SECURITIES AND EXCHANGE COMMISSION
Washington, D.C. 20549

FORM 6-K

REPORT OF FOREIGN ISSUER
Pursuant to Rule 13a-16 or 15d-16 of
the Securities Exchange Act of 1934

For the month of May 2006

Commission File Number 000-51122

pSivida Limited

(Translation of registrant's name into English)

Level 12 BGC Centre
28 The Esplanade
Perth WA 6000
(Address of principal executive offices)

(Indicate by check mark whether the registrant files or will file annual reports under cover Form 20-F or Form 40-F).

Form 20-F ☒ Form 40-F o

Indicate by check mark if the registrant is submitting the Form 6-K in paper as permitted by Regulation S-T Rule 101(b)(1):

Indicate by check mark if the registrant is submitting the Form 6-K in paper as permitted by Regulation S-T Rule 101(b)(7):

Indicate by check mark whether the registrant by furnishing the information contained in this Form is also thereby furnishing the information to the Commission pursuant to Rule 12g3-2(b) under the Securities Exchange Act of 1934.

Yes o No ☒

If "Yes" is marked, indicate below the file number assigned to the registrant in connection with Rule 12g3-2(b): 82- ___.

SIGNATURE

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

Date: May 30, 2006

pSivida Limited

By: /s/ Aaron Finlay

Aaron Finlay
Company Secretary

EXHIBIT INDEX

EXHIBIT 99.1:	BrachySil™ Pancreatic Program: Regulatory Approval for European Human Trial
EXHIBIT 99.1:	BrachySil™ Liver Programme: Expansion of Multicentre Clinical Trial
EXHIBIT 99.1:	Despatch for Rights Issue Prospectus

ASX/MEDIA RELEASE **30 May 2006**

BrachySil™ Pancreatic Program: Regulatory Approval for European Human Trial

---

Boston, MA. and Perth, Australia – Global bio-nanotech company pSivida Limited **(ASX:PSD, NASDAQ:PSDV, Xetra:PSI)** is pleased to announce that the Regulatory Agency in the U.K. (The Medicines and Healthcare Products Regulatory Agency or MHRA) has granted approval to proceed with the first human study of BrachySil™ in pancreatic cancer through a phase IIa clinical trial.

Pancreatic cancer is a second clinical indication for BrachySil™, currently in Phase IIb clinical trials for the treatment of inoperable primary liver cancer. Pancreatic cancer has one of the lowest cancer survival rates (5 year overall survival rate of approximately 5%). There is significant clinical and market demand for effective therapies to treat this aggressive form of cancer. According to *GLOBOCAN, there were over 230,000 new cases and nearly as many deaths from pancreatic cancer worldwide in 2002. Approximately 50% of these new cases were in North America and Europe.

The six month clinical study which has been approved to proceed by the MHRA is a phase IIa trial in patients with inoperable pancreatic cancer and will be undertaken at two leading centres for cancer treatment, Guys & St Thomas' Hospital in London (UK) and Singapore General Hospital. The trial represents the "first-in-Man" safety study for BrachySil™ in this indication and is designed to enrol a total of 15 patients. The primary objective is to determine the safety of the targeted image-guided implantation of pSivida's BrachySil™ product. Efficacy, as determined by CT scanning of the tumour size and overall survival, will be secondary endpoints. The findings will provide a platform for further multicentre efficacy and safety trials.

Pre-clinical evaluation of BrachySil™ into pancreatic cancers has provided the teams involved in the clinical trial with valuable feedback allowing optimisation of BrachySil™ for this indication and standardisation between the two study centres. The clinical program will utilise product sourced from the same GMP manufacturing process and supply chain which is currently being utilised for the liver cancer program. The manufacturing process is established and scaled up to support clinical development and early launch volumes.

A phase IIa study for advanced inoperable liver cancer completed in June 2005 on eight patients showed BrachySil™ to be both safe and well tolerated. All patients experienced a decrease in the size of their tumours, with some experiencing complete tumour regression.

"We have recently announced progress with the development of BrachySil™, our targeted oncology product, in primary liver cancer. We believe that because the BrachySil™ primary liver program is in Phase IIb clinical trials, it will provide valuable clinical information for the development and commercialisation of BrachySil™ for pancreatic cancer and other indications," said pSivida CEO, Mr Gavin Rezos. "We also believe that gaining regulatory agency approval in Europe to begin the first clinical study for the pancreatic cancer indication represents a very positive milestone towards broader value generation from BrachySil™."

