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IASCC

Indian Association of Supportive Care in Cancer

In collaboration with



2020

26th - 27th Sep

03rd - 04th Oct

05 pm - 09 pm

**SUPPORTIVE CARE
MAKES EXCELLENT
CANCER CARE POSSIBLE**

**LIVE
WEBINAR**



Annual Meeting of IASCC 2020



Click on below link for registration

<http://iascc.co.in/registration>



Annual Meeting of IASCC 2020



Dear Colleague,

IASCC (Indian Association of Supportive Care in Cancer) is an affiliate organization of MASCC, dedicated to improve the supportive care in India. The inception of the organization was declared in September 2018 during Best of MASCC by Dr. Sudeep Gupta in the presence of Dr. Matti Aapro (Past President of MASCC). This organization is completely focused on supportive care required during cancer directed therapies. Other important aspects would be to support good quality research in the same field.

On **September 26th - 27th & October 3rd - 4th**, we are organizing the **2nd Annual Meeting of IASCC 2020**. This IASCC annual meeting will be held on a virtual platform.

The exciting program planned for this conference will have presentations, workshops, and discussions on treatments and techniques for minimising the symptoms of cancers and side effects of cancer directed and related therapy. The conference also includes a diverse array of presentations on the psychological, social, and economic dimensions of cancer diagnosis and treatment. The meeting will feature sessions for doctors and paramedics associated with supportive care in cancer.

We look forward to having your active participation in the niche field of oncology.



Organizing Chairpersons



Dr. Sudeep Gupta
Director, ACTREC
Prof. of Medical Oncology,
Tata Memorial Hospital,
Mumbai



Dr. Shona Nag
Director of Oncology
Sr. Consultant Medical Oncologist,
Sahyadri Group of Hospitals,
Pune



Dr. D C Doval
Chair Medical Oncology & Chief of
Breast Cancer & Thoracic Services,
Rajiv Gandhi Cancer Institute,
Delhi

Organizing Secretaries



Dr. Vikas Ostwal
Prof. of Medical Oncology,
Tata Memorial Hospital,
Mumbai



Dr. Anant Ramaswamy
Asst. Prof. of Medical Oncology,
Tata Memorial Hospital,
Mumbai



Dr. Anant Gokarn
Asst. Prof. of Hemato Oncology,
Tata Memorial Hospital,
Mumbai

Joint Organizing Secretaries



Ms. Anita D'Souza
Professor & Principal,
College of Nursing,
Tata Memorial Hospital,
Mumbai



Ms. Irene Sunder
Nursing Officer in Charge,
CVAD Clinic,
Tata Memorial Hospital,
Mumbai



Highlights of the Meeting

- ④ Second official joint meeting of MASCC and IASCC
- ④ Conference to cover important aspects of supportive care in cancer
- ④ MASCC new annual membership applicants will be entitled to get free life time membership of IASCC and free conference registration for Best of MASCC 2020, India
- ④ Research grant, e-poster and oral presentation awards
- ④ Collaboration with experts in supportive care
- ④ Free one year membership of MASCC and life membership of IASCC for oral or e-poster selected as complimentary



Scientific Program

Day 3 - Saturday, 3rd October

Time

Topic

16.30 - 17.30 Session 1- Oral Abstract Presentation & Research Funding Proposals

16.30 - 17.30 Oral Abstract Presentation
 Judges: Dr. Biswajit Dubashi, Dr. Sadashivadu Gundeti,
 Dr. K. Pavithran, Dr. Ashish Bakshi

17.30 - 18.50 Session 2 : Nutrition in supportive care

Chairpersons Dr. S. D. Banavali, Dr. Mehul Bhansali, Dr. Sudeep Sarkar

Evaluation of nutrition status of cancer patient:
 Tools and modules

17.30 - 17.38 - Pre / Post Chemotherapy
 Speaker: Dr. Mansi Sharma

17.38 - 17.46 - Pre / Post Surgery
 Speaker: Dr. Sanket Mehta

17.46 - 17.54 Interactions of nutrition and cancer and its treatment
 Speaker: Dr. Divya Choudhary

17.54 - 18.02 Myths about nutrition and cancer
 Speaker: Dr. Rajeswari Alagirimunuswamy

18.02 - 18.10 Recent evidences in cancer nutrition in last 3 years
 Speaker: Dr. Rakesh Pinninti

