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PHARMACYDAILY.COM.AU

Today's issue of PD

Pharmacy Daily today has two pages of news, plus a full page featuring the **MIMS August update**.

RA market blows out

WITH the increased use of biologics, drugs with newer mechanisms of action and combination therapies, the biosimilar sector of the global rheumatoid arthritis (RA) market is trending strongly with a compound annual growth rate (CAGR) of 71% according to BCC Research.

While the global market for therapies for RA is expected to grow at a CAGR of 1.3%, to \$17.2b by 2020, biosimilars for major biologic products etanercept, infliximab and rituximab are dominating growth.

CLICK HERE to access the data.

Did you know
that goat's milk
contains prebiotic
oligos which are
structurally closer
to breast milk*



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* When compared to standard cow's milk;
Source: Urashima and Taufik, 2010.

Health Care Homes report

THE government's proposed Patient Centred 'Health Care Home' (PCHCH) concept should be extended to all applicable patients, not just those with chronic conditions, according to a report released yesterday by a coalition of organisations including the Consumers Health Forum of Australia and the Royal Australian College of General Practitioners.

The report summarises consensus principles from a roundtable convened in Melbourne last month attended by stakeholders from across the sector, including the PSA.

The PCHCH would place multidisciplinary teams in a one-stop health care hub to help coordinate care and treatment focused around the patient.

In the May 2016 Budget funding for a first stage of the Health Care Home was announced, and from Jul 2017 up to 65,000 patients should be able to voluntarily enrol in a trial of Health Care Homes in up to 200 practices across seven Primary

Lung health list

LUNG Foundation Australia has developed Australia's first national Lung Health Checklist for the Indigenous community, together with the Qld govt's Indigenous Respiratory Outreach Care Program.

Eight simple questions help recognise symptoms and risks of lung disease and the need to act promptly - for details see lungfoundation.com.au.

MA fines BMS \$10k

DUE to Bristol-Myers Squibb (BMS) being found in breach of sections of the Medicines Australia (MA) Code of Conduct on certain matters relating to a doctor meeting, the company has been ordered to pay a fine of \$10,000.

Bizarrely, when rival MSD appealed the sanction as too limited, but lost the appeal, the Medicines Australia Appeals Committee retained the company's \$20,000 appeal bond, meaning the matter cost MSD twice as much as the original BMS fine.

CLICK HERE for the ruling.

Health Network regions.

The effectiveness of the trial will then be reviewed after two years.

The report, which summarises models in the US and Canada, said an integrated system would "provide patients with more individualised attention including tailored care, alternative consultation modes, and online access to their own health information, tests & appointments.

Recommendations include building IT infrastructure to enable shared care planning and coordinated care, including health care team access to pharmacy, pathology and diagnostic imaging.

View the report at chf.org.au.

Certificate for Leave

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AHPRA ad sanction

THE Australian Health Professions Regulatory Agency has charged a NSW chiropractor with breaching advertising requirements, alleging his website advertised chiropractic services in a way that was likely to be "false, misleading or deceptive".

Without commenting directly on the matter which is now before the courts, AHPRA ceo Martin Fletcher said the agency took its role of protecting the public very seriously.

"Anyone advertising a regulated health service, regardless of whether they are registered health practitioners or not, must meet the requirements of the law," he said.

AHPRA said chiropractors "must practise in an evidence-based way".

Cincotta Mascot award

ANTHONY Vass, owner of the Cincotta Discount Chemist Mascot was named NSW Pharmacist of the Year for 2016 at a recent PSA (NSW) dinner at Sydney's Oatlands House.

The pharmacy is a reference site for the PSA's Health Destination Pharmacy program.

Vass said a "lightbulb moment" was a turning point for him when he attended the AIM High Adherence training program initiated by Cincotta Discount Chemist in conjunction with the University of Technology Sydney.

He said at that point he realised he hadn't fully engaged with patients, but by "Taking a moment to ask customers simple questions

about how and when they take their medication has opened up a whole new understanding of how I can become more effective as a health professional."

The Aim High Adherence and HDP programs were an "unqualified success," he said and his store now has growth in all areas and 200 more community Webster-pak customers than before.



