Porous metal is a lightweight and porous material that has been developed through advancements in material preparation and processing technology. This innovative material combines the advantages of both structural and functional materials, exhibiting exceptional qualities such as high thermal conductivity, resistance to extreme temperatures, porosity, a large surface area, and the ability to customize its porous structure (Lakhdar et al., 2021; Wang et al., 2013; Xu et al., 2016; Li et al., 2020). In China’s economy, the iron and steel industry play a significant role as a foundation sector that is promising for potential applications in filtration, catalysis, and chemical reactions. To meet the demand for efficient filtration and separation matrices, there is a need for innovative designs in porous metal materials. The development of gradient porous materials with superior separation matrices can greatly improve the overall quality of life (Zhu et al., 2022; Adeniran et al., 2017; Hashe and Jen, 2020).

By utilizing porous materials made from Ni-Cr-Mo-Cu alloys, it is possible to improve air filtration efficiency and effectively remove harmful substances. The porous structure of these materials allows for pollutants to be trapped and retained, preventing their release into the atmosphere. Additionally, their high thermal conductivity allows for efficient heat transfer, contributing to their excellent filtration performance. Furthermore, the ability to customize the pore structure of these materials opens numerous possibilities for tailoring their filtration capabilities to specific requirements (Umamungu et al., 2022). Moreover, they have the potential to achieve precise control over particle size and distribution, making them ideal for use in air and water filtration systems, as well as in separating various substances.

In recent times, researchers have been exploring various techniques for creating porous materials with improved filtration and separation properties. One approach is to utilize advanced manufacturing techniques such as 3D printing and electrodeposition (Ameta et al., 2022). These methods allow for accurate control over the design and fabrication of the porous structures, enabling the creation of materials with customized properties. Another method involves the integration of functional additives into the porous metal matrix. These additives enhance the filtering efficiency by promoting specific interactions with the target substances. For instance, the addition of nanoparticles with catalytic properties can effectively eliminate harmful pollutants by facilitating chemical reactions. To meet the demand for efficient filtration and separation materials, there is a need for innovative designs in porous metal materials. The development of gradient porous materials with superior filtering quality holds great promise (Li et al., 2015).
Ti-Al gradient porous materials were effectively created by He et al. utilizing high purity Al powder as the matrix and mixed Ti and Al powder as the gradient layer (He et al., 2010). It had higher filtering accuracy and permeability compared to pure porous aluminum or pure porous titanium materials. However, pore clogging between the matrix and gradient layer is important because of the significant variation in particle size between the Al powder and the Ti-Al mixed powder, which lowers permeability and filtering precision. Ti powder and high purity quartz sheet were used as the raw materials in Liu et al. successful in-situ reaction technique preparation of Ti-TiSi2 gradient composite porous material (Liu et al., 2022).

The findings shown that the maximum pore diameters of the matrix and gradient porous films produced by fine powder sintered at 1050°C under 30 kPa pressure are 7.9 μm and 6.4 μm, and the relative air permeability coefficients of matrix and film are 56.6 m³·m⁻²·h⁻¹·KPa⁻¹ and 94.4 m³·m⁻²·h⁻¹·KPa⁻¹. The maximum pore size difference between the porous film and the matrix is negligible, but the relative air permeability coefficient, which is brought on by the reduction in porous film thickness, drops quickly. Researchers have been quite concerned about the Ni-Cr-Mo-Cu alloy porosity wear resistance and ability to do filtration tasks in highly corrosive settings. In a Ni-Cr-Mo-Cu alloy, the Cr element can create a Cr₂O₃ oxide coating according to (Wen et al., 2022). Ni-Cr-Mo-Cu alloy corrosion resistance may be enhanced by this. However, there are not many publications on how well Ni-Cr-Mo-Cu porous materials filter.

In this paper, Ni, Cr, Mo, Cu powder was used as raw material to prepare porous material matrix by activation reaction sintering, and then Ni, Cr, Mo and Cu with larger particle size were painted on the surface of the matrix, and a gradient degree Ni-Cr-Mo-Cu porous material was obtained by vacuum sintering. Ni-Cr-Mo-Cu porous material with first gradient pore size after sintering was used as raw material, and Ni-Cr-Mo-Cu porous material with second gradient pore size was obtained by painting the surface with fine particle size Ni, Cr, Mo and Cu. The data of element distribution and pore structure were obtained by characterization.

2. EXPERIMENT

The purity is 99.9%, the particle size is 38 – 74 μm, and the powder mass ratio is w (Ni) : w (Cr) : w (Mo) : w (Cu) = 55:30:10:5 of each element powder, through the roller ball mill evenly mixed 48 hours after removal, (Zou et al., Cr at 30%, porous Ni-Cr-Mo-Cu material pore structure is the best), and then the mixed powder into the mold, with a pressure of 60MPa pressed to form a Φ25 mm raw compact (Xide et al., 2021). By maintaining heat for two hours in a 1100°C vacuum sintering furnace, a porous material matrix made of Ni-Cr-Mo-Cu was produced.

