

UNITED STATES
SECURITIES AND EXCHANGE COMMISSION
Washington, D.C. 20549

FORM 8-K

CURRENT REPORT

Pursuant to Section 13 OR 15(d) of The Securities Exchange Act of 1934

Date of Report (Date of earliest event reported) December 27, 2007

PSIVIDA LIMITED

(Exact name of registrant as specified in its charter)

Western Australia, Commonwealth of Australia
(State or other jurisdiction
of incorporation)

000-51122
(Commission File Number)

Not applicable
(IRS Employer
Identification No.)

**Level 12 BGC Centre
28 The Esplanade
Perth WA 6000
Australia**

**400 Pleasant Street
Watertown, MA 02472
U.S.A.**

(Address of principal executive offices) (Zip Code)

Registrant's telephone number, including area code: (617) 926-5000

Not applicable

(Former name or former address, if changed since last report.)

Check the appropriate box below if the Form 8-K filing is intended to simultaneously satisfy the filing obligation of the registrant under any of the following provisions:

- Written communications pursuant to Rule 425 under the Securities Act (17 CFR 230.425)
- Soliciting material pursuant to rule 14a-12 under the Exchange Act (17 CFR 240.14a-12)
- Pre-commencement communications pursuant to rule 14d-2(b) under the Exchange Act (17 CFR 240.14d-2(b))
- Pre-commencement communications pursuant to rule 13e04(c) under the Exchange Act (17 CFR 240.13e-4(c))

Item 3.01. Notice of Delisting or Failure to Satisfy a Continued Listing Rule or Standard; Transfer of Listing.

On December 27, 2007, the Company received a letter from the NASDAQ Listing Qualifications Department notifying the Company that, for the last 30 consecutive business days, the bid price of the Company's American Depositary Shares ("ADSs") has closed below the minimum \$1.00 per share required for continued listing on the NASDAQ Global Market under Marketplace Rule 4450(a)(5).

In accordance with Marketplace Rule 4450(e)(2), NASDAQ will provide the Company with 180 calendar days, or until June 24, 2008, to regain compliance. If at any time before June 24, 2008 the bid price of the Company's ADSs closes at \$1.00 per share or more for a minimum of 10 consecutive business days, NASDAQ will provide written notification that the Company has achieved compliance with Marketplace Rule 4450(a)(5). If compliance with Marketplace Rule 4450(a)(5) cannot be demonstrated by June 24, 2008, NASDAQ will provide written notice that the Company's securities will be delisted from the NASDAQ Global Market. At that time, the Company may appeal NASDAQ's determination or, if the Company satisfies the requirements of Marketplace Rule 4310(c) other than the minimum bid requirement, the Company may apply to transfer its securities to the NASDAQ Capital Market, in which case, if the Company's application is approved, the Company will be afforded the remainder of the Capital Market's second 180 calendar day period to regain compliance while on the NASDAQ Capital Market.

During the provided compliance periods, the Company will seek to regain compliance. The Company will also continue to monitor its stock price closely and will consider its options for regaining compliance in the event that its stock price remains below \$1.00. No assurances can be made at this point as to whether the Company will regain compliance.

On December 31, 2007, the Company issued a press release announcing the NASDAQ letter. A copy of this press release is filed as Exhibit 99.1 to this report and is incorporated herein by reference.

Item 9.01. Financial Statements and Exhibits

99.1 Press Release dated December 31, 2007

Incorporation by Reference

pSivida Limited hereby incorporates by reference this Current Report on Form 8-K in the Company's registration statements (Nos. 333-132776, 333-132777, 333-135428, 333-141083, 333-141091 and 333-143225) on Form F-3.

SIGNATURE

Pursuant to the requirements of the Securities Exchange Act of 1934, the registrant has duly caused this report to be signed on its behalf by the undersigned hereunto duly authorized.

PSIVIDA LIMITED

By: /s/ Michael J. Soja

Name: Michael J. Soja

Title: Vice President, Finance and
Chief Financial Officer

Dated: January 2, 2008

EXHIBIT INDEX

**Exhibit
Number**

Exhibit Description

99.1

Press Release dated December 31, 2007



Media RELEASE

31 December 2007

pSivida Receives NASDAQ Notification Related to Minimum Bid Price

Boston, MA and Perth, Australia (December 31, 2007) – pSivida Limited (NASDAQ: PSDV, ASX: PSD, FF: PSI), a global drug delivery company, today announced that on December 27, 2007, the Company received a letter from the NASDAQ Listing Qualifications Department notifying the Company that, for the last 30 consecutive business days, the bid price of the Company’s American Depositary Shares (“ADSs”) has closed below the minimum \$1.00 per share required for continued listing on the NASDAQ Global Market under Marketplace Rule 4450(a)(5). Under the rules set forth by the NASDAQ Listing Qualifications Department, issuing this notice is customary practice when a NASDAQ quoted company’s closing bid price has been less than \$1.00 per share for 30 consecutive trading days.

The NASDAQ letter does not affect the listing of the Company at this time, and the Company’s shares will continue to trade on the NASDAQ Global Market under the symbol “PSDV.”