18.10 - 18.20 Role of nutrition as supportive care in gastric cancer
 Speaker: Dr. Y. T. Shivshankar



Scientific Program

Day 3 - Saturday, 3rd October

Time

Topic

18.20 - 18.45

Panel discussion : Evaluation of nutrition status of cancer patient - Tools and modules

Chairpersons

Dr. Rajendra Toprani, Dr. Ashwin Dabhi

Moderator: Dr. Nilesh Chordiya

Panelists: Dr. K M Mohandas

Dr. Mahesh Goel

Dr. Ritika Samaddar

Dr. Dibyendu Sharma

Dr. Mudit Agarwal

Dr. Anil Heroor

Dr. Prathmesh Pai

18.45 - 18.50

Audience Q&A

18.50 - 19.45

Session 3 : Procedures in supportive care

Chairpersons

Dr. Kamran Khan, Dr. Sanjay Dudhat

18.50 - 19.00

Diversion stoma / palliative bypass

Speaker: Dr. Deepak Chhabra

19.00 - 19.10

Angioembolization/ PTBD/ IR procedures as a part of supportive care

Speaker: Dr. Ashwin Polnaya

19.10 - 19.20

Endoscopic procedures as a part of supportive care

Speaker: Dr. Prachi Patil



Scientific Program

Day 3 - Saturday, 3rd October

Time

Topic

19.20 - 19.45 Panel discussion on procedures in supportive care

Chairpersons **Dr. Raman Deshpande, Dr. R. C. Mistry**

Moderator: Dr. Sanjay Sharma

Panelists: Dr. Manish Bhandare

Dr. Deep Goel

Dr. Dushyant Mandalik

Dr. Prasanth Penumadu

Dr. Suyash Kulkarni

Dr. Abhay Kapoor

Dr. Indusekhar Subbanna

Dr. Akash Shukla

19.45 - 19.50 Audience Q&A

19.50 - 20.40 **Session 4 : Digital health for optimal supportive care in oncology: Benefits, limits and future perspectives**

Chairpersons **Dr. Arvind Krishnamurthy, Dr. Saroj Kanta Mishra**

19.50 - 20.00 Tele compliance to treatment

Speaker: Dr. Vijay Patil

20.00 - 20.10 Artificial Intelligence-based tool to generate directional integrative treatment reports for cancer patients (ZIOPAR: ZenOnco.io Integrative Oncology Preliminary Assessment Report)

Speaker: Mr. Kishan Shah

20.10 - 20.20 Cancer Care Management in Digital Era

Speaker: Dr. Amit Jotwani

20.20 - 20.40 Audience Q&A

Moderator: Dr. Bhawna Sirohi

Experts: Dr. Vijay Patil, Dr. Amit Jotwani,

Mr. Kishan Shah, Dr. Swapnil Rane



Scientific Program

Day 4 - Sunday, 4th October

Time

Topic

17.00 - 18.30

Research Funding Proposals

Judges: Dr. S. D. Banavali, Dr. Atul Sharma,
Dr. Sudeep Gupta, Dr. Shona Nag,
Dr. D C Doval, Dr. Sreenivas V

17.30 - 18.50

Session 5 : Supportive care rehab (specific issues)

Chairpersons

Dr. Sanjay Sharma, Dr. Arun Goel

17.30 - 17.40

Lymphedema management

Speaker: Dr. Manjusha Vagal

17.40 - 17.50

Physiotherapy and occupational therapy

Speaker: Dr. Neha Smriti

17.50 - 18.00

Systemic therapy related cardiac toxicity management

Speaker: Dr. Avinash Pandey

18.00 - 18.10

DVT/VTE/PTE

Speaker: Dr. Adwaita Gore

18.10 - 18.20

Vascular access management in cancer patient-global
guidelines and Indian experience

Speaker: Dr. Vineet Talwar

18.20 - 18.26

Taste bud changes with different chemotherapy:
Modification of nutrition and outcome

