SOLARA
Active Pharma Sciences

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July 8, 2019

The BSE Limited
Phiroze Jeejeebhoy Towers
Dalal Street, Mumbai – 400 001

Scrip Code: 541540

The National Stock Exchange of India Limited
Exchange Plaza, Bandra-Kurla Complex
Bandra (E), Mumbai – 400 051

Scrip Code: SOLARA

Dear Sir / Madam,

Sub: Announcement under Regulation 30 of SEBI (Listing Obligations and Disclosure Requirements) Regulation, 2015

We are enclosing herewith a press release issued by the Company titled:

"Solara Cuddalore facility completes USFDA Inspection with zero 483s"

Thanking you,

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Yours faithfully,
For Solara Active Pharma Sciences Limited

S. Murali Krishna
Company Secretary

Encl:- as above

SOLARA ACTIVE PHARMA SCIENCES LIMITED

Solara's Cuddalore facility completes USFDA Inspection with ZERO 483s

Chennai, India – July 8, 2019: Solara Active Pharma Science Limited (Solara), a leading Active Pharmaceutical Ingredient provider today announced the successful completion of US FDA inspection at the multi-product manufacturing facility at Cuddalore with ZERO 483s inspectional observations.

The Cuddalore multi-product API manufacturing facility was established in the year 1991 and is equipped with appropriate infrastructure to include several independent production blocks and related packaging sections. This site is inspected by various Regulatory Authorities including US FDA, MHRA, EDQM, WHO, PMDA, TGA, KFDA, and COFEPRIS. The current US FDA inspection at the facility was scheduled and completed between 1st July and 5th July 2019.

Solara continues to stay focused on maintaining the highest level of compliance across its manufacturing facilities. This is Company's third consecutive US FDA audit with "Zero 483s". In Dec'18 and Jan'19, the company also completed US FDA inspections at Solara Research Centre(SRC), Chennai and Ambernath manufacturing facility with zero 483s.

About Solara

Solara Active Pharma Sciences Ltd (BSE-541540, NSE-SOLARA) headquartered in Bengaluru, India offers a basket of diversified, high-value Commercial APIs and Contract manufacturing services in over 75 countries. It has a manufacturing base comprising five API facilities in India with approvals including the USFDA, EU GMP, KFDA and PMDA in Japan.

Investor / Analyst contact

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Disclaimer: Certain statements in this document that are not historical facts are forward-looking statements. Such forward-looking statements are subject to certain risks and uncertainties like government actions, local, political or economic developments, technological risks, and many other factors that could cause actual results to differ materially from those contemplated by the relevant forward-looking statements. Solara Active Pharma Sciences Ltd will not be in any way responsible for any action taken based on such statements and undertakes no obligation to publicly update these forward-looking statements to reflect subsequent events or circumstances.

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