unusual or surprising results," not "synergism."⁷²

After its creation in 1982 with essentially exclusive appellate jurisdiction, the Federal Circuit swept away such variants.⁷³ It did so even though two post-*Graham* Supreme Court decisions, *Anderson's-Black Rock, Inc. v. Pavement Salvage Co.*⁷⁴ and *Sakraida v. Ag Pro, Inc.*⁷⁵ had seemingly reaffirmed pre-1952 Act "invention" rules.

In KSR International Co. v. Teleflex Inc., the Supreme Court held the Federal Circuit itself guilty of applying a "rigid" rule on obviousness, one requiring that the prior art provide a "teaching, suggestion or motivation" to combine prior art elements. The Court's opinion discussed, without disapproval, Anderson's-Black Rock and Sakraida. It also rejected a general rule against using "obvious to try."

After KSR, Federal Circuit panels reiterated that KSR did not eliminate a requirement that there be a reason (or motivation) to combine or modify the prior art. They also focused on a requirement that there be a reasonable expectation of success.

Given the importance of the non-obviousness condition, the frequency with which it arises, and the difficulty of applying it in various fields of technology, it is not surprising that differences of opinion arose among Federal Circuit judges and were reflected in panel opinions with varying if not conflicting

⁷¹ *Graham*, 383 U.S. at 17.

⁷² Sarkisian v. Winn-Proof Corp., 688 F.2d 647, 651 (9th Cir. 1982) (en banc).

See, e.g., Am. Hoist & Derrick Co. v. Sowa & Sons, Inc., 725 F.2d 1350, 1360–61 (Fed. Cir. 1984) (determining that synergism is essential for patentability, although predecessor courts rejected that notion); Richdel, Inc. v. Sunspool, Corp., 714 F.2d 1573, 1579–80 (Fed. Cir. 1983); Stratoflex, Inc. v. Aeroquip Corp., 713 F.2d 1530, 1540 (Fed. Cir. 1983).

⁷⁴ See Anderson's-Black Rock, Inc. v. Pavement Salvage Co., 396 U.S. 57, 61–62 (1969).

⁷⁵ See Sakraida v. Ag Pro, Inc., 425 U.S. 273, 279–80 (1976).

See KSR Int'l Co. v. Teleflex Inc., 550 U.S. 398, 407 (2007). On KSR, see 2 CHISUM ON PATENTS, supra note 2, § 5.02[9].

⁷⁷ See KSR Int'l, 550 U.S. at 414.

language. The opinions cited different statements in *KSR* to support conclusions of obviousness and no obviousness.

To date, only one en banc decision has attempted half-heartedly to resolve these conflicts. In *Apple Inc. v. Samsung Electronics Co.*, the Federal Circuit overturned a panel decision that had reversed a jury verdict of non-obviousness.⁷⁸ The majority stressed that it was *not* addressing any "important legal questions about the inner workings of the law of obviousness."⁷⁹ The majority did affirm the relevance of "objective evidence" of non-obviousness, including commercial success, industry praise, copying, and long-felt need.⁸⁰

Despite its recognition that there should be no special, extra-statutory rules on what is an "invention" in applying the non-obviousness condition for patentability, the Federal Circuit effectively created such a rule in applying the § 112 requirement that a patent specification include, as of its priority date, a written description of "the invention" in addition to an enabling disclosure of how to make and use it. The written description requirement undoubtedly plays a key role in preventing an applicant from retroactively claiming to have invented subject matter by changing claims through post-filing amendments or in continuing applications.81 In this priority policing mode, a written description analysis compares a claim to the description. However, in Regents of University of California v. Eli Lilly & Co., a Federal Circuit panel held that an application specification could fail to provide a written description of the invention recited in a claim in that specification (i.e., an "original claim").82 The panel announced its new rule despite the explicit provision in § 112 making claims part of the specification and the unquestioned principle that a claim defines an invention. Regents reasoned that a specification failed to show "possession" of an invention, even one explicitly claimed, when it delineated the invention generically and in terms of function rather than structure, providing only a "mere wish" or "plan" for obtaining a claimed invention rather than examples of it (working or constructive).83 The subsequent 2010 en banc Ariad Pharmaceuticals, Inc. v. Eli Lilly

See Apple Inc. v. Samsung Elecs. Co., 839 F.3d 1034, 1039–40, 1057 (Fed. Cir. 2016) (en banc).

