

Available online at www.sciencedirect.com

SciVerse ScienceDirect

E**≣≋**₹S

Journal of the European Ceramic Society 32 (2012) 2187-2191

www.elsevier.com/locate/jeurceramsoc

## Novel fabrication of injection-moulded ceramic parts with large section via partially water-debinding method

Wei Liu<sup>a</sup>, Xianfeng Yang<sup>b</sup>, Zhipeng Xie<sup>a,\*</sup>, Cui Jia<sup>c</sup>, Linlin Wang<sup>a</sup>

<sup>a</sup> State Key Laboratory of New Ceramics and Fine Processing, Department of Materials Science and Engineering, Tsinghua University, Beijing 100084, PR China <sup>b</sup> College of Physics and Electronic Science, Changsha University of Science & Technology, Changsha 410114, PR China

<sup>c</sup> The Palace Museum, Beijing 100009, PR China

Received 25 November 2011; received in revised form 26 February 2012; accepted 1 March 2012 Available online 26 March 2012

## Abstract

In this work, we adopt a combination of low molecular weight PEG (L-PEG) and high molecular weight PEG (H-PEG) as water-soluble binder to fabricate injection-moulded ceramic parts with large section. The mechanism of the combination of PEGs removal was proposed for the first time. Defect-free near gear parts with large-sized-section (thickness – 16 mm) were successfully fabricated through water extraction (15 h) followed by rapid thermal pyrolysis (4.5 h). It solves the difficulty of fabricating injection-moulded ceramic parts with large section and our approach is energy saving and high-efficiency as compared with conventional thermal debinding. The results demonstrate that our approach of partially water-debinding followed by rapid thermal pyrolysis could solve the problems of conventional thermal debinding, providing an effective route for the production of injection moulded ceramic parts with large section.

© 2012 Elsevier Ltd. All rights reserved.

Keywords: Injection moulding; Shaping; Defects; PEGs

## 1. Introduction

In recent years, ceramic injection moulding (CIM) has been recognised as a cost-effective process for the production of precise and complex-shaped ceramics with high performance in massive form.<sup>1–4</sup> This method usually comprises of four basic stages: mixing, moulding, debinding and sintering. Among the debinding methods, thermal debinding was the first process developed and is still widely used in the CIM industry because of its simplicity and the low investment of equipment.<sup>4</sup> However, the process is time-consuming because the debinding rate must be slow in order to avoid internal pressure buildup from decomposed binders, which causes cracking, blistering and exfoliation during debinding.<sup>5–7</sup> In addition, the successful manufacturing of injection-moulded ceramics by thermal debinding is only appropriate for components of relatively small cross-section

0955-2219/\$ – see front matter © 2012 Elsevier Ltd. All rights reserved. doi:10.1016/j.jeurceramsoc.2012.03.005 (usually $\sim$ 3–5 mm).<sup>8,9</sup> For injection-moulded ceramic parts with large section, defects such as cracking, deformation and blistering still occur quite easily even if long thermal pyrolysis time is adopted. Because the length of mass transfer path for binder removal is proportional to the size of cross section for thermal debinding, it is difficult to fabricate defect-free ceramic parts with large section via thermal debinding.<sup>9</sup> Consequently, in practical production, if the cross section size of the ceramic parts is larger than  $\sim 10$  mm, almost all the manufactories adopt the moulding approaches such as dry pressing, slip casting and cold isostatic pressing. The organic binders used in the above approaches are much less than those used in injection moulding, so binder removal for such approaches is much easier than that for CIM. However, subsequent mechanical processing (grinding, lathing, cutting etc.) after pressing or sintering is necessary for such technologies because unlike CIM, they are not near net shaping technology, and this leads to high cost with brittle and rigid ceramic materials. In addition, as is known to us, the degree of continuous production of the above shaping technology is far away from that of CIM, which also increases the cost of the

<sup>\*</sup> Corresponding author. Tel.: +86 10 6279 9031; fax: +86 10 6277 1160. *E-mail address:* xzp@mail.tsinghua.edu.cn (Z. Xie).

production. Hence, it is necessary to seek a feasible debinding route to fabricate large-section-sized ceramic parts via injection moulding.

Partially water-debinding followed by rapid thermal pyrolysis appeared as a good alternative, for it presents both high efficiency and environmental acceptability.<sup>10</sup> For this approach, water-soluble binder is firstly removed while the interconnected pore channels are formed from exterior to interior, leaving the insoluble binders in the contact region and the pore channels could serve as escape paths for decomposed gas during subsequent thermal debinding for insoluble binders.<sup>11</sup> The binder system for water debinding mainly consists of water soluble components, which dissolve in water and insoluble backbone binder which keeps the strength of the green body in the whole debinding process.<sup>12</sup> Therefore, binder removal of water-based binder system is much easier than that of wax-based one since the pore channels originated from water-soluble binder could serve as escape paths for the following thermal pyrolysis as indicated above. Therefore, it may be feasible for water debinding to replace thermal debinding to fabricate ceramic parts with large section.

