ANNUAL REPORT FOR IPR 2020-21

S.NO.	NAME OF THE SEMINAR/CONFERENCE/ WORKSHOP	NUMBER OF PARTICIPANTS	DATE	RESOURCE PERSON
1.	Assurance of Quality and Regulatory Issues	25	17/8/2020	Dr.K.V.Subramanyam
2.	ANDAs: The Development and Review of Pharmaceuticals	30	21/02/2021	Dr.Sandhya Rani
3.	CGMP_CFR section on 211 and 210.	22	11/2/2020	Dr.Prahlad
4.	Directives for the European Union and Australia	32	24/09/2021	Dr.Raju Srivastav
5.	Drug and cosmetic regulations in the Japanese market	25	21/4/2021	Dr.Anmol Kher
6.	Emerging market drug registration and approval procedures	32	27/3/2021	Dr.Shirin Sultana
7.	Methods for Filing a Patent	31	5/11/2020	Dr.Soujanya Reddy
8.	Rights to intellectual property	45	11/12/2020	Dr.Srihari

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CIRCULAR

Date: 14/8/2020

An IPR activity titled " **Assurance of Quality and Regulatory Issues**" is scheduled to take place on our campus on 17/8/2020. All of the staff members have been notified that this event will take place. Everyone is admonished to attend the Program on each and every occasion in order to acquire information.

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EVENT REPORT

Name Of The Activity	Assurance of Quality and Regulatory Issues	
Type Of Activity	Seminar	
Date And Time Of Activity	17/8/2020	10.30AM
No. Of Participants	25	
Resource Person	Dr.K.V.Subramanyam	
Coordinators	Mr. M. Naveen Kumar	
Description	The principal addressed the gathering and welcomed the guest. The HOD introduced about the guest and handled over to the speaker. He says that It's not uncommon for regulatory affairs and quality assurance to be discussed in the same breath, as though the two industries are the same. But, although the two fields intersect, they are indeed different. The line between regulatory affairs and quality assurance can sometimes be difficult to distinguish, but there are fundamental differences that set the two disciplines apart. Regulatory affairs is an industry tasked with overseeing how certain products are developed, tested, manufactured, marketed, and distributed to ensure each process is compliant with the relevant regulatory statutes implemented by various regulatory agencies. These professionals often work in the biopharmaceutical, medical devices, and food safety industries. Finally HOD delivered the vote of thanks and the guest was felicitated with a momentum	
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Date: 16/02/2021

It is therefore announced to all staff that an IPR activity on **ANDAs: The Development and Review of Pharmaceuticals** Process will be held at our campus on 21/02/2021. Everyone is urged to attend the Program without fail in order to acquire the knowledge.

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EVENT REPORT

Name Of The Activity	ANDAs: The Development and Review of Pharmaceuticals	
Type Of Activity	Seminar	
Date And Time Of Activity	21/02/2021	11.30AM
No. Of Participants	30	
Resource Person	Dr.Sandhya Rani	
Coordinators	Mr. M. Naveen Kumar	
Description	The principal addressed the gathering and welcomed the guest. The HOD introduced about the guest and handled over to the speaker. The speaker started with the an abbreviated new drug application (ANDA) contains data which is submitted to FDA for the review and potential approval of a generic drug product. Once approved, an applicant may manufacture and market the generic drug product to provide a safe, effective, lower cost alternative to the brand-name drug it references. Finally the HOD delivered the vote of thanks and the guest was felicitated with a momentum.	
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CIRCULAR

Date: 04/02/2020

It is therefore announced to all staff that an IPR activity on **CGMP_CFR section on 211 and 210.** will be held at our campus on **11/02/2020**. Everyone is urged to attend the Program without fail in order to acquire the knowledge.