*GLOBOCAN is a worldwide database of cancer incidence and mortality rates.

-ENDS-

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

US Public Relations
Beverly Jedynak
President
Martin E. Janis & Company, Inc
Tel: +1 (312) 943 1100 ext. 12
bjedynak@janispr.com

UK & Europe Public Relations
Mark Swallow / Helena Podd
Citigate Dewe Rogerson
Tel: +44 (0)20 7638 9571
mark.swallow@citigatedr.co.uk

NOTES TO EDITORS:

pSivida is a global bio-nanotech 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 own the trademarks Vitrasert® and Retisert™. pSivida has licensed the technologies underlying both of these products to Bausch & Lomb. The technology underlying Medidur™, a treatment for diabetic macular edema, is licensed to Alimera Sciences and is in Phase III clinical trials.

pSivida owns the rights to develop and commercialise a modified form of silicon (porosified or nano-structured silicon) known as BioSilicon™, which has applications in drug delivery, wound healing, orthopaedics, and tissue engineering. pSivida's subsidiary, AION Diagnostics Limited is developing diagnostic products and the subsidiary pSiNutria is developing food technology products both using BioSilicon™.

pSivida's intellectual property portfolio consists of 70 patent families, 74 granted patents and over 290 patent applications. pSivida conducts its operations from offices and facilities near Boston in the United States, Malvern in the United Kingdom, Perth in Australia and Singapore.

pSivida is listed on NASDAQ (**PSDV**), the Australian Stock Exchange (**PSD**) and on the Frankfurt Stock Exchange on the XETRA system (**German Symbol: PSI. Securities Code (WKN) 358705**). pSivida is a founding member of the NASDAQ Health Care Index and the Merrill Lynch Nanotechnology Index.

The Company's largest shareholder and a strategic partner is QinetiQ, a leading international defence, security and technology company, formed in 2001 from the UK Government's Defence Evaluation & Research Agency (DERA). QinetiQ was instrumental in discovering BioSilicon(TM) and pSivida enjoys a strong relationship with, including access to its cutting edge research and development facilities.

For more information, visit www.psivida.com

This document contains forward-looking statements that involve risks and uncertainties. The statements are indicated by the use of words such as "believes", "expects", "anticipates" and similar words and phrases. Although we believe that the expectations reflected in such forward-looking statements are reasonable at this time, we can give no assurance that such expectations will prove to be correct. Given these uncertainties, readers are cautioned not to place undue reliance on such forward-looking statements. Actual results could differ materially from those anticipated in these forward-looking statements due to many important factors including: failure to complete negotiations for new centers for the BrachySil™ phase IIb clinical trial for inoperable primary liver cancer; the failure of our discussions with the FDA for BrachySil™ to continue or to lead to FDA approval; the failure of the BrachySil™ phase IIb clinical trial for inoperable primary liver cancer to determine the optimal dose, provide key safety data or support future pivotal efficacy trials or product registration or approval; failure to commence Phase IIa BrachySil™ trials for the treatment of pancreatic cancer; the failure of the results of the Retisert™ for DME trial to be a good indicator of the results of pSivida's ongoing Phase III Medidur™ for DME trial; failure of the Medidur™ trials in DME to show a very similar improvement in visual acuity and diabetic retinopathy severity score as Retisert™ for DME; inability to recruit patients for the Phase III Medidur™ for DME trial; our failure to develop applications for BioSilicon™ due to regulatory, scientific or other issues, our inability to successfully integrate pSivida Inc's operations and employees; the failure of the pSivida Inc's products to achieve expected revenues and the combined entity's inability to develop existing or proposed products; the failure of the Bausch & Lomb/Novartis co-promotion arrangement to provide faster royalty growth; failure of the slower progression or reduction of diabetic retinopathy resulting from the Retisert™ implant to have significant implications for Retisert™ and Medidur™; failure of our evaluation agreements to result in license agreements; failure of Medidur™ to release the same drug as Retisert™ at the same rate; failure of the Medidur™ trials in DME to show a very similar stabilization or improvement diabetic retinopathy as Retisert™ for DME; failure to achieve cost savings; failure to execute on US growth strategy; failure of the findings of the pancreatic cancer phase IIa trial to provide a platform for further multicentre efficacy and safety trials; failure of there to be optimisation and standardisation between the two pancreatic cancer study centres; failure of the BrachySil™ primary liver program that is in Phase IIb clinical trials to provide a valuable platform for the development and commercialisation of BrachySil™ for pancreatic cancer and other indications. Other reasons are contained in cautionary statements in the Annual Report on Form 20-F filed with the U.S. Securities and Exchange Commission, including, without limitation, under Item 3.D, "Risk Factors" therein. We do not undertake to update any oral or written forward-looking statements that may be made by or on behalf of pSivida.

