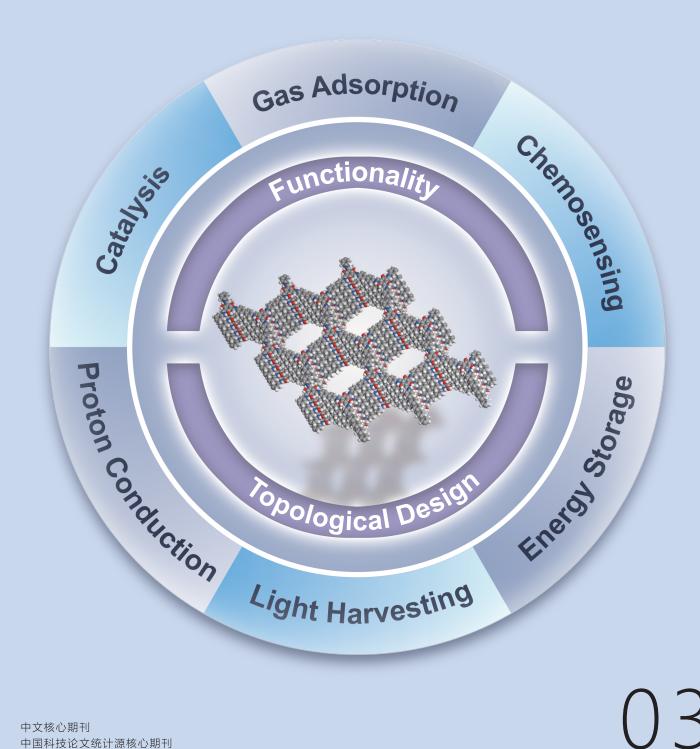
# 功能高分子学报

# JOURNAL OF FUNCTIONAL POLYMERS



中文核心期刊

中国科技论文统计源核心期刊

中国科学引文数据库来源期刊

中国学术期刊文摘来源期刊

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## 功 能 高 分 子 学 报 第 31 卷 第 3 期 2018 年 6 月

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## 目 次

## 特约综述

共价有机骨架的设计、制备及应用 ……………………………… 周 婷, 龚祎凡, 郭 佳 (189) 研究论文 还原响应型嵌段共聚物自组装纳米胶束作为 siRNA 运输载体的研究 (216)两亲刚性三嵌段聚多肽的合成及其自组装行为 …………………………… 陈 智, 姚 远, 唐颂超 (225)柠檬酸改性壳聚糖水凝胶的制备与性能 (232)基于遥爪型大分子交联剂的动态化学交联水凝胶的黏弹性质 (241)接枝型镉离子印迹膜的制备及其识别选择性能 (248)透明质酸聚合物胶束的制备及其内涵体的 pH 敏感性 ………… 刘艳华, 周成铭, 杨 (255)聚乳酸纤维对热塑性淀粉塑料性能的影响 (261)多孔载药 n-HA/PEEK/CS 复合材料的制备与性能 ·························· 周浩浩, 曹慧群, 周 莉, 罗仲宽, 胡惠媛, 黄丹榕 (267)高内相乳液模板法制备淀粉基大孔材料及表征 …………… 张守村, 孙 武, 皮 茂 (273)熔融聚合氨基酸的体外生物活性 ………………………………… 樊晓霞, 严永刚 (279)《功能高分子学报》第一届青年编辑委员会 ......(封二) 《功能高分子学报》征稿简则 ......(封三)

### JOURNAL OF FUNCTIONAL POLYMERS

Vol .31 No .3 June 2018 CONTENTS

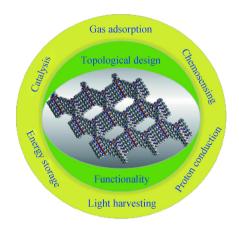
#### Special Review

#### Covalent Organic Frameworks: Design, Synthesis and Applications

ZHOU Ting, GONG Yi-fan, GUO Jia

Journal of Functional Polymers, 2018, 31(3):189-215.

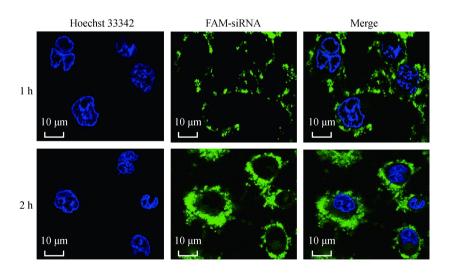
COFs show outstanding physical and chemical properties by virtue of the designability, tailorability and functionality of the extended ordering structure. The diversity of building blocks, specific topological structure, controllable pore environment and flexible functional design make COFs become a fantastic platform for broad applications such as gas storage, catalysis, chemosensing, light harvesting, proton conduction and energy storage.



