

CPHI & P-MEC India comes with the backing of the Delhi-based Pharmaceuticals Export Promotion Council of India (Pharmexcil) – the organisation behind key Government international strategies such as 'Made in India'. Mr. Yogesh Mudras, Managing Director, UBM India commented "the shift to the Delhi-NCR region, in close geographical proximity to the policy makers, consulates and government bodies will enhance our community building efforts, as well as the creation of a robust pharma eco-system."

Government investment plans – including a US\$ 640 million venture capital fund to boost drug discovery and strengthen pharmaceutical infrastructure – and the 'Pharma Vision 2020' initiative are creating new benefits for government engagement with industry.

UBM cited rising international interest in the event as a key factor in the move, with numbers of both international and domestic exhibitors requiring a far bigger venue – and on a single site.

The venue change is also seen as a boost to domestically focused markets by bringing a wider remit of attendees and greater access to national regulatory pathways. The country registers the second largest Abbreviated New Drug Applications (ANDAs) globally and is the world's leader in Drug Master Files (DMFs) applications, so access to regulatory expertise has never been greater.

The 2018 edition will also feature India Pharma Week (returning for a third year), with 10 unique events and activities spanning the Indian Capital. These include dynamic engagements such as the Pharma Leader's Golf Pre-Connect Congress, Plant Visits, Women in Pharma – Power Breakfast, India Pharma Awards, Networking Evening, and a closed-door CEO Roundtable, amongst others.

The ultimate goal of all these intertwined events is to help nurture a complete pharma ecosystem in India and better enable networking with the country's corridors of power. In fact, the recent CPHI Global Pharma Index showed India represents global pharma's second fastest growing market and praised the Delhi based regulator, the CDSCO (Central Drugs Standard Control Organization) for its efforts in introducing a certification programme and initiatives for increased compliance. Impressively, 52% of international respondents believed the CDSCO is moving toward comparability with the regulatory standards of the EMA and FDA.