**Figure 1:** Sintering process

Ni, Cr, Mo, and Cu powders with a purity of 99.9% and a particle size of 20 μm were employed as the raw ingredients for the gradient porous film, adopting the same ratio as the matrix. To prepare, 1% stearic acid and alcohol were added to the uniformly mixed powder. It was evenly coated on the surface of the porous support body with a soft brush. After sintering at 1100°C and holding for 2 h, the first-order gradient porous material was obtained and labeled as #1 sample (Jin et al., 2023; Shi et al., 2022; Li et al., 2020; Alcaraz et al., 2023). Then, sample #1 was selected as the original matrix, the element powder mixture with particle size of 7 μm was brushed, and the second-order gradient pore size porous material was obtained after vacuum sintering labeled as sample #2. The sintering process is shown in Figure 1. 150°C for 30 minutes is to remove impurities such as water vapor, the final sintering temperature is 900°C, 200 degrees lower than the matrix sintering temperature is because of the use of ultra-fine-grained powder, the temperature is too high will lead to sample sintering densification and even make the pores clogged. According to GB 5164-85, the porosity of the porous material samples was measured, and the phase composition and pore structure of the graded Ni-Cr-Mo-Cu porous material were characterized by X-Ray Diffraction (XRD), Scanning Electron Microscope (SEM) and other testing methods.

3. RESULTS AND DISCUSSION

3.1 Phase Analysis

Figure 2 shows the XRD patterns of Ni-Cr-Mo-Cu porous material matrix and #1 and #2 samples. As shown in Figure 1, the phase composition of the matrix after vacuum sintering is Ni, NiCu and Cr₁₂Ni₂₈₃. The diffraction peaks of #1 and #2 samples do not shift, indicating that their phase composition is the same.

3.2 First Order Gradient Pore Size Ni-Cr-Mo-Cu Porous Material

The SEM surface and side morphology of the first-gradient Ni-Cr-Mo-Cu porous material are shown in Figure 3.

**Figure 3:** #1 SEM images of sample: (a) Surface; (b) Cross-section

It can be seen from Figure 3 (a) that the film thickness of the #1 sample is uniform and the pores are abundant. The maximum pore size of the material measured by the instrument is 10.4 μm, and the permeability is 83.7 m³·m⁻²·h⁻¹·KPa⁻¹. This lays a good foundation for the gradient pore size porous materials to have good filtration. It can be seen from Figure 3 (b) that the film layer of sample #1 has a good bond with the support body, and the film thickness is uniform.

3.3 Second Order Gradient Pore Size Ni-Cr-Mo-Cu Porous Materials

In order to systematically study the effect of different particle size differences on gradient porous films, Ni, Cr, Mo, Cu powder with raw material particle size of 20 μm and 7 μm were used to prepare a gradient porous film and a second gradient porous film, respectively. After testing, the pore structure parameters of the gradient pore size Ni-Cr-Mo-Cu porous material are listed in Table 1.

<table>
<thead>
<tr>
<th>Sample</th>
<th>Permeability m³·m⁻²·h⁻¹·KPa⁻¹</th>
<th>Maximum pore size / μm</th>
</tr>
</thead>
<tbody>
<tr>
<td>matrix</td>
<td>96.3</td>
<td>13.3</td>
</tr>
<tr>
<td>#1</td>
<td>83.7</td>
<td>10.4</td>
</tr>
<tr>
<td>#2</td>
<td>78.9</td>
<td>7.2</td>
</tr>
</tbody>
</table>
It can be seen from Table 1 that compared with the support body, the permeability and maximum pore size of the 1# sample showed a small change. The permeability decreased by 13.1% from 96.3 m⁻¹·h⁻¹·KPa⁻¹ to 83.7 m⁻¹·h⁻¹·KPa⁻¹, and the maximum pore size decreased by 10.4 μm. The permeability of 2# sample decreased by 19.1% compared with the matrix, but the filtration accuracy was greatly improved, and the maximum pore size was maintained at 7.2 μm. According to the analysis, the first gradient degree film plays a key role in the preparation of second-order gradient pore size porous materials. Figure 4 shows the SEM plane and side morphology of 2#.

![Figure 4: 2# SEM morphology of sample: (a) Surface; (b) Cross-section](image)

It can be seen from Figure 4 (a) that the pore size is significantly smaller than that of the first-gradient porous material, the distribution between pores is tight, and the pore structure is incomplete. It can be seen from Figure 4 (b) that the second order gradient pore size porous material consists of a matrix and two membrane layers, and the bond between the membrane layer and the membrane layer is good. It can be seen that the pore size decreases from left to right, indicating that the second-order gradient pore size porous material has a good gradient film structure.

Figure 5 shows the line scanning energy spectrum of porous materials with second-order gradient pore size.

![Figure 5: (a)2# sample linear scanning spectrum diagram; (b) 2# sample Cross-section](image)

From figure 5 (a), it can be seen that the content of Mo element at the boundary increases sharply, which may be caused by the rapid diffusion of Mo element at the boundary, while the content of elements such as Ni Cr Mo Cu in the middle part drops to close to 0 scale, which may be because the element here is a hole and cannot be scanned. It can be seen that the film boundary is clear, the four elements of Ni, Cr, Mo and Cu change alternately, and the changes of the four elements are basically in line with the structure of the film layer, and the content of the four elements is roughly the same as that of the element powder in Figure 5 (b).

Figure 6 shows the profile morphology and energy spectrum analysis diagram of the second-order gradient pore size porous material.

![Figure 6: (a)2#The cross-section morphology; (b) The Element content diagram at position 1; (c