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New Products

Fraxiparine (nadroparin calcium)

is a low molecular weight heparin made by depolymerisation of standard heparin. It is a glycosaminoglycan with a mean molecular weight around 4,500 daltons. It possesses a high ratio of anti-Xa activity to anti-IIa activity between 2.5 to 4.0 compared to unfractionated heparin for which this ratio is one. Nadroparin has both immediate and prolonged antithrombotic action. Nadroparin exhibits a high-affinity binding to the plasma protein anti-thrombin III (ATIII). This binding leads to an accelerated inhibition of factor Xa and to a lesser extent, factor IIa (Anti-Xa:Anti-IIa ratio of 3.6:1), which contributes to the antithrombotic potential of nadroparin. Fraxiparine is indicated for prophylaxis against deep vein thrombosis (DVT) associated with general or orthopaedic surgery and for treatment of DVT and in the prevention of clotting during haemodialysis. Fraxiparine is contraindicated in patients with: a history of thrombocytopenia with nadroparin; an increased risk of haemorrhage including those with bleeding disorders (except for disseminated intravascular coagulation not induced by heparin); active bleeding or organic lesions likely to bleed (such as active peptic ulceration), haemorrhagic cerebrovascular accident or infective endocarditis and severe renal failure (creatinine clearance < 30 mL/min) receiving treatment for DVT. Fraxiparine is available as 1900 IU/0.2 mL, 2850 IU/0.3 mL, 3800 IU/0.4 mL, 5700 IU/0.6 mL, 7600 IU/0.8 mL and 9500 IU/1 mL in single use prefilled glass syringe, packs of 2's.

Herceptin SC (trastuzumab (rch))

is a recombinant DNA derived humanized monoclonal antibody that selectively targets the extracellular domain of the human epidermal growth factor receptor 2 protein (HER2). Herceptin SC is indicated for the following. The treatment of HER2-positive early breast cancer following surgery, and in association with chemotherapy and, if applicable, radiotherapy. The treatment of

HER2-positive locally advanced breast cancer in combination with neoadjuvant chemotherapy followed by adjuvant Herceptin. The treatment of patients with metastatic breast cancer who have tumours that overexpress HER2: as monotherapy for the treatment of those patients who have received one or more chemotherapy regimens for their metastatic disease; in combination with taxanes for the treatment of those patients who have not received chemotherapy for their metastatic disease; or in combination with an aromatase inhibitor for the treatment of post-menopausal patients with hormone-receptor positive metastatic breast cancer. Herceptin is contraindicated in the treatment of early or locally advanced breast cancer. It is also contraindicated in patients with a left ventricular ejection fraction of less than 45% and those with symptomatic heart failure. It is also contraindicated in patients with known hypersensitivity to Chinese hamster ovary cell proteins. Herceptin SC is available as a ready to use solution (600 mg/5 mL) in a single use vial in packs of 1's.

Lonquex (lipegfilgrastim (rbe))

is a long acting form of recombinant human granulocyte colony stimulating factor (G-CSF). Human G-CSF is a glycoprotein that regulates the production and release of functional neutrophils from the bone marrow. Filgrastim is an unglycosylated recombinant methionyl human G-CSF. Lipegfilgrastim is a sustained duration form of filgrastim due to decreased renal clearance. Lipegfilgrastim binds to the human G-CSF receptor like filgrastim and pegfilgrastim. Lonquex is indicated for reduction in the duration of neutropenia and the incidence of febrile neutropenia in adult patients treated with cytotoxic chemotherapy for malignancy (with the exception of chronic myeloid leukaemia and myelodysplastic syndromes). Lonquex is contraindicated in patients with known hypersensitivity to other G-CSF products including lenograstim, pegfilgrastim and filgrastim. Lonquex is available as

a single use prefilled glass syringe containing a solution volume of 0.6 mL with 6 mg of lipegfilgrastim, for a 10 mg/mL solution in packs of 1's.

Sylvant (siltuximab) is a chimeric (human murine) immunoglobulin G1k (IgG1k) monoclonal antibody against human Interleukin-6 (IL-6) produced in a Chinese hamster ovary (CHO) cell line. Siltuximab prevents the binding of human IL-6 to both soluble and membrane bound IL-6 receptors (IL-6R), thus inhibiting the formation of the hexameric signalling complex with gp130 on the cell surface. Overproduction of IL-6 has been hypothesised to play a central role in driving plasma cell proliferation and systemic manifestations in patients with Castleman's disease. Sylvant is indicated for the treatment of patients with multicentric Castleman's disease (MCD) who are human immunodeficiency virus (HIV) negative and human herpesvirus-8 (HHV-8) negative. Sylvant is available as a 100 mg and 400 mg powder for infusion in a single use glass vial in packs of 1's.

Safety Related Changes

Eviplera (300 mg tenofovir disoproxil fumarate/200 mg emtricitabine/25 mg rilpivirine) should not be coadministered with rilpivirine unless required for dose adjustment (e.g. with rifabutin).

Humira (adalimumab (rch)) is now indicated for the treatment of active moderate to severe hidradenitis suppurativa (acne inversa) in adult patients with an inadequate response to conventional systemic hidradenitis suppurativa therapy.

Nolvadex (tamoxifen citrate)

is now indicated for the primary reduction of breast cancer risk in women either at moderately increased risk (lifetime breast cancer risk 1.5 to 3 times the population average) or high risk (lifetime breast cancer risk greater than 3 times the population average).

This list is a summary of only some of the changes that have occurred over the last month. Before prescribing, always refer to the full product information.