⁷⁹ *Id.* at 1039.

⁸⁰ Id.

⁸¹ See Quake v. Lo, 928 F.3d 1365, 1373 (Fed. Cir. 2019).

Regents of Univ. of Cal. v. Eli Lilly & Co., 119 F.3d 1559, 1562 (Fed. Cir. 1997).

⁸³ *Id.* at 1566.

& Co. decision confirmed *Regents*.⁸⁴ Thus, decisions after *Regents* applied the § 112 written description requirement to hold unpatentable claims deemed to be generic and functional even when the claims were in a prior application as filed and even assuming that the application provided an enabling disclosure.⁸⁵

With § 112, as with § 103, the proper course would have stuck to the statute and not indulged in judicial speculation on what was an "invention" and whether an invention, which was described by a claim and supported by an enabling disclosure, was sufficiently completed. No doubt unduly broad, functional claims should be held improper. However, the separate statutory requirement of enablement was available and had been interpreted by the Supreme Court since 1854 as precluding such claims. To avoid confusion, there should be one set of rules for evaluating the adequacy of disclosure to support a broad claim, not two. *Non multiplicantur res extra necessitatem*.

3. Creation of Federal Circuit

In 1982, Congress created the Federal Circuit by combining the seven judgeships of the Court of Claims with the five of the Court of Customs and Patent Appeals. It gave the Federal Circuit near exclusive appellate jurisdiction over patent cases, including appeals from district court decisions as well as those from the PTO and the International Trade Commission. No change in appellate structure had attained such significance since Congress created the intermediate "regional" courts of appeal in 1892 (which greatly relieved the Supreme Court of routine appeals in patent cases).⁸⁸

Why did Congress do it? And has the experiment succeeded?

Suggestions that Congress gave the Federal Circuit exclusive jurisdiction to "strengthen" the patent system overlooked a fundamental principle of the Constitution: an Article III court is independent of the political branches of

See Ariad Pharms., Inc. v. Eli Lilly & Co., 598 F.3d 1336, 1349 (Fed. Cir. 2010) (en banc).

See Nuvo Pharms. (Ireland) Designated Activity Co. v. Dr. Reddy's Lab'ys Inc., 923 F.3d 1368, 1381–82 (Fed. Cir. 2019).

See Juno Therapeutics, Inc. v. Kite Pharma, Inc., 10 F.4th 1330, 1342 (Fed. Cir. 2021) (overturning jury verdict of no written description violation by broad claim to chimeric antigen receptor ("CAR") T-cell therapy, the verdict including a \$1,200,322,551.50 damage award).

⁸⁷ See O'Reilly v. Morse, 56 U.S. (15 How.) 62, 112 (1854).

See 2 Chisum on Patents, *supra* note 2, § 5.02[2] (examining the 1892 Act and its significance for patent litigation).

government and cannot be given any task other than deciding judicial cases applying the law. If Congress desired to strengthen patents, it needed to have amended the statutes. It did not and has not (with the exception of 1984 and 2011 amendments that altered what constitutes prior art).⁸⁹

Did creation of the Federal Circuit nevertheless have the *effect* of strengthening patents? After the 1970s, the percentage of patents held not invalid rose. But whatever "anti-patent" bias was shown in some of the regional circuits might well have changed in the 1980s (without creation of a Federal Circuit) due to increased perception of the value of intellectual property, especially with the growing impact of international trade on the U.S. economy.

Two related and more defensible purposes for taking patent cases out of the hands of the regional circuits were to increase consistency and predictability in the application of patent law and to reduce "forum-shopping," that is, parties seeking to maneuver a case into a district court in a favorable circuit. The regional circuits had reputations, whether deserved or not, for widely-varying attitudes about patentability. For example, patents were almost always upheld in the Fifth Circuit and almost never in the Eighth. Indeed, a conflict in the rulings of the Fifth and Eighth Circuits on the validity of the same patent caused the Supreme Court to grant certiorari in the 1966 *Graham* case. The Court held that neither circuit had applied the correct test for obviousness.

The Federal Circuit's exclusive jurisdiction eliminated circuit shopping. However, another form of forum of shopping developed: to obtain a favored district court, such as the Eastern or Western districts in Texas, a phenomenon the Circuit had facilitated with its *VE Holdings* venue ruling.