Speaker: Dr. Amish Vora

18.26 - 18.30

Scalp cooling and exercise to prevent neuropathy

Speaker: Dr. Amish Vora



Scientific Program

Day 4 - Sunday, 4th October

Time

Topic

18:30 - 19:00 Panel Discussion: Establishing an ideal supportive care department

Chairpersons **Dr. Raghunadharao D, Dr. Kaustubh Patel**

Moderator: Dr. Shona Nag

Panelists: Dr. Rebecca Dsouza, Ms. Anita Dsouza, Dr. Sushma Bhatnagar, Ms. Irene Sunder, Dr. Seema Gulia, Dr. Amol Kothekar, Dr. R Ramanjulu, Dr. Sanjay M H, Ms. Rumana Hamied, Dr. Neha Bidvai

19.00 - 20.05 **Session 6 : Toxicity Management**

Chairpersons **Dr. Shekar Patil, Dr. Shyam Aggarwal**

19.00 - 19.10 Skin: HFS / Rash

Speaker: Dr. Vivek Agarwala

19.10 - 19.20 Immunological common toxicities liver/ lung and Gut

Speaker: Dr. Amol Dongre

19.20 - 19.30 CINV management: New strategies ?

Speaker: Dr. Joydeep Ghosh

19.30 - 19.40 Management of oral mucositis secondary to cancer therapy

Speaker: Dr. Rajesh Lalla

19.40 - 20.05 Panel Discussion: Is CTCAE adequate to measure toxicity of systemic therapy?

Chairpersons **Dr. D. C. Doval, Dr. Meenu Walia**

Moderator: Dr. T P Sahoo

Panelists: Dr. Atul Sharma, Dr. Manikandan Dhanushkodi, Dr. Poonam Patil, Dr. Roshni Pandey, Dr. Anubha Bharthuar, Dr. Shailesh Bondarde, Dr. Atul Narayankar



Scientific Program

Day 4 - Sunday, 4th October

Time

Topic

20.05 - 21.00 Session 7 : Management of critical care infections - What to delete, What to switch?

Chairpersons Dr. S H Advani, Dr. Tapan Saikia

20.05 - 20.15 Contraindications of specific drugs based on disease condition
Speaker: Dr. Bhuvan Chugh

20.15 - 20.25 Drug - Drug Interaction
Speaker: Dr. Nandini Menon

20.25 - 20.35 Choosing the right antibiotic
Speaker: Dr. Abdul Ghafur

20.35 - 21.00 Panel discussion on Management of critical care infections: What to delete, What to switch?

Chairpersons Dr. K. Pavithran, Dr. Chanchal Goswami

Moderator: Dr. Kumar Prabhash
*Panelists: Dr. Vamshi Krishna, Dr. Vasant Nagvekar,
Dr. Abdul Ghafur, Dr. Nikhil Ghadyalpatil,
Dr. Vashishth Maniar, Dr. Amol Dongre,
Dr. Bhuvan Chugh, Nandini Menon*

21.00 - 21.05 Vote of thanks

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**ACCESS ABSTRACT POSTER & VIDEO
OF IASCC ANNUAL MEETING 2020**

Chemotherapy-induced bone loss among postmenopausal women with non-metastatic breast cancer

Author: Yadav Nisha

View

Frequency of oral mucositis in elderly patients on Cancer therapy

Author: Prudvi Gundala

View

Study of frequency of myalgia in patients receiving chemotherapy

Author: Praisey Samuel

View

Frequency and severity of oral mucositis in patients receiving chemotherapy

Author: Abhijna Yergolkar

View



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**Clinical patterns of chemotherapy drug related cutaneous eruptions -
prospective observational single center study**

Author: Sai Preeth K

View

Frequency of myalgia in patients with breast cancer

Author: Sahana Sreenivasan

View

**Symptom clusters in patients with gastrointestinal cancer attending a
palliative care setting: a cross sectional study from eastern India**

Author: Chaitanya Patil

View

**Oral cryotherapy: too cool way for prevention of oral mucositis for concurrent
chemo-radiotherapy**

Author: Anusha Thomas

View



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Identification of symptom clusters in advanced head and neck cancer patients attending supportive care clinic: A cross sectional study

Author: Nilesh Dhamne

[View](#)

A study to assess the severity, risk factors and quality of life of patients associated with chemotherapy-induced peripheral neuropathy

Author: Saumya Srivastava

[View](#)

Anemia requiring transfusion, a frequent toxicity of dose-dense chemotherapy, in young indian women on chemotherapy for early breast cancer.