Most previous researches on water-debinding were focused on debinding dynamics, debinding mechanism, optimisation of binder system and microstructure evolution. Yang et al. proposed the solvent debinding mechanism and investigated microstructure evolution for alumina injection-moulded compacts with water soluble binders.<sup>13</sup> Tsai et al. investigated solvent debinding kinetics of alumina green bodies shaped by powder injection moulding.<sup>14</sup> Kim et al. investigated pore structure evolution and binder distribution during both solvent extraction and wicking.<sup>15</sup> Shivashankar and German introduced an effective length scale (the volume to surface area ratio) to understand the effect of the component's shape and dimensions on the critical heating rates for polymer burnout.<sup>16</sup> Yang et al. used the effective length raised by Shivashankar et al. to explore the water-debinding dynamics.<sup>12</sup> In addition, Krauss, Bakan, Park, Liang et al. have done some researches on different water-based binder systems with different kinds of powders.<sup>17-20</sup>. However, the practical production of injection-moulded ceramic parts with large section via partially water-debinding has seldom been reported and researched.

In this study, we adopt a combination of low molecular weight PEG (L-PEG) and high molecular weight PEG (H-PEG) as water-soluble binder to fabricate injection-moulded ceramic parts with large section, which has never been reported. The mechanism of the combination of PEG removal was proposed for the first time. Defect-free near gear parts with large-sized-section (thickness - 16 mm) were successfully fabricated through water extraction (15 h) followed by rapid thermal pyrolysis (4.5 h). It solves the difficulty of fabricating injectionmoulded ceramic parts with large section and the approach is energy saving and high-efficiency as compared with conventional thermal debinding. The results demonstrate that our approach of partially water-debinding followed by rapid thermal pyrolysis could solve the problems of conventional thermal debinding, providing an effective route for the production of injection-moulded ceramic parts with large section.

Table 1	
The used water-based binder system	1.

Components	PEG	PMMA	SA	DBP	Phenothiazine
Weight percentage (%)	66	20	8.5	5	0.5

## 2. Experimental procedure

The ceramic powder used was a commercial zirconia  $(3 \text{ mol}\% \text{ Y}_2\text{O}_3)$  with average particle size  $(d_{50})$  of 0.16 µm and Brunauer-Emmett-Teller specific area of 8 m<sup>2</sup>/g (grade YSZ-F-DM-3.0, Farmeiya Advanced Materials Co., Ltd., Jiujiang, China). The powders were mixed with the binders in a twin screw kneader (SK-160, ShangHai Rubber Machinery, Shanghai, China) at 150–170 °C for a period of 30–45 min. The binders used in water-based binder system for injection moulding in the experiments were: polymethyl methacrylate (PMMA, Plexiglas-8n, Degussa Co., Ltd., Beijing, China), polyethylene glycol (PEG) (Sinopharm Chemical Reagent Beijing Co., Ltd., Beijing, China), stearic acid (SA, Shantou Xilong Chemical Factory, Guangdong, China), dibutyl phthalate (DBP, Beijing Modern Eastern Fine Chemical) and phenothiazine (Sinopharm Chemical Reagent Beijing Co., Ltd., China) with the compositions shown in Table 1. At the same time, injection moulding using wax-based binder system was conducted simultaneously for contrast. The binders used in wax-based binder system for injection moulding were: polypropylene (PP, K8303, Beijing Yanshan Petrochemical Co., Ltd., China), ethylene-vinyl acetate copolymer (EVA, VA content was 14%, Beijing Chemical Factory, China), paraffin wax (Shenyang Paraffin-wax Chemical Co., Ltd., China), stearic acid (SA, Shantou Xilong Chemical Factory, Guangdong, China) and dibutyl phthalate (DBP, Beijing Modern Eastern Fine Chemical) with the compositions shown in Table 2.

The specimens were shaped using an injection moulding machine (JPH30C/E, Qinchuan Hengyi Plastics Machinery Co., Ltd., China). For water-based binder system, debinding was performed in two steps: water leaching to remove the PEG and thermal pyrolysis to remove residual binders. The green (asmoulded) specimens were immersed in distilled water which was held at 40 °C for the first stage. The second stage involved a thermal treatment. Because the pore nets first formed as a result of water debinding, heating rate of the second stage could be quite fast. For wax-based binder system, the as-prepared compact was debound for 48 h in air atmosphere in a muffle furnace.

After debinding, the compacts were finally sintered at a rate of 4 °C/min to 1000 °C and held there for 1 h and then sintered at a rate of 2 °C/min to 1500 °C and held for 2 h to finish sintering.

Table 2The used wax-based binder system.

Components	PP	EVA	PW	SA	DBP
Weight percentage (%)	10	10	60	15	5

Download English Version:

https://daneshyari.com/en/article/1475291

Download Persian Version:

https://daneshyari.com/article/1475291

Daneshyari.com