The Federal Circuit provided less consistency and predictability than might have been hoped for because of panel variation. Cases are decided by rotating panels of three of the up to twelve judges. Prior opinions might show that judges N and S tended to find patents not obvious but judges D and P tended to find them obvious. A given patent's chances would then be better before a panel of N, S and D than before one with N, D and P. Parties could not "shop" for a panel as they previously shopped the circuits because a party could not predict which judges would be on a panel. But panel variation ran counter to a fundamental principle of the law: like cases should be decided alike and without regard to which judges are on a panel.

Varying views of the judges also left some important issues of law unsolved or subject to conflicting resolution for long periods until resolved en banc or by the Supreme Court. Examples included, as discussed above, the tests

⁸⁹ See id. §§ 5.02[7], 5.03[3][c][vi]; 3A CHISUM ON PATENTS, supra note 2, § 9.05[4] (examining the 1984 amendment).

for the doctrine of equivalents, prosecution history estoppel, and the proper approach to claim construction. A particularly stark example was a schism on whether a patent's product-by-process claim was infringed when an accused infringer made the same product using a different process. The schism arose in 1992.⁹⁰ It went unresolved until 2009!⁹¹

Another example is the on-sale bar to patentability. The judges split on whether a reduction to practice was required of an invention to be "on sale." The Supreme Court finally resolved the issue in *Pfaff v. Wells Electronics*. ⁹² A further controversy concerned whether experimental use negating a public use bar ended with a reduction to practice. In an en banc decision, *Medicines Co. v. Hospira, Inc.*, the Federal Circuit declined to address the issue. ⁹³

Other divisions remain subtle but important. When does a generic drug maker's label induce infringement of a treatment method claim?⁹⁴ When does an unclaimed feature in a successful product preclude a presumption that a nexus connects the product's success and the merits of the claimed invention?⁹⁵

2. Section 101 Ineligibility: Death and Revival

A development of undeniable importance during the fifty-year period was the Supreme Court's erratic and irrational interpretation of the § 101 definition of patent eligible subject matter. The Court's ineligibility decisions were

Compare Atl. Thermoplastics Co., Inc. v. Faytex Corp., 970 F.2d 834, 837 (Fed. Cir. 1992), with Scripps Clinic & Rsch. Found. v. Genentech, Inc., 927 F.2d 1565, 1583 (Fed. Cir. 1991) (agreeing with Scripps that the same product being made with a different process would still infringe under precedent), overruled by Abbott Lab'ys v. Sandoz, Inc., 566 F.3d 1282, 1293 (Fed. Cir. 2009).

See Pfaff v. Wells Elecs., 525 U.S. 55, 67–68 (1998); see also 2A CHISUM ON PATENTS, supra note 2, §§ 6.02[2][1], 6.02[6][a].

⁹¹ See Abbott, 566 F.3d at 1293.

⁹³ See Medicines Co. v. Hospira, Inc., 827 F.3d 1363, 1381 (Fed. Cir. 2016) (en banc); see also 2A CHISUM ON PATENTS, supra note 2, § 6.02[6][d][ii].

Ompare GlaxoSmithKline LLC v. Teva Pharms. USA, Inc., 7 F.4th 1320 (Fed. Cir. 2021), with Takeda Pharms. U.S.A., Inc. v. West-Ward Pharm. Corp., 785 F.3d 625 (Fed. Cir. 2015).

⁹⁵ Compare Quanergy Sys., Inc. v. Velodyne Lidar USA, Inc., 24 F.4th 1406, 1417 (Fed. Cir. 2022), with Teva Pharms. Int'l GMBH v. Eli Lilly & Co., 8 F.4th 1349, 1361 (Fed. Cir. 2021), and Fox Factory, Inc. v. SRAM, LLC, 944 F.3d 1366, 1374 (Fed. Cir. 2019).

surprising as well as disturbing because the statutory language, which covers any machine, manufacture, composition of matter, or process, has remained essentially unchanged since 1791.

In the 1972 *Gottschalk v. Benson* case, a truncated Court held that a patent's claim to a mathematical algorithm useful for converting numbers was an unpatentable abstract idea. ⁹⁶ The short opinion by Justice Douglas was unanimous but only six justices participated.

The *Benson* opinion was poorly reasoned, as I demonstrated in a 1986 law review article.⁹⁷ The decision effectively validated the Patent Office's de facto policy not to allow "software" patents, a policy encouraged by a computer hardware manufacturer that later accumulated a huge portfolio of such patents.⁹⁸ The Office's policy probably had long-term negative effects on the quality of such patents when they did begin to emerge. If the Office had earlier examined and issued appropriately narrow software patents, their full disclosures would have been available as prior art in examining later applications.