Author: Parth Sharma

[View](#)

Prevalence of distress among cancer patients and factors associated with it - A prospective observational study

Author: Saadvik Raghuram

[View](#)

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A retrospective analysis on the pattern of g-csf usage during the dose dense doxorubicin – Cyclophosphamide and paclitaxel chemotherapy

Author: Gurram Sreeram

[View](#)

Support services infection control at St. Jude India

Author: Vaishali Parte

[View](#)

Effectiveness of game intervention on oral hygiene status among pediatric cancer clients

Author: Supriya Ghanbahadur

[View](#)

Chemotoxicity profile and its associated clinical and socio? Demographical factors among elderly cancer patients receiving chemotherapy

Author: Sindhu Dahagama

[View](#)

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A prospective study to compare caregivers' knowledge and perception of cancer pain with patients' pain assessment and to evaluate their quality of life

Author: Divyesh Kumar

View

Assessment of pain, acute stress disorder, depression and anxiety in patients of cervical carcinoma undergoing brachytherapy to improve supportive care during brachytherapy

Author: Shylini P

View

A phenomenological study on lived experiences of the patients with bladder cancer undergone urinary diversion

Author: Sukhpal Kaur

View





Scientific Program

Day 1 - Saturday, 26th September

Time

Topic

17.00 - 18.10 **Session 1 : Assessment of geriatric cancer patient**

Chairpersons **Dr. Ramesh Sarin, Dr. Madhuchanda Kar**

- 17.00 - 17.10 Role of comprehensive geriatric assessment tools in toxicity prevention
Speaker: Dr. Bhawna Sirohi
- 17.10 - 17.20 Development of geriatric support care unit: Urgent need of the hour
Speaker: Dr. Anant Ramaswamy
- 17.20 - 17.30 Polypharmacy in geriatric oncology patients
Speaker: Dr. Chetan Deshmukh
- 17.30 - 17.40 Role of a geriatric nurse in geriatric oncology
Speaker: Ms. Prathepa Jagdish
- 17.40 - 17.50 Scope and impact of comorbidities in geriatric cancers
Speaker: Dr. Nidhhi Tandon
- 17.50 - 18.10 Panel discussion on above topics
Moderator: Dr. Rohit Pai
Panelists: Dr. Raja Pramanik
 Dr. Rejiv Rajendranath
 Dr. Tejinder Singh
 Dr. Shruti Kate
 Dr. Lalit Mohan Sharma
 Dr. Manisha Singh
 Dr. Kiran Kattimani



Scientific Program

Day 1 - Saturday, 26th September

Time

Topic

18.10 - 18.40 Session 2 : Management of pain

Chairperson Dr. P. N. Jain

18.10 - 18.20 Pain management in older patients: The age effect
Speaker: Dr. Raghu Thota

18.20 - 18.30 Newer strategies in the management of chronic pain
Speaker: Dr. Madhuri Lokapur

18.30 - 18.40 Early integration of palliative care into mainstream oncology practice
Speaker: Dr. Prakash Fernandes

18.40 - 18.50 Session 3 : Informational needs of older adults with cancer

Chairpersons Mrs. Vandana Gupta, Dr. Mary Muckaden

18.40 - 18.50 Holistic supportive care in elderly
Speaker: Dr. Lakshmi Haridas

18.50 - 19.50 Session 4 : Stoma care and central venous access device

Chairperson Ms. Irene Sunder

18.50 - 19.20 Stoma care
Speaker: Ms. Sonal Rane

19.20 - 19.50 Central venous access workshop
Speaker: Ms. Priyanka Lad



Scientific Program

Day 2 - Sunday, 27th September

Time

Topic

17.00 - 17.50 **Session 5 : Needs of adolescents and young adults with cancer - Beyond no man's land**

Chairpersons **Dr. N. K. Warriar, Dr. Amit Joshi**

17.00 - 17.10 Young cancer patients: Are we supporting them enough ?

