- Magazine
 - Current/Back Issues
 - **Features**

 - Columns
 - <u>Digital Editions</u>
 - Subscribe Now
 - Advertise Now
- News
- All Companies
 ALL CATEGORIES
 - Industry Associations
 - Company Capabilities
 - Add Your Company
- Supply Chain
 - 3D/Additive Manufacturing
 - Contract Manufacturing
 - <u>Electronics</u>
 - Machining & Laser Processing
 - Materials
 - Molding
 - Packaging & Sterilization
 - R&D & Design
 - Software & IT
 - <u>Testing</u>
 - Tubing & Extrusion
- **Device Sectors**
 - Cardiovascular
 - **Diagnostics**
 - <u>Digital Health</u>
 - Neurological
 - Patient Monitoring
 - o Surgical
- Top 30 Company Report
- **Expert Insights**
- Slideshows
- Videos
- **Podcasts**
- Resources
- Infographics
- Whitepapers
- Research
 - White Papers
 - Case Studies
 - Product Spec Sheets
 - Salary Survey
 - Market Data Datawatch Column
- **MPO Summit**
- **Events**
 - Industry Events
 - Live From Show Events
 - Webinars
- **Microsite**
 - Companies
 - Product Releases
 - Product Spec Sheets Services
 - White Papers / Tech Papers
 - Press Releases
 - Videos
 - Literature / Brochures Case Studies
- About Us About Us
 - Contact Us
 - Advertise with Us
 - eNewsletter Archive Privacy Policy
 - Terms of Use

Five Response Options to a Patent Infringement Demand Letter

Baldassare Vinti, Proskauer Rose04.06.16

In recent years, there has been an influx of patent litigation. Although not a requirement of litigation, these disputes often begin with a demand letter from the patent holder. These letters can take a variety of forms, but generally, they state or imply the recipient is infringing a patent, and they conclude by offering a license and/or demanding the recipient cease the activity (such as the sale of a medical device) that is allegedly infringing. Some letters may threaten a lawsuit if the activity does not cease.

Related CONTENT

- Watered-Down Version of Patent Reform Bill Could Have Far Reach

- Varian Settles Patent Case with University of Pittsburgh
 ResMed Wins Patent Infringement Lawsuit Against BMC Medical
 Arthrex Pays KFx Medical, Smith & Nephew Millions in Patent Laws
 CorMatrix Cardiovascular Receives Two U.S. Patents for Biomateria

While receiving such letters can be upsetting or confusing, the recipient has recourse to minimize or eliminate the risk and disruption to its business. Responding to demand letters involves strategic decisions based on a number of factors, including:

The patent owner's litigation history, the propensity for litigation, and sophistication

- The commercial relationship, if any, between the patent owner and the recipient of the demand letter.
- The established profitability of the product alleged to be infringing, its commercial success, and its current popularity.

6/1/2018 Five Response Options To A Patent Infringement Demand Letter - Your online source for medical device product information - Medical Pro...

The strength of the infringement allegations. Often, a patent attorney will be able to assess the strength of the patent infringement claims, at least to a certain extent, prior to the com a lawsuit.

The demand of the patent owner. Is the patent owner demanding the recipient cease the allegedly infringing activity or that the recipients take a license?

With these factors in mind, there are some common courses of action to consider upon receipt of a demand letter, each with different pros and cons.

Option 1: Request More Information

Pro: Allows more time to gather information, investigate the seriousness of the demand, and assess any potential exposure. Of course, if the demand letter provides sufficient information, may choose to simply deny infringement of any valid patent claim provided there is a good faith basis to do so.

Con: Requesting additional information, as opposed to a denial of infringement (assuming there is a basis to do so), may lead the patent owner to assume (perhaps incorrectly) that the recunsure whether it is infringing. Moreover, the patent owner is not legally obligated to respond, or respond adequately, to such a request.

Option 2: Do Nothing

Pro: Some patent owners who are not committed to actually filing a patent infringement lawsuit may nonetheless cast a wide net when sending demand letters in the hope that some recipi nominal amount to avoid a disruption to their business. If this is the case, there is a chance that by ignoring the demand letter, there will be no serious follow up by the patent owner.

Con: Ignoring a demand letter may cause the patent owner to file a patent infringement lawsuit.

Option 3: File a Lawsuit for Declaratory Judgment

In certain instances, a party accused of infringement may file a lawsuit against the patent owner for a declaration that it does not infringe the patent claims, or that the patent claims are invathe patent owner's litigation history is informative when considering this option.