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EVENT REPORT

Name Of The Activity	CGMP_CFR section on 211 and 210.	
Type Of Activity	Workshop	
Date And Time Of Activity	11/02/2020	10.00AM
No. Of Participants	22	
Resource Person	Dr.Prahlad	
Coordinators	Mr. M. Naveen Kumar	
Description	The principal addressed the gathering and welcomed the guest. The HOD introduced about the guest and handled over to the speaker. The speaker said that FDA ensures the quality of drug products by carefully monitoring drug manufacturers' compliance with its Current Good Manufacturing Practice (CGMP) regulations. The CGMP regulations for drugs contain minimum requirements for the methods, facilities, and controls used in manufacturing, processing, and packing of a drug product. The regulations make sure that a product is safe for use, and that it has the ingredients and strength it claims to have. The approval process for new and generic drug marketing applications includes a review of the manufacturer's compliance with the CGMPs. FDA assessors and investigators determine whether the firm has the necessary facilities, equipment, and ability to manufacture the drug it intends to market. Finally the HOD delivered the vote of thanks and the guest was felicitated with a momentum.	
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Date: 20/09/2021

It is therefore announced to all staff that an IPR activity on **Directives for the European Union and Australia** will be held at our campus on **24/09/2021**. Everyone is urged to attend the Program without fail in order to acquire the knowledge.

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Name Of The Activity	Directives for the European Union and Australia	
Type Of Activity	Workshop	
Date And Time Of Activity	24/09/2021	10.00AM
No. Of Participants	32	
Resource Person	Dr.Raju Srivastav	
Coordinators	Mr. M. Naveen Kumar	
Description	The principal addressed the gathering and welcomed the guest. The HOD introduced about the guest and handled over to the speaker. The speaker said that sthe European Union and Australia enjoy a strong, dynamic, and continuously evolving partnership. The relationship is currently based on the 2008 European Union-Australia Partnership Framework, a comprehensive statement of shared values and close historical, political, economic and cultural ties. As our relationship evolved, the EU and Australia have moved to upgrade bilateral ties. To this end, in 2017 the EU, its Member States and Australia signed the EU Australia Framework Agreement. Finally the HOD delivered the vote of thanks and the guest was felicitated with a momentum	
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Date: 18/4/2021

It is therefore announced to all staff that an IPR activity on **Drug and cosmetic regulations in the Japanese market** will be held at our campus on **21/4/2021**. Everyone is urged to attend the Program without fail in order to acquire the knowledge.

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EVENT REPORT

Name Of The Activity	Drug and cosmetic regulations in the Japanese market	
Type Of Activity	Workshop	
Date And Time Of Activity	21/4/2021	10.00AM
No. Of Participants	25	
Resource Person	Dr.Anmol Kher	
Coordinators	Mr. M. Naveen Kumar	
Description	The principal addressed the gathering and welcomed the guest. The HOD introduced about the guest and handled over to the speaker. The speaker said that with Japanese cosmetics are regulated under the *Pharmaceuticals and Medical Devices Act* (hereinafter referred to as "the Act"), which is issued by the competent authority, the Ministry of Health, Labour and Welfare (MHLW), and supported by a series of subsidiary rules, standards and guidance documents. Under the Act, Japan legally classifies cosmetics (in the broad sense of beauty products) into two categories: cosmetics and quasi drugs. The regulations governing each category differ greatly Finally the HOD delivered the vote of thanks and the guest was felicitated with a momentum.	
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Date: 23/3/2021

It is therefore announced to all staff that an IPR activity on **Emerging market drug registration and approval procedures** will be held at our campus on **27/3/2021**. Everyone is urged to attend the Program without fail in order to acquire the knowledge.

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EVENT REPORT

Name Of The Activity	Emerging market drug registration and approval procedures	
Type Of Activity	Seminar	
Date And Time Of Activity	27/3/2021	11.00AM
No. Of Participants	32	
Resource Person	Dr.Shirin Sultana	
Coordinators	Mr. M. Naveen Kumar	
Description	The principal addressed the gathering and welcomed the guest. The HOD introduced about the guest and handled over to the speaker. The speaker stated that Asia is expected to overtake Europe in pharmaceutical market within the next decade and sales are driven by growth in key emerging markets. e.g., China is deemed to be the second largest pharmaceutical market after the United States by 2015. More than 85% population lives in the emerging market and so the real economic growth has come from these markets. This promotes many MNC's switched to these emerging countries particularly in China, India, Russia, Korea and Mexico Finally the HOD delivered the vote of thanks and the guest was felicitated with a momentum.	
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Date: 3/11/2020

It is therefore announced to all staff that an IPR activity on **Methods for Filing a Patent** will be held at our campus on **5/11/2020**. Everyone is urged to attend the Program without fail in order to acquire the knowledge.