Speaker: Dr. Jyoti Bajpai

17.10 - 17.20 Development of ACT Clinic

Speaker: Dr. Amol Patel

17.20 - 17.50 Panel discussion on above topics

Moderator: Dr. Gauri Kapoor

Panelists: Ms. Rashmi Methry, Dr. Biswajit Dubashi
Ms. Ruby Ahluwalia, Dr. Ghanshyam Biswas

17.50 - 18.30 **Session 6 : Cognitive and endocrine disorders in adolescents and young adult's cancers**

Chairpersons **Dr. Prakash Chitalkar, Dr. K. Sambasivaiah**

17.50 - 18.00 Cognitive issues

Speaker: Dr. Prabhat Bhargava

18.00 - 18.10 Endocrine disorders and cancer treatment

Speaker: Dr. Vaishali Deshmukh

18.10 - 18.30 Panel discussion on above topics

Moderator: Dr. Amit Joshi

Panelists: Dr. Venkatraman Radhakrishnan
Dr. Prabhat Bhargava, Dr. Pritam Kataria
Dr. Simran Kaur, Dr. Vaishali Deshmukh
Dr. Bhavesh Poladia, Dr. Ashay Karpe
Dr. Ravi Tippeswamy



Annual Meeting of IASCC 2020



Scientific Program

Day 2 - Sunday, 27th September

Time

Topic

18.30 - 18.50

Session 7 : Fertility and sexuality related issues in adolescents and young adult patients across cancer care continuum

Chairpersons

Dr. Shona Nag, Dr. Sudeep Gupta

18.30 - 18.40

Fertility and cancer treatment

Speaker: Dr. Amita Maheshwari

18.40 - 18.50

Psychosexual interventions in adolescents and young adult cancer survivors

Speaker: Dr. Savita Goswami

18.50 - 19.10

Chemotherapy associated toxicity and supportive care- Role of APN (Advanced Practice Nurse)

Speaker: Ms. Kim Lee



Annual Meeting of IASCC 2020



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Annual membership fees for MASCC is \$25 USD

To pay MASCC annual membership, log on to www.mascc.org

Form to be filled by MASCC new members

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Qualification : _____

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Mobile number : _____ Email Id: _____

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MASCC membership fees amount paid : _____

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Cama Industrial Premises Co-Op Society Ltd., Sunmill Compound, Lower Parel (W),
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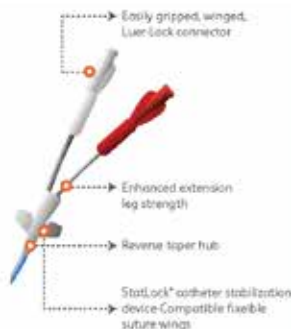
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1000 mg iron can be given in a single dose

Convenient IV infusion of 15 minutes and flexible dosing**
Up to 200 mg as a IV bolus injection
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Reference: 1. Hedenus M, Karlsson T, Ludwig H, Rzychon B, Felder M, Roubert B, et al. Intravenous iron alone resolves anemia in patients with functional iron deficiency and lymphoid malignancies undergoing chemotherapy. Medical Oncology. 2014 Jun; 31(12).
*Data on file ** Encicarb prescribing information
CIA: Chemotherapy-induced Anemia; FID: Functional Iron Deficiency; FCM: Ferric Carboxymaltose ESA: Erythropoiesis-stimulating Agent Hb: Hemoglobin