Pro: If a party believes that the patent owner is likely to file a lawsuit, filing a declaratory judgment lawsuit may allow the recipient of the demand letter to file in its home court.

Con: May prove to be a more costly approach if the patent owner had no intention of actually commencing litigation.

Option 4: Seek to Invalidate the Patent

In certain circumstances, a party may seek to invalidate the asserted patent(s) by petitioning the United States Patent & Trademark Office to review the patent, such as by filing a petition fc Review, Covered Business Method, or Post Grant Review.

Pro: In certain circumstances, this option is a possible cost-effective alternative to litigation.

Con: If the patent survives the challenge, the patent owner may aggressively pursue its claims in litigation. Additionally, not all of these procedures are available for all patents. A patent attrable to assess the available and appropriate procedures.

Option 5: License the Patent

Negotiate with the patent owner for a license to use the patent

Pro: This option is a possible cost-effective resolution to litigation.

Con: In some circumstances, the license cost outweighs the cost of litigating the claim, especially if the claim can be disposed of early in litigation.

Conclusion

Damage awards in the medical device space can be quite large in certain instances. Therefore, it is important to take demand letters seriously. The correct approach to follow will depend o facts and circumstances of the case.

Company employees should be instructed to bring demand letters to the attention of the legal department as soon as they are received. A patent attorney will be able to assess the merits of and recommend a strategic approach to maximize the chances of a successful and cost-effective resolution.

Author Note: Fabio Tarud and Daniel Werb, associates at the firm, assisted in the preparation of this article.

Baldassare Vinti is a partner in the Patent Law and Intellectual Property Groups at Proskauer. He can be reached at bvinti@proskauer.com. Related Searches

- patent
- lifepak 1000 defibrillator
- trademark
- medical device

Suggested For You



Positive Results from Boston Scientific's Bronchial Thermoplasty Study

Portable Transcranial Doppler Tech Assesses Early Strokes with High Accuracy



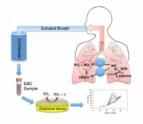
Masimo Introduces Rad-G Pulse Oximete



NeuroMetrix Reports Substantial Expansion of International IP Assets for Quell Technology



GI Dynamics Announces Scientific Advisory Board Members to Further Development of EndoBarrier



Graphene-Based Sensor Could Improve Asthma Treatment



Nevro Files U.S. Patent Infringement Lawsuit Against Boston Scientific



Glytec Receives Three Additional Patent Notices of Allowance for Diabetes Technologies



The Value of Medical Device Patents

CorMatrix Cardiovascular Receives Two U.S. Patents for Biomaterial Compositions



Arthrex Pays KFx Medical, Smith & Nephew Millions in Pater



ResMed Wins Patent Infringement Lawsuit Against BMC Med



Varian Settles Patent Case with University of Pittsburgh



Watered-Down Version of Patent Reform Bill Could Have Far Reaching Effects

Related Columns

Diagnostics
Theranos: Not Long for the Business World?
It won't be long now. With a skeleton staff, dwindling finances, a sullied reputation, and a disgraced leader under criminal investigation, embattled blood testing firm Theranos Inc. se destined for a terrible, miserable, no goo...
Michael Barbella, Managing Editor 05.03.18

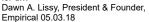
Pharma Industry's Growth—What Can Medtech Learn from It?
Global revenues for our sister industry—pharmaceuticals—continues to grow (Table 1). By 2022, worldwide pharmaceutical sales are projected to reach an estimated \$1.4 trillion at a CAGR of 6.4 percent from 2001 sales of \$390 billion.2 That. Maria Shepherd, President and Founder, Medi-Vantage 05.03.18





The Right Vendors Foster **Growth: Why Firms of All Sizes Need to Outsource**

I was working as a product development engineer at AcroMed (a spinal implants manufacturer that was acquired by DePuy in 1998), when I noticed a gap in the medical device industry. There were plenty of companies coming up with great ideas for d..





Embracing the Digital Evolution for Content Management

Digitalization, global expansion, and changing compliance requirements are driving medical device manufacturers to rethink their commercial models with a specific focus on content management. The industry's move toward digital has caused a prol. Terri Howard, Director of Commercial Strategy, Medical Device and Diagnostics, Veeva Systems 05.03.18



13485:2016—Beyond the Risk Management File

Many companies currently are transitioning from ISO 13485:2003 to ISO 13485:2016, the updated standard drafted to help medtech firms evolve with the industry and address changes in the underlying ISO 9001 benchmark. Some companies, however, are findi... James A. Dunning, Principal, QPC Services LLC 05.03.18



Medical Devices and the Political Climate of 2018: An Update from the Swamp

The November midterm elections have the potential to shift the power balance, leadership, and legislative agenda of Congress, which would certainly have a sizable impact on the medical device industry. In addition to what lies ahead, current actions.