ASX/MEDIA RELEASE **25 May 2006**

BrachySil™ Liver Programme: Expansion of Multicentre Clinical Trial

---

Boston, MA. and Perth, Australia – Global bio-nanotech company pSivida Limited **(ASX:PSD, NASDAQ:PSDV, Xetra:PSI)** is pleased to announce that the scope of the BrachySil™ phase IIb clinical trial for inoperable primary liver cancer has expanded to encompass a roll-out to further clinical centres in new Asian territories. In addition to the five centres currently active in Singapore, and centres in Vietnam and Malaysia, new centres are being negotiated in the Philippines and Taiwan.

A phase IIb clinical trial commenced in October 2005 with BrachySil™ (32-P BioSilicon™) as a potential new treatment for inoperable primary liver cancer (hepatocellular carcinoma, HCC). pSivida is also progressing a formal dialogue with the FDA relating to the U.S. programme for HCC and pancreatic cancers. This active discourse has provided valuable information for pSivida to advance its programme planning in the U.S. market.

The study, which was designed in collaboration with Singapore General Hospital (SGH) and approved by the Singaporean regulatory authority (Health Sciences Authority), is designed to determine the optimal dose of BrachySil™ in treating inoperable HCC. Patients will be evaluated up to 12 months after treatment. The endpoints will be based on evaluations of patient safety and target tumour responses, as well as overall survival.

A phase IIa study conducted at SGH that commenced in June 2004 on eight patients with advanced liver cancer showed BrachySil™ to be both safe and well tolerated. All patients experienced a decrease in the size of their tumours, with some experiencing complete regression. To date, six of the eight patients are still alive.

Phase IIa BrachySil™ trials for the treatment of pancreatic cancer to be conducted in hospitals in London and Singapore are expected to commence next month. Pancreatic cancer typically has one of the lowest cancer survival rates and represents a further important clinical indication for BrachySil™ with a high unmet need.

“We are delighted to announce that the liver cancer programme will now be supported by a larger number of clinical centres in the key Asian region, and that clinical evaluation of the BrachySil™ product in pancreatic cancer remains on track,” said pSivida CEO, Mr. Gavin Rezos. “Our target is to generate value for our lead oncology product in all key markets, particularly the US for which the active dialogue with the FDA is a key component.”

-ENDS-

---

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

US Public Relations
Beverly Jedynak
President
Martin E. Janis & Company, Inc
Tel: +1 (312) 943 1100 ext. 12
bjedynak@janispr.com

UK & Europe Public Relations
Mark Swallow / Helena Podd
Citigate Dewe Rogerson
Tel: +44 (0)20 7638 9571
mark.swallow@citigatedr.co.uk

NOTES TO EDITORS:

pSivida is a global bio-nanotech 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 own the trademarks Vitrasert® and Retisert™. pSivida has licensed the technologies underlying both of these products to Bausch & Lomb. The technology underlying Medidur™, a treatment for diabetic macular edema, is licensed to Alimera Sciences and is in Phase III clinical trials.

pSivida owns the rights to develop and commercialise a modified form of silicon (porosified or nano-structured silicon) known as BioSilicon™, which has applications in drug delivery, wound healing, orthopaedics, and tissue engineering. pSivida's subsidiary, AION Diagnostics Limited is developing diagnostic products and the subsidiary pSiNutria is developing food technology products both using BioSilicon™.

pSivida's intellectual property portfolio consists of 70 patent families, 74 granted patents and over 290 patent applications. pSivida conducts its operations from offices and facilities near Boston in the United States, Malvern in the United Kingdom, Perth in Australia and Singapore.

pSivida is listed on NASDAQ (**PSDV**), the Australian Stock Exchange (**PSD**) and on the Frankfurt Stock Exchange on the XETRA system (**German Symbol: PSI. Securities Code (WKN) 358705**). pSivida is a founding member of the NASDAQ Health Care Index and the Merrill Lynch Nanotechnology Index.