Abridged Prescribing Information

ENCICARB INJECTION

Composition: Each ml contains Ferric Carboxymaltose equivalent to elemental iron 50 mg. **Presentation:** Vials of 20 ml. For further details, please consult the full prescribing information. **Indications:** For treatment of iron deficiency when oral iron preparations are ineffective or cannot be used. **Dosage:** The cumulative dose for repletion of iron using ferric carboxymaltose is determined based on the patient's body weight and haemoglobin (Hb) level and must not be exceeded. For Hb = 10 g/dl - body weight 35 kg to <70 kg: 1500 mg, body weight ≥70 kg: 2000 mg. For Hb = 10 g/dl - body weight 35 kg to <70 kg: 1000 mg, body weight ≥70 kg: 1500 mg. A cumulative iron dose of 500 mg should not be exceeded for patients with body weight < 35 kg. For overweight patients, a normal body weight/blood volume relation should be assumed when determining the iron requirement. Maximum tolerated single dose: 1000 mg of iron (20 ml) per day or 15 mg of iron (0.3 ml) per kg body weight. Do not administer 1000 mg of iron (20 ml) more than once a week. **Intravenous injection:** Undiluted solution up to 1000 mg iron. For doses greater than 200 and up to 500 mg iron, ferric carboxymaltose should be administered at a rate of 100 mg/min. For doses greater than 500 and up to 1000 mg iron, ferric carboxymaltose should be administered over 15 minutes. **Intravenous drip infusion:** Intravenous infusion up to a maximum single dose of 20 ml of Ferric Carboxymaltose Injection (1000 mg of iron). Ferric Carboxymaltose Injection must be diluted only in sterile 0.9% sodium chloride solution. A single maximum daily injection dose of 200 mg iron should not be exceeded in haemodialysis-dependent chronic kidney disease patients. **Contraindications:** Contraindicated in cases of known hypersensitivity to Ferric Carboxymaltose Injection or to any of its excipients, anemia not attributed to iron deficiency (e.g. other microcytic anaemias), evidence of iron overload or disturbances in utilization of iron, and in pregnancy in the first trimester. **Adverse reactions:** Headache, dizziness, nausea, abdominal pain, constipation, diarrhea, injection site reactions and rash are commonly reported adverse reactions. **Use in special populations:** Pregnancy: A careful risk/benefit evaluation is required before use during pregnancy. Use during pregnancy may influence skeletal development in the fetus. **Lactation:** Based on limited data on nursing women it is unlikely that Ferric Carboxymaltose Injection represents a risk to the nursing child. **Overdosage:** May lead to accumulation of iron in storage sites eventually leading to haemosiderosis. Monitoring of iron parameters such as serum ferritin and transferrin saturation may assist in recognizing iron accumulation.



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Abbreviated prescribing information (Lenvima):

Trade Name: Lenvima. **Active ingredient:** Lenvatinib. **Pharmaceutical form:** 4 mg and 10 mg hard capsule. **Pharmacology:** Lenvima (Lenvatinib) is a receptor tyrosine kinase inhibitor that inhibits the kinase activities of vascular endothelial growth factor (VEGF) receptors VEGF1 (FLT1), VEGFR2 (KDR) and VEGFR3 (FLT4). Lenvatinib also inhibits other RTKs that have been implicated in pathogenic angiogenesis, tumour growth and cancer progression in addition to their normal cellular functions including fibroblast growth factor (FGF) receptors FGFR1, 2, 3 and 4; the platelet derived growth factor receptor alpha (PDGFR α), KIT and RET. **Indications:** Lenvima is indicated for the treatment of patients with locally recurrent or metastatic, progressive, radioactive iodine-refractory differentiated thyroid cancer (DTC). Lenvima is also indicated for the first-line treatment of patients with unresectable hepatocellular carcinoma (HCC). **Dose and Administration:** **DTC:** The recommended daily dose of Lenvatinib is 24 mg (two 10 mg capsules and one 4 mg capsule) taken once daily. The daily dose is to be modified as needed according to the dose/toxicity management plan. **Recommended Dosage for HCC:** based on actual body weight: 12 mg for patients greater than or equal to 60 kg (three 4mg capsules) or 8 mg for patients less than 60 kg (two 4mg capsules). The daily dose is to be modified as needed according to the dose/toxicity management plan. Take LENVIMA orally once daily until disease progression or until unacceptable toxicity. **Precautions:** Hypertension is commonly manifested side effect. Control blood pressure prior to treatment with Lenvima. Withhold Lenvima for grade 3

hypertension. **Contraindication:** None. **Warning:** Hypertension; proteinuria; renal failure and impairment/ GI toxicity; cardiac failure; reversible posterior leukoencephalopathy syndrome (RPLS); hepatotoxicity; hemorrhagic events; arterial thromboembolic events (ATES); fistula formation and gastrointestinal perforation; QT interval prolongation; hypocalcaemia; thyroid dysfunction and impairment of thyroid stimulating hormone suppression; wound healing complications; etc. **Drug interactions:** No drug interactions are expected with CYP3A, P-glycoprotein (P-gp), breast cancer resistance protein (BCRP) inhibitors and CYP3A and P-gp inducers. **Adverse reactions:** In **DTC patients:** The most common adverse reactions observed in LENVIMA-treated patients ($\geq 30\%$) were, in order of decreasing frequency, hypertension, fatigue, diarrhea, arthralgia/myalgia, decreased appetite, decreased weight, nausea, stomatitis, headache, vomiting, proteinuria, palmar-plantar erythrodysesthesia (PPE) syndrome, abdominal pain, and dysphonia. In **HCC patients:** The most common adverse reactions observed in the LENVIMA-treated patients ($\geq 20\%$) were, in order of decreasing frequency, hypertension, fatigue, diarrhea, decreased appetite, arthralgia/myalgia, decreased weight, abdominal pain, palmar-plantar erythrodysesthesia syndrome, proteinuria, dysphonia, hemorrhagic events, hypothyroidism, and nausea. **Overdose:** There is no specific antidote for overdose with Lenvatinib. In case of suspected overdose, Lenvatinib should be withheld and supportive care initiated. Storage: Lenvima should be stored below 30°C. Full prescribing information is available on request.