Jeffrey J. Kimbell and David C. Rudloff, Jeffrey J. Kimbell & Associates Inc. 05.03.18



ISO 13485:2016—Upgrading to the New Edition (Part 2)

For the first time in 13 years, the International Organization for Standardization (ISO) has updated ISO 13485, the medical device industry's framework for quality management systems (QMS). With an emphasis on risk management in the quality sys...

Maria Fagan, President and Co-Founder, Regulatory and Quality Solutions LLC (R&Q) 05.03.18



Care Chain Consolidation: Tail Wagging the Dog?

For the past 35 years or so, supply chain consolidation has become an integral part of the medtech industry. Far less common but increasing in frequency is the desire by nonhealthcare entities like Amazon, Apple, CVS, Google, and most recently, Walm Chris Oleksy, Founder and CEO, Oleksy Enterprises and Next Life Medical; CEO, Emergent Respiratory 05.03.18



To Clinical or Not to Clinical, That Is the Question

For simple or well-characterized medical products, it is risk-appropriate and most cost effective to immediately design and engineer the commercial configuration of the device. In these cases, the design would presumably be sufficiently "locked... David C. Robson, Principal, Robson Advisors 05.03.18



Medtech M&A Off to a Strong Start in 2018

The medical device industry seems to be experiencing a bit of déjà vu this year. Though the start of 2018 was tinged with uncertainty over changes in the global healthcare market and the fate of Obamacare's medical device tax,... Mark Bonifacio, President, Bonifacio Consulting Services 05.03.18

Software & IT | Surgical **Extolling the Virtues of Surgical Simulation**

Gaining surgical competence for a new technique has a drastic learning curve. When surgeons are trained to perform a new procedure, however, the curriculum appears remarkably straightforward. Typically, the surgeon will attend a one- or two-day cours... Sam Brusco, Associate Editor 04.03.18



There are many opinions about the scope and scale of future medtech mergers after the M&A boom seen in 2016 and 2017. By June 2017, medical device M&A value rose 178 percent,1 propelled by the strategic imperative for medical device companies..

Maria Shepherd, President and Founder, Medi-Vantage 04.03.18







VED!

Learn from My Mistakes: Why You Need a Quality System

I've made my share of stupid mistakes. When I was younger, most of them stemmed from just not thinking things through. I still cringe to admit I had some really big hair in the 1980s, one of many reasons I'll be forever grateful my younge... Dawn A. Lissy, President & Founder, Empirical 04.03.18

Brexit: D-Day or the Y2K for Global Supply Chains?

On the same day Britons went to the polls to determine their country's future with the European Union (EU), the Irish Business & Employers Confederation (IBEC) held a manufacturing conference in Dublin. IBEC represents the policy interests. Dan O'Mahony, Lifesciences Manager, Exertis Supply Chain Services Limited 04.03.18

Planning for Brexit

Are you ISO 13485 certified by a United Kingdom Notified Body? Perhaps you're a contract manufacturer serving OEM customers who are certified by a U.K. Notified Body. Either way, medical device companies must start preparing now to mollify the ..

James A. Dunning, Principal, QPC Services LLC 04.03.18







Breaking News

- First-in-Human Treatment with J-Valve TAVI Device for Aortic Regurgitation
- Olympus Launches DualKnife J ESD Electrosurgical Knife to Shorten Proced FDA Clears Branchpoint Technologies' AURA Intracranial Pressure Monitoring
- System
- FDA Green Lights First Ever Artificial Iris
- Samsung Receives FDA Clearance for Premium Ultrasound System RS85

CURRENT ISSUE May 2018

- A Services Convergence: A Review of Medtech's Consolidating Supply Chain Developing the True Partnership: A Full-Service Outsourcing Roundtable
- So Fruitful Together: An M&A Roundtable
- **Designing from Finish to Start**
- View More >

View Breaking News >

Copyright © 2018 Rodman Media. All rights reserved. Use of this constitutes acceptance of our privacy policy The material on this site may not be reproduced, distributed, transmitted, or otherwise used, except with the prior written permission of Rodman Media.