The Company's largest shareholder and a strategic partner is QinetiQ, a leading international defence, security and technology company, formed in 2001 from the UK Government's Defence Evaluation & Research Agency (DERA). QinetiQ was instrumental in discovering BioSilicon(TM) and pSivida enjoys a strong relationship with it having access to its cutting edge research and development facilities. For more information visit www.QinetiQ.com

For more information, visit www.psivida.com

This document contains forward-looking statements that involve risks and uncertainties. The statements are indicated by the use of words such as "believes", "expects", "anticipates" and similar words and phrases. Although we believe that the expectations reflected in such forward-looking statements are reasonable at this time, we can give no assurance that such expectations will prove to be correct. Given these uncertainties, readers are cautioned not to place undue reliance on such forward-looking statements. Actual results could differ materially from those anticipated in these forward-looking statements due to many important factors including: failure to complete negotiations for new centers for the BrachySil™ phase IIb clinical trial for inoperable primary liver cancer; the failure of our discussions with the FDA for BrachySil™ to continue or to lead to FDA approval; the failure of the BrachySil™ phase IIb clinical trial for inoperable primary liver cancer to determine the optimal dose, provide key safety data or support future pivotal efficacy trials or product registration or approval; failure to commence Phase IIa BrachySil™ trials for the treatment of pancreatic cancer; the failure of the results of the Retisert™ for DME trial to be a good indicator of the results of pSivida's ongoing Phase III Medidur™ for DME trial; failure of the Medidur™ trials in DME to show a very similar improvement in visual acuity and diabetic retinopathy severity score as Retisert™ for DME; inability to recruit patients for the Phase III Medidur™ for DME trial; our failure to develop applications for BioSilicon™ due to regulatory, scientific or other issues, our inability to successfully integrate pSivida Inc's operations and employees; the failure of the pSivida Inc's products to achieve expected revenues and the combined entity's inability to develop existing or proposed products; the failure of the Bausch & Lomb/Novartis co-promotion arrangement to provide faster royalty growth; failure of the slower progression or reduction of diabetic retinopathy resulting from the Retisert™ implant to have significant implications for Retisert™ and Medidur™; failure of our evaluation agreements to result in license agreements; failure of Medidur™ to release the same drug as Retisert™ at the same rate; failure of the Medidur™ trials in DME to show a very similar stabilization or improvement diabetic retinopathy as Retisert™ for DME; failure to achieve cost savings; failure to execute on US growth strategy. Other reasons are contained in cautionary statements in the Annual Report on Form 20-F filed with the U.S. Securities and Exchange Commission, including, without limitation, under Item 3.D, "Risk Factors" therein. We do not undertake to update any oral or written forward-looking statements that may be made by or on behalf of pSivida.

ASX/MEDIA RELEASE **24 May 2006**

Despatch of Rights Issue Prospectus

Boston, MA. and Perth, Australia – Global bio-nanotech company pSivida Limited **(ASX:PSD, NASDAQ:PSDV, Xetra:PSI)** is pleased to confirm that the despatch to eligible shareholders of the Rights Issue prospectus and entitlement and acceptance forms was completed today.

As announced on 2 May 2006, the Rights Issue is on a non-renounceable 1 for 8 basis to raise approximately A$29 million at A$0.60 per ordinary share. The Rights Issue has an incorporated top up facility whereby eligible shareholders may apply for additional new ordinary shares in excess of their entitlement at the same price.

The Rights Issue is not underwritten. To the extent there is any shortfall under the Rights Issue, pSivida has agreed to place such shortfall through Janney Montgomery Scott LLC, its US based Lead Manager for this issue, to institutional and sophisticated investors.

Any ordinary shares issued in the U.S. will be issued in a private placement. These shares will not be registered under the U.S. Securities Act of 1933, as amended, or any U.S. state securities laws and may not be offered, sold or transferred in the U.S. absent registration or an applicable exemption from registration requirements.