1D-11-09B-05



For full prescribing information please write to :

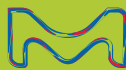
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Right
time

In Locally
advanced &
1st line
RM SCCHN

In 1st line
RAS WT
mCRC

Right
patient

Right
response

our Mission
never stops

Erbitux (Cetuximab) Abbreviated Prescribing Information:

SCHEDULE H PRESCRIPTION DRUG – CAUTION Not to be sold by retail without the prescription of a Registered Medical Practitioner

Warning: To be sold by retail on the prescription of a Oncologist only. Before prescribing ERBITUX, please consult full prescribing information. **Presentation:** ERBITUX 5 mg/mL solution for infusion. Excipients: sodium chloride, glycine, polysorbate 80, citric acid monohydrate, sodium hydroxide, water for injections. **Indications:** Epidermal growth factor receptor-expressing, RAS wild-type metastatic colorectal cancer (mCRC) in combination with irinotecan-based chemotherapy (CT), or in first-line in combination with FOLFOX, or as a single agent in patients who have failed oxaliplatin- and irinotecan-based therapy and who are intolerant to irinotecan. Squamous cell carcinoma of the head and neck (SCCHN): in combination with radiation therapy (RT) for locally advanced (LA) disease or with platinum-based chemotherapy (p-CT) for recurrent and/or metastatic (RM) disease. **Dosage and administration:** Once a week, intravenously with an infusion pump, gravity drip or a syringe pump; separate infusion line. Initial dose 400 mg/m² (should be given slowly with max. infusion rate: 5 mg/min); the recommended infusion period is over 120 mins; subsequent weekly doses 250 mg/m² (Max. infusion rate: 10 mg/min; recommended over 60 mins). Supervision/monitoring by a physician experienced in antineoplastic therapy throughout infusion and for at least one hour afterwards is required. Resuscitation equipment must be ensured. Prior to first infusion, premedication with antihistamines and corticosteroids at least 1 hour prior to administration of ERBITUX; also recommended for all subsequent infusions. Administer CT not earlier than one hour after ERBITUX infusion. mCRC: administer ERBITUX until disease progression. Wild-type RAS tumor status must be verified prior to first infusion by an experienced laboratory using validated test methods. LA SCCHN: start ERBITUX therapy one week before RT and continue throughout treatment. RM SCCHN: administer ERBITUX in combination with p-CT and continue until disease progression. **Special Populations:** Elderly: no dose adjustment required (limited experience in patients >75 years). Pediatric patients (<18 years): efficacy not established, no new safety signals. (Only patients with adequate renal, hepatic and hematological parameters have been investigated. **Contraindications:** Known severe hypersensitivity reactions (grade 3/4 NCI-CTCAE). In combination with oxaliplatin-containing CT in mutated/unknown RAS status. Contraindications for concomitancy used CT or RT must be considered. **Special warnings and precautions:** Severe infusion-related reactions (IRR): including anaphylactic reactions. May commonly occur, in some cases with fatal outcome; immediate and permanent discontinuation of ERBITUX therapy, may necessitate emergency treatment. May be anaphylactic or anaphylactoid in nature or represent a cytokine release syndrome. Symptoms may occur during the first infusion and for up to several hours afterwards or with subsequent infusions and may include bronchospasm, urticaria, increase or decrease in blood pressure, loss of consciousness or shock. In rare cases, angina pectoris, myocardial infarction or cardiac arrest have been observed. The risk for anaphylactic reactions is much increased in patients with a history of allergy to red meat or tick bites or positive results of tests for IgE antibodies against fibrinogen. Mild/moderate IRRs: decrease infusion rate, also for all subsequent infusions. Closely monitor patients with reduced performance status (PS) and pre-existing cardio-pulmonary disease. Skin reactions: oral tetracyclines and topical 1% hydrocortisone cream with moisturizer may be considered for prophylactic