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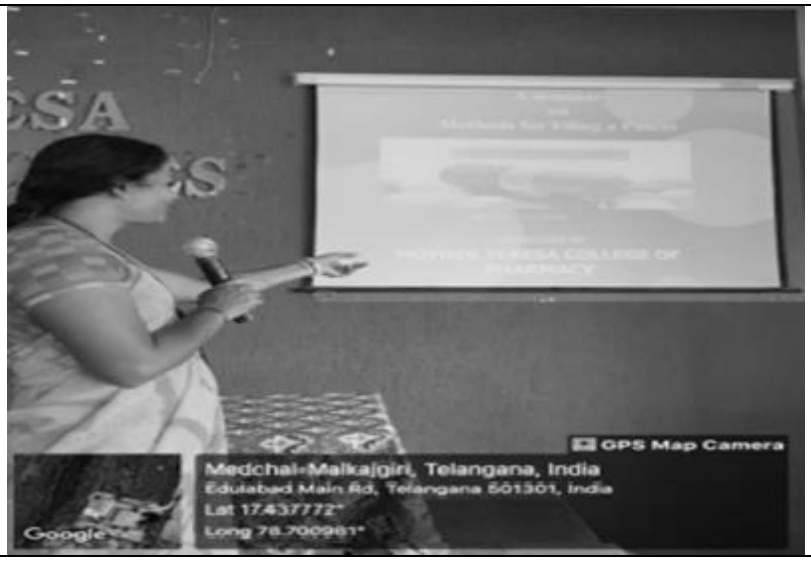
EVENT REPORT

Name Of The Activity	Methods for Filing a Patent	
Type Of Activity	Seminar	
Date And Time Of Activity	5/11/2020	10.30AM
No. Of Participants	31	
Resource Person	Dr.Soujanya Reddy	
Coordinators	Mr. M. Naveen Kumar	
Description	The principal addressed the gathering and welcomed the guest. The HOD introduced about the guest and handled over to the speaker. The speaker explained about the patent as an exclusive right granted for an invention, which is a product or a process that provides, in general, a new way of doing something, or offers a new technical solution to a problem. To get a patent, technical information about the invention must be disclosed to the public in a patent application. Finally the HOD delivered the vote of thanks and the guest was felicitated with a momentum.	
Photo		

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Date: 07/12/2020

It is therefore announced to all staff that an IPR activity on **Rights to intellectual property** will be held at our campus on **11/12/2020**. Everyone is urged to attend the Program without fail in order to acquire the knowledge.

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EVENT REPORT

Name Of The Activity	Rights to intellectual property	
Type Of Activity	Seminar	
Date And Time Of Activity	11/12/2020	10.30AM
No. Of Participants	45	
Resource Person	Dr.Srihari	
Coordinators	Mr. M. Naveen Kumar	
Description	The principal addressed the gathering and welcomed the guest. The HOD introduced about the guest and handled over to the speaker. The speaker stated that the Innovare intellectual property services enables clients to develop comprehensive IP protection plans that take full advantage of patents, trademarks, copyrights and other forms of intellectual property. To create awareness about Intellectual Property Rights (IPR) through conducting a workshop on patent and IPR that enable Universities, Industries, Government Department and Research development institutions for patent searches, patent drafting. Innovare Academic Sciences Pvt Ltd (IAS) works personally with persons to meet their requirements for patent, copyright and trademark information. Finally the HOD delivered the vote of thanks and the guest was felicitated with a momentum.	
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