In accordance with Marketplace Rule 4450(e)(2), NASDAQ will provide the Company with 180 calendar days, or until June 24, 2008, to regain compliance. If at any time before June 24, 2008 the bid price of the Company’s ADSs closes at \$1.00 per share or more for a minimum of 10 consecutive business days, NASDAQ will provide written notification that the Company has achieved compliance with Marketplace Rule 4450(a)(5). If compliance with Marketplace Rule 4450(a)(5) cannot be demonstrated by June 24, 2008, NASDAQ will provide written notice that the Company’s Securities will be delisted from the NASDAQ Global Market. At that time, the Company may appeal NASDAQ’s determination or, if the Company satisfies the requirements of Marketplace Rule 4310(c) other than the minimum bid requirement, the Company may apply to transfer its securities to the NASDAQ Capital Market, in which case, if the Company’s application is approved, the Company will be afforded the remainder of the Capital Market’s second 180 calendar day period to regain compliance while on the NASDAQ Capital Market.

“The Company is committed to regaining compliance, and are pursuing various strategies including the potential of non-dilutive capital,” said pSivida’s Managing Director, Dr Paul Ashton. “The Company believes its current cash position, together with the expected development funding from Pfizer and the remaining US\$1.5 million due in April related to the sale of a subsidiary, is sufficient to continue operations until at least September 30th, 2008”.

-ENDS-

Released by:

pSivida Limited

Brian Leedman
Vice President, Investor Relations
pSivida Limited
Tel: +61 8 9226 5099
brianl@psivida.com

US Public Relations

Beverly Jedynek
President
Martin E. Janis & Company, Inc
Tel: +1 (312) 943 1100 ext. 12
bjedynek@janispr.com

European Public Relations

Eva Reuter
Accent Marketing Limited
Tel: +49 (254) 393 0740
e.reuter@dr-reuter.eu

NOTES TO EDITORS:

pSivida is a global drug delivery company committed to the biomedical sector and the development of drug delivery products. Retisert® is FDA approved for the treatment of uveitis. Vitrasert® is FDA approved for the treatment of AIDS-related CMV Retinitis. Bausch & Lomb owns the trademarks Vitrasert® and Retisert®. pSivida has licensed the technologies underlying both of these products to Bausch & Lomb. The technology underlying Medidur™ for diabetic macular edema is licensed to Alimera Sciences and is in Phase III clinical trials. pSivida has a worldwide collaborative research and license agreement with Pfizer Inc. for other ophthalmic applications of the Medidur™ technology.

pSivida owns the rights to develop and commercialize a modified form of silicon (porosified or nano-structured silicon) known as BioSilicon™, which has applications in drug delivery, wound healing, orthopedics, and tissue engineering. The most advanced BioSilicon™ product, BrachySil™ delivers a therapeutic, P32 directly to solid tumors and is presently in Phase II clinical trials for the treatment of pancreatic cancer.

pSivida's intellectual property portfolio consists of 71 patent families, 99 granted patents, including patents accepted for issuance, and over 300 patent applications. pSivida conducts its operations from facilities near Boston in the United States, Malvern in the United Kingdom and Perth in Australia.

pSivida is listed on NASDAQ (**PSDV**), the Australian Stock Exchange (**PSD**) and on the Frankfurt Stock Exchange on the XETRA system (**PSI**). pSivida is a founding member of the NASDAQ Health Care Index and the Merrill Lynch Nanotechnology Index.

Various statements made in this release are forward-looking and involve a number of risks and uncertainties. All statements that address activities, events or developments that we intend, expect or believe may occur in the future are forward-looking statements. The following are some of the factors that could cause actual results to differ materially from the forward-looking statements: the risks that we will not be able to raise additional capital; that we will continue to incur losses and may never become profitable; that we will be required to pay penalties pursuant to registration agreements with securities holders and not have sufficient funds to do so; that we will be unable to develop new products; that we will be unable to protect our own intellectual property or will infringe on others' intellectual property; that we will not receive regulatory approvals necessary to commercialize products; that we will be unable to secure partners necessary to develop and market products; that our current licensees will terminate their agreements with us; that our competitors' products will receive regulatory approval before, reach the market before, or otherwise receive better market acceptance than, our product candidates; that our international business operations will result in increased costs or delays; that manufacturing problems will delay product development and commercialization; that third-party reimbursement and health care providers will not cover the costs of our products; that we will fail to retain some or all of our key personnel; we will be subject to product liability suits and not have sufficient insurance to cover damages; that we will fail to effectively manage changes in our business; that we will fail to comply with environmental laws and regulations; that we will fail to achieve and maintain effective internal control over financial reporting; that amortization or impairment of other intangibles will adversely affect our operating results; that our being headquartered outside of the United States will make it difficult to effect legal services against us or our management, lead to adverse shareholder tax consequences, or otherwise limit shareholder rights; that we will be delisted from the ASX or NASDAQ; that our expectation to not pay cash dividends will decrease our stock price; that exercise of outstanding warrants and stock options will dilute ownership and reduce stock price; that future stock issuances could dilute ownership, restrict operations, encumber assets, or otherwise cause a decline in stock price; and the risk that Pfizer will influence our business in non-beneficial ways; and other factors that may be described in our filings with the Securities and Exchange Commission. Given these uncertainties, readers are cautioned not to place undue reliance on such forward-looking statements. We do not undertake to publicly update or revise our forward-looking statements even if experience or future changes make it clear that any projected results expressed or implied in such statements will not be realized.