use and medium to high-potency topical corticosteroids or oral tetracyclines for treatment (see clinical practice guidelines). Severe skin reaction (grade 3): interrupt treatment, only resume if reaction resolves to grade 2. Second or third occurrence of severe skin reactions: resume at lower dose (200 mg/m² after second, 150 mg/m² after third) only if reaction resolves to grade 2. Fourth occurrence or failure to resolve to grade 2 during interruption: permanent discontinuation. **Intermittent lung disease:** if diagnosed, discontinuation and appropriate treatment. Electrolyte disturbances: determination of serum electrolytes recommended prior to and periodically during treatment. Electrolyte depletion (e.g. hypomagnesemia, hypokalaemia as a consequence of diarrhea), hypocalcemia, particularly in combination with p-CT is recommended. **Neuropathy and related infectious complications:** careful monitoring is recommended particularly in patients experiencing skin lesions, mucositis or diarrhea that may facilitate the occurrence of infections. **Severe and sometimes fatal cardiovascular events:** increased frequency associated with age ≥ 65 years or PS has been observed. Patient cardiovascular status, PS and concomitant administration of cardiotoxic compounds (e.g. fluoropyrimidines) should be taken into account. **Acute or warning symptoms of keratitis:** refer promptly to an ophthalmologist, consider benefit/risk of continuing use. **Confirmed ocular keratitis:** interruption or discontinuation of ERBITUX. Use with caution in patients with history of keratitis, ulcerative keratitis or severe dry eye (e.g. use of contact lenses). CRC patients with mutated/unknown RAS status: ERBITUX should not be used since negative effects on PS and OS as well as on FOLFOLX have been reported in RAS mutated tumors. There is limited experience in combination with RT in mCRC. **Fertility, pregnancy and lactation:** Only use during pregnancy or in women with inadequate contraception if potential benefits justify potential risks to fetus. Breast-feeding during treatment and 2 months later is not recommended. Effects on male/female fertility have not been evaluated. **Undesirable effects:** Very common (>1/10): skin reactions (e.g. acne-like rash and/or pruritus), dry skin, desquamation, hyperhidrosis, or nail disorders, single cases of skin necrosis, hypomagnesemia, mild/moderate IRRs (e.g. fever, chills, dizziness, dyspnea), increased liver enzyme levels and mucositis, in some cases severe. Mucositis may lead to ulcers. Common (<1/10): headache, conjunctivitis, diarrhea, nausea, vomiting, hypocalcemia, anorexia, weight loss, severe IRRs. **Discontinuation (<1/100, <1/1000):** pleuritis, keratitis, deep vein thrombosis, pulmonary embolism or interstitial lung disease. Very rare (<1/10,000): Stevens-Johnson syndrome/toxic epidermal necrolysis. **Frequency not known:** superinfection of skin lesions with subsequent complications (e.g. cellulitis, erysipelas, staphylococcal scalded skin syndrome, necrotizing fasciitis, sepsis, septic meningitis). In combination with local RT in SCCHN: typical undesirable effects of RT (e.g. mucositis, radiation dermatitis, dysphagia or leukopenia, mainly as a myelosuppressive). In combination with ERBITUX: slightly higher rates of severe acute radiation dermatitis, mucositis and late RT-related events. **Interactions:** Fluoropyrimidines: increased frequency of hand-foot syndrome and cardiac ischaemia (e.g. myocardial infarction and congestive heart failure). **Capacitans and oxaliplatin (CCEQs):** frequency of severe diarrhea may be increased. **Ac-T:** increased frequency of severe leukopenia/neutropenia, which may lead to a higher rate of febrile neutropenia, pneumonia and sepsis. Storage: Store in a refrigerator (2°C – 8°C) Shelf life: 48 months

Date of Information: June 2019 Based on CDEs of Cetuximab V.16.0 dated 28th June 2019.

For further information refer to full prescribing information or write to:

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Godrej One, 8th Floor, Pirojsha Nagar, Eastern Express Highway, Vikhroli (East) Mumbai – 400079

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