#### **Papers**

#### Redox-Responsive Micelles Self-assembled from Multi-block Copolymer for Delivery of siRNA

LIU Ben-xin, HE Chang-yu, TAN Lian-jiang, LIU Bing-ya, ZHU Zheng-gang, GONG Bing, SHEN Yu-mei Journal of Functional Polymers, 2018, 31(3):216-224.



Amphiphilic cationic multi-block copolymers were synthesized and self-assembled into micelles for delivery of siRNA. Confocal laser scanning microscopy results demonstrate that the FAM-siRNA can be delivered into SGC7901 cells by micelles.

#### Synthesis and Self-assembly of Amphiphilic Rigid Triblock Copolypeptides

CHEN Zhi, YAO Yuan, TANG Song-chao

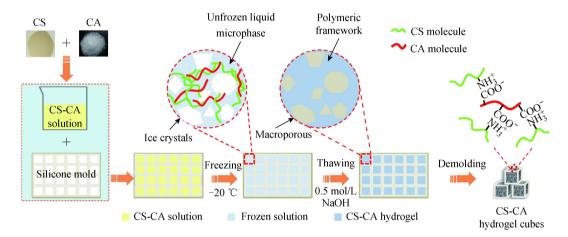
Journal of Functional Polymers, 2018, 31(3):225-231.

N-H + N-H 
$$\frac{N\text{-TMS}}{\text{DMF}}$$
  $\frac{V(\text{Water}): V(\text{DMF})=11:1}{\text{Self-assembly}}$   $\frac{V(\text{Water}): V(\text{DMF})=11:1}{\text{Self-assembly}}$ 

Amphiphilic rigid triblock polypeptides were synthesized via NCA ring opening polymerization using N-TMS as the initiator, and self-assembled into rod-like large compound micelles in mixed system of DFM-water. The unusual all-rigid skeletons and the confined conformation of the amphiphilic triblock copolypeptides might be the reason of these results.

#### Fabrication and Characterization of Citric Acid Modified Chitosan Hydrogel

YANG Shang-ying, YUAN Hui-hua, YI Bing-cheng, WANG Xian-liu, ZHOU Ying, ZHANG Yan-zhong *Journal of Functional Polymers*, 2018, 31(3):232-240.



Chitosan-citric acid (CS-CA) hydrogel was formed after a freeze-gelation process with the formulated CS-CA solutions. The newly developed CS-based hydrogel with improved wet mechanical properties could be applied in wound dressings or as scaffolds for engineering different tissues.

# Viscoelastic Properties of Dynamic Chemical Crosslinking Hydrogel Based on Telechelic Macromolecular Crosslinker YE Zhi-peng, XIE Man-qing, XIE Jia-li, ZHAO Chuan-zhuang

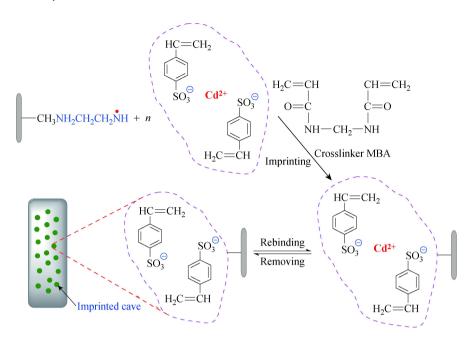
Journal of Functional Polymers, 2018, 31(3):241-247.

$$\begin{array}{c} \text{Chitosan} \equiv \\ \text{DF-PEG} \end{array} \end{array} \equiv \begin{array}{c} \text{OH} \\ \text{NH}_2 \\ \text{O} \\ \text{O} \end{array} \longrightarrow \begin{array}{c} \text{OH} \\ \text{NH}_2 \\ \text{O} \end{array} \longrightarrow \begin{array}{c} \text{Dangling} \\ \text{Elastic-active Intrachain} \end{array}$$

The density of elastic-active cross-linker, which determines the modulus, relaxation time and the shear thickening behavior of hydrogel, can be tuned by molecular weight and concentration of chitosan or DF-PEG.

#### Preparation of Graft Cd2+ Ion-Imprinted Membrane and Its Recognition Selectivity

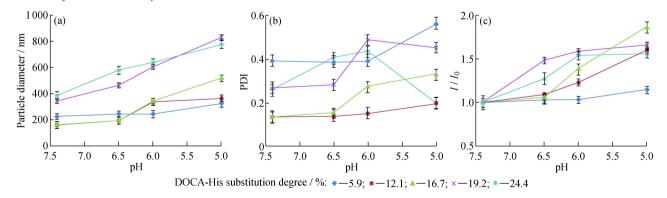
LIU Chun-yan, ZHANG Zheng-guo, LEI Qing-juan, GAO Bao-jiao, AN Fu-qiang, YANG Lei *Journal of Functional Polymers*, 2018, 31(3):248-254.