The Record Date for the Rights Issue was 22 May 2006. The ordinary shares to be issued in connection with the Rights Issue are expected to commence trading on the Australian Stock Exchange on 8 June 2006. Applications will close on 7 June 2006.

-ENDS-

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

US Public Relations
Beverly Jedynak
President
Martin E. Janis & Company, Inc
Tel: +1 (312) 943 1100 ext. 12
bjedynak@janispr.com

UK & Europe Public Relations
Mark Swallow / Helena Podd
Citigate Dewe Rogerson
Tel: +44 (0)20 7638 9571
mark.swallow@citigatedr.co.uk

NOTES TO EDITORS:

pSivida is a global bio-nanotech 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 own the trademarks Vitrasert® and Retisert™. pSivida has licensed the technologies underlying both of these products to Bausch & Lomb. The technology underlying Medidur™, a treatment for diabetic macular edema, is licensed to Alimera Sciences and is in Phase III clinical trials.

pSivida owns the rights to develop and commercialise a modified form of silicon (porosified or nano-structured silicon) known as BioSilicon™, which has applications in drug delivery, wound healing, orthopaedics, and tissue engineering. pSivida's subsidiary, AION Diagnostics Limited is developing diagnostic products and the subsidiary pSiNutria is developing food technology products both using BioSilicon™.

pSivida's intellectual property portfolio consists of 70 patent families, 74 granted patents and over 290 patent applications.

pSivida conducts its operations from offices and facilities near Boston in the United States, Malvern in the United Kingdom, Perth in Western Australia and Singapore.

pSivida is listed on NASDAQ (**PSDV**), the Australian Stock Exchange (**PSD**) and on the Frankfurt Stock Exchange on the XETRA system (**German Symbol: PSI. Securities Code (WKN) 358705**). pSivida is a founding member of the NASDAQ Health Care Index and the Merrill Lynch Nanotechnology Index.

The Company's largest shareholder and a strategic partner is QinetiQ, a leading international defence, security and technology company, formed in 2001 from the UK Government's Defence Evaluation & Research Agency (DERA). QinetiQ was instrumental in discovering BioSilicon$^{(TM)}$ and pSivida enjoys a strong relationship with it having access to its cutting edge research and development facilities. For more information visit www.QinetiQ.com

For more information, visit www.psivida.com

This document contains forward-looking statements that involve risks and uncertainties. The statements are indicated by the use of words such as "believes", "expects", "anticipates" and similar words and phrases. Although we believe that the expectations reflected in such forward-looking statements are reasonable at this time, we can give no assurance that such expectations will prove to be correct. Given these uncertainties, readers are cautioned not to place undue reliance on such forward-looking statements. Actual results could differ materially from those anticipated in these forward-looking statements due to many important factors including: the failure of the results of the Retisert for DME trial to be a good indicator of the results of pSivida's ongoing Phase III Medidur™ for DME trial; failure of the Medidur™ trials in DME to show a very similar improvement in visual acuity and diabetic retinopathy severity score as Retisert™ for DME; inability to recruit patients for the Phase III Medidur™ for DME trial; our failure to develop applications for BioSilicon™ due to regulatory, scientific or other issues, our inability to successfully integrate pSivida Inc's operations and employees; the failure of the pSivida Inc's products to achieve expected revenues and the combined entity's inability to develop existing or proposed products; the failure of the Bausch & Lomb/Novartis co-promotion arrangement to provide faster royalty growth; failure of the slower progression or reduction of diabetic retinopathy resulting from the Retisert™ implant to have significant implications for Retisert™ and Medidur; failure of our evaluation agreements to result in license agreements; failure of Medidur™ to release the same drug as Retisert™ at the same rate; failure of the Medidur™ trials in DME to show a very similar stabilization or improvement diabetic retinopathy as Retisert™ for DME; failure to achieve cost savings; failure to execute on US growth strategy. Other reasons are contained in cautionary statements in the Annual Report on Form 20-F filed with the U.S. Securities and Exchange Commission, including, without limitation, under Item 3.D, "Risk Factors" therein. We do not undertake to update any oral or written forward-looking statements that may be made by or on behalf of pSivida.