At almost precisely the time I finished work on the treatise, a line of decisions by the Court of Customs and Patent Appeals had effectively cabined *Benson*. However, just as the treatise appeared in 1978, the Supreme Court extended *Benson*, in *Parker v. Flook*, a 5-4 decision, holding that an unpatentable mathematical formula did not become patentable subject matter by the addition of "conventional, post-solution applications." Thus were ineligible applicant Flook's claims to a method for updating the value of an "alarm limit" on a variable involved in a process of catalytic chemical conversion of hydrocarbons. That a specific improvement in an industrial process should be per se excluded from patenting was a truly disturbing result.

But despair at perpetuated irrationality abated for a time when, shortly after *Flook*, the Court rendered two decisions. In *Diamond v. Chakrabarty*, it held

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⁹⁶ See Gottschalk v. Benson, 409 U.S. 63, 73 (1972).

⁹⁷ See Donald S. Chisum, The Patentability of Algorithms, 47 U. PITT. L. REV. 959, 961–62 (1986).

See Donald S. Chisum, Patenting Intangible Methods: Revisiting Benson (1972) After Bilski (2010), 27 SANTA CLARA COMPUT. & HIGH TECH. L.J. 445, 447–48 (2011).

⁹⁹ See, e.g., In re Musgrave, 431 F.2d 882, 894 (C.C.P.A. 1970) (representing the culminating decision in this line of decisions from the Court of Customs and Patent Appeals).

¹⁰⁰ See Parker v. Flook, 437 U.S. 584, 585, 596 (1978).

¹⁰¹ See id. at 585–86.

that the PTO could not reject as ineligible claims to a genetically-modified bacterium. ¹⁰² Importantly, the Court noted that it was up to Congress to provide exceptions to the Patent Act's broad § 101 definition of patentable subject matter. The dissent did not dispute that § 101 was broad but argued only that two plant protection statutes indicated a Congressional intent to protect only some kinds of "animate inventions." ¹⁰³

In *Diamond v. Diehr*, another 5-4 decision, the Court held eligible a claim to a process for curing synthetic rubber, which included in one of its steps the use of a long-known mathematical formula and a programmed digital computer.¹⁰⁴ Essentially, in relevant respects, the claims in *Diehr* were indistinguishable from those in *Flook*. If anything, there was a stronger case for the patentability of the *Flook* claims because the claim's calculation method or algorithm was asserted to be new whereas the formula in the *Diehr* claims was admittedly known. Unfortunately, apparently in deference to the principle of stare decisis (precedent), the majority in *Diehr* nominally distinguished *Flook* rather than overruling it (and *Benson*) as inconsistent with the principle *Chakrabarty* recognized.

Thus, by the time the Federal Circuit came into being in 1982, *Benson* was, effectively, dead. And so it remained until a trio of Supreme Court decisions revived it: *Bilski v. Kappos*, ¹⁰⁵ *Mayo Collaborative Services v. Prometheus Laboratories, Inc.*, ¹⁰⁶ and *Alice Corp. v. CLS Bank International*. ¹⁰⁷ *Mayo* added "law of nature" to "abstract idea" as an ineligible concept. ¹⁰⁸ In contrast, in a partial victory for rationality, the Court in *Association for Molecular Pathology v. Myriad Genetics, Inc.* held that isolated DNA was not eligible because it was a product of nature but that synthetically-created (complementary) DNA was eligible. ¹⁰⁹

This is not the place to excessively parse the Court's cases. But note an interesting fact: the 1978 to 1981 cases all involved appeals from a reluctant PTO (as did *Bilski*) whereas the 2012-2014 cases involved invalidating issued patents.

¹⁰² See Diamond v. Chakrabarty, 447 U.S. 303, 318 (1980).

¹⁰³ See id. at 319 (Brennan, J. dissenting).

¹⁰⁴ See Diamond v. Diehr, 450 U. S. 175, 191–92 (1981).

¹⁰⁵ See Bilski v. Kappos, 561 U.S. 593, 612 (2010).

See Mayo Collaborative Servs. v. Prometheus Lab'ys, Inc., 566 U.S. 66, 72 (2012).