Free radicals generated on the AMPSF membrane surface lead to the occurrence of graft crosslinking-polymerization between SSS monomer surrounding  $Cd^{2+}$  and cross-linker MBA, thus embedding the  $Cd^{2+}$  into the cross-linked network and synchronizing graft/cross-linking polymerization with ion imprinting. The  $Cd^{2+}$  template was washed away and  $Cd^{2+}$  ion-imprinted membranes with grafting type (GIIM) were prepared, which have sound performances in binding affinity, recognition selectivity and osmotic separation of  $Cd^{2+}$ .

#### Preparation and Endosome pH Sensitivity of Hyaluronic Acid Polymeric Micelles

LIU Yan-hua, ZHOU Cheng-ming, YANG Tong Journal of Functional Polymers, 2018, 31(3):255-260.

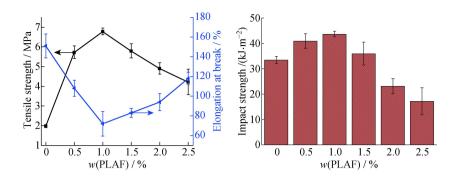


HA-DOCA-His micelles could be stable at physiological (pH=7.4) and tumor extracellular (pH>6.5), and be endocytosed as the micelle entity into tumor cells, then would disintegrate and trigger PTX quick release at endosomal microenvironment ( $pH=5.0\sim6.0$ ).

#### Effect of Polylactic Acid Fiber on the Properties of Thermoplastic Starch Plastics

GUO Bin, ZHA Dong-dong, XUE Can, YIN Peng, LI Pan-xin

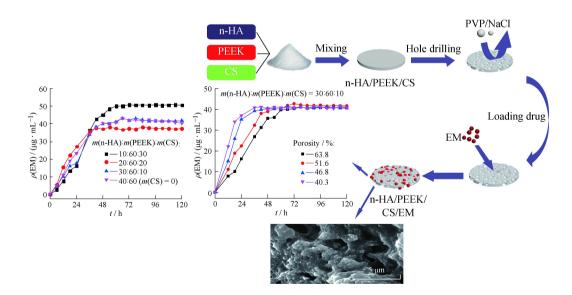
Journal of Functional Polymers, 2018, 31(3):261-266.



Compared with 1.98 MPa and 33.45 kJ/m² of the unreinforced TPS, the maximum values of tensile strength and impact strength of 1.0% PLAF/TPS can reach 6.79 MPa and 43.71 kJ/m² respectively, which due to the uniformly dispersed PLA fibers in the TPS matrix and the interaction between PLA fibers and starch macromolecules.

#### Preparation and Properties of Porous Drug-Delivery n-HA/PEEK/CS Composite Materials

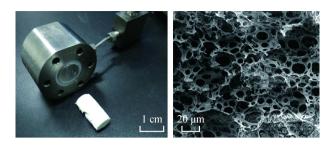
ZHOU Hao-hao, CAO Hui-qun, ZHOU Li, LUO Zhong-kuan, HU Hui-yuan, HUANG Dan-rong Journal of Functional Polymers, 2018, 31(3):267-272.



The novel drug-delivering bone-repair materials were synthesized via forming holes using polyvinylpyrrolidone (PVP) and sodium chloride (NaCl) as the porogen and loading the antibiotic erythromycin (EM) into the n-HA/PEEK/CS composite materials .

# Preparation and Characterization of Starch-Based Macroporous Materials by High Internal Phase Emulsion Templating Method ZHANG Shou-cun, SUN Wu, PI Mao

Journal of Functional Polymers, 2018, 31(3):273-278.

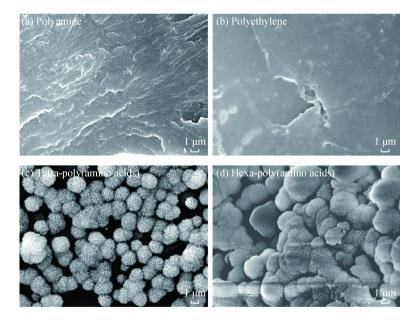


The C/W HIPEs emulsion can be used to produce the macroporous materials with special construction by the emulsion-templating polymerization method. Using this method, a novel starch-based macroporous material was produced.

#### In vitro Bioactivity of Poly (amino acids) Synthesized by Melting Polycondensation

FAN Xiao-xia, YAN Yong-gang

Journal of Functional Polymers, 2018, 31(3):279-284.



The hexa-poly (amino acids) and tetra-poly (amino acids) based on 6-aminohexanoic acid were synthesized by *in-situ* melting polycondensation. Hexa-poly (amino acids) and tetra-poly (amino acids) induced the form of apatite on their surface after soaking into simulated body fluid (SBF) for 5 d, showing good bioactivity. However, polyethylene and polyamide are not bioactive.