¹⁰⁷ See Alice Corp. Pty. Ltd. v. CLS Bank Int'l, 573 U.S. 208, 219 (2014).

¹⁰⁸ See Mayo, 566 U.S. at 78.

See Ass'n for Molecular Pathology v. Myriad Genetics, Inc., 569 U.S. 576, 580 (2013).

Sufficing to show the uncertainty of what the Court expounded on ineligibility is the number of Federal Circuit decisions reaching varying results. 110 After six years of applying and attempting to clarify Alice and Mayo, the Federal Circuit in 2020 issued eleven precedential opinions: seven held claims ineligible, 111 six eligible. 112

Will the Court revisit *Benson-Mayo-Alice*? A good candidate would have been *American Axle & Manufacturing, Inc. v. Neapco Holdings LLC*, in which a panel majority held that a method of making a car part was an ineligible natural law.¹¹³ Unfortunately, the Court recently denied certiorari review of that case.

If the Court does eventually take another § 101 eligible subject matter case, one can hope that the Court will not attempt to simply refine and clarify. Instead, it should drive a stake into the heart of *Benson*.

The Supreme Court is the primary culprit in the crime of § 101 confusion, but Congress has been compliant. In several sections of the AIA, it evidenced awareness of the problem but explicitly declined to address it. For example, the AIA's Section 18 provided for post-grant review of business method patents but cautioned: "Nothing in this section shall be construed as amending or interpreting categories of patent-eligible subject matter set forth under section 101 of title 35, United States Code." 114

For a discussion of all these cases, see 1 Chisum on Patents, *supra* note 2, § 1.03.

See In re Rudy, 956 F.3d 1379, 1380 (Fed. Cir. 2020); Bozeman Fin. LLC v. Fed. Rsrv. Bank of Atlanta, 955 F.3d 971, 974 (Fed. Cir. 2020); Customedia Techs., LLC v. Dish Network Corp., 951 F.3d 1359, 1360–61 (Fed. Cir. 2020); Simio, LLC v. Flexsim Software Prods., 983 F.3d 1353, 1356 (Fed. Cir. 2020); Am. Axle & Mfg., Inc. v. Neapco Holdings LLC, 967 F.3d 1285, 1299 (Fed. Cir. 2020); Elec. Commc'n v. ShoppersChoice.com, LLC, 958 F.3d 1178, 1180 (Fed. Cir. 2020); Ericsson Inc. v. TCL Commc'n Tech., 955 F.3d 1317, 1320 (Fed. Cir. 2020).

^{See TecSec, Inc. v. Adobe Inc., 978 F.3d 1278, 1282 (Fed. Cir. 2020); XY, LLC v. Trans Ova Genetics, LC, 968 F.3d 1323, 1326 (Fed. Cir. 2020); Packet Intel. LLC v. NetScout Sys., Inc., 965 F.3d 1299, 1303 (Fed. Cir. 2020); Uniloc USA, Inc. v. LG Elecs. USA, Inc., 957 F.3d 1303, 1305 (Fed. Cir. 2020); CardioNet, LLC v. InfoBionic, Inc., 955 F.3d 1358, 1362 (Fed. Cir. 2020); Illumina, Inc. v. Ariosa Diagnostics, 952 F.3d 1367 (Fed. Cir. 2020), modified, 967 F.3d 1319, 1321 (Fed. Cir. 2020).}

¹¹³ See Am. Axle, 967 F.3d at 1293–94 (concluding that a claim applying Hooke's law was directed to a law of nature).

¹¹⁴ Leahy-Smith America Invents Act § 18(e).

1. AIA: Post-Issuance Review

The 2011 enactment of the AIA by Congress was the number one development in U.S. patent law over the past fifty years.¹¹⁵

On substantive patent law, the AIA prospectively switched from a first-to-invent to a first-to-file priority system and revised the definition of prior art in § 102. In that respect, the AIA was a third stage in the shift of the U.S. patent system toward the model adopted by most other countries. The first stage was the 1995 adoption of the twenty-year-from-effective-filing date patent term to replace the prior seventeen-year-from-issuance term. The second stage was the 1999 adoption of eighteen-month publication of patent applications. Those changes ended the unfortunate phenomenon of "submarine" patents issuing many years after their filing date. However, the changes were prospective, and, for over two decades, patents continued to issue with seventeen-year terms based on pre-June 8, 1995, filing dates. 116

Even more significant than its substantive law change were the AIA's provisions on post-issuance review by a PTO Board.

The AIA's importance is confirmed by one simple fact. The Supreme Court decides few cases at all and very few on patent law, but the Court has a sense for "where the action is." In the first eight years of AIA post-grant review, it

Integrating a major statutory revision into a multiple volume treatise presents challenges. When Congress enacted a new copyright statute in 1976, Professor Nimmer chose to stop revising his original version of Nimmer on Copyrights and took the time to prepare a second edition. The AIA did not so comprehensively change patent law, and the vast bulk of case law on patent law remains applicable. Therefore, I prepared a special section entitled "America Invents Act of 2011: Analysis and Cross-References," which analyzed in detail the statute with its legislative history and the PTO's implementing regulations. That section remains unchanged. I also added sections in the relative parts of the existing chapters. For example, § 11.07[5] covers inter partes and post-grant review. It is regularly revised to account for case law and for changes. It shows how quickly the body of case law addressing procedural and jurisdictional issues on IPR and PGR has grown.

See, e.g., Immunex Corp. v. Sandoz Inc., 964 F.3d 1049, 1054 (Fed. Cir. 2020) (patent issuing 2011 based on 1990 priority application and May 1995 divisional application).

granted certiorari in six cases: five on aspects of post-issuance review, 117 and one on whether the AIA altered the § 102 "on sale" bar (holding that it did not). 118

A major attraction of inter partes review ("IPR") and post-grant review ("PGR") to a challenger and potential accused infringer is the opportunity to have an adjudication of issues of patentability (anticipation and obviousness), without discovery on the full ranges of issues in an infringement suit and before an expert tribunal instead of a jury in a district court suit.

In enacting the AIA's post-issuance procedures, Congress expressed its hope that they would, unlike the prior inter partes reexamination procedure, "serve as an effective and efficient alternative to often costly and protracted district court litigation." ¹¹⁹ Have they succeeded?

0. The Promised Bonus: Growth in the Profession

And now the bonus. A significant development over the past fifty years was the growth and change in the profession. Before, patent practice was concentrated in relatively small firms located primarily in a few cities. Today, it is vastly larger and more diverse. The growth was attributable in a significant part to the emergence of commercial biotechnology and, less positively, to the proliferation of patent litigation, including suits by "non-practicing entities."

Of particular relevance to my work as a scholar is the extent of academic focus on the patent system. As of the 1970s, there were few law schools that offered even a single course covering patent law, and there were almost no full-time professors who listed patent law as among their interests. Now, there are many schools offering programs and multiple courses.

¹¹⁷ See Cuozzo Speed Techs., LLC v. Lee, 579 U.S. 261, 266 (2016) (holding that the decision of whether to institute review is not appealable, and that the agency may issue reasonable regulations about how to conduct post-grant review); SAS Inst., Inc. v. Iancu, 138 S. Ct. 1348, 1352–53 (2018) (holding that the Patent Office must resolve all claims in the case—not just some of them); Oil States Energy Serv., LLC v. Greenes Energy Grp., LLC, 138 S. Ct. 1365, 1370 (2018) (holding that post grant proceedings at the PTO do not violate Article III or the seventh amendment); Return Mail, Inc. v. U.S. Postal Serv., 139 S. Ct. 1853, 1859 (2019) (holding that a federal agency is not a "person" who may challenge patent validity post-issuance); Thryv, Inc. v. Click-To-Call Techs., LP, 140 S. Ct. 1367, 1370 (2020) (holding that a party's failure to timely institute review at the PTAB is not appealable).

Helsinn Healthcare S.A. v. Teva Pharms. USA, Inc. 139 S. Ct. 628, 630 (2019).

¹¹⁹ H.R. REP. No. 112-98 at 45 (2011).

As a young law professor, I began work on a treatise on United States patent law rather than concentrate on the type of law review articles generally expected for advancement in legal academia. I had become fascinated by the history of the patent system and its role in the workings of the federal judiciary but was frustrated by the absence of up-to-date treatises and reference texts comparable to those on copyright, trademark, bankruptcy, and other areas of federal law. But today, we are blessed with an outpouring of legal scholarship on patents by full time professors and others. 120

Particularly significant to me is the excellent two-volume treatise by my partner at the Academy (and spouse), Janice Mueller. See Janice M. Mueller, Mueller on Patent Law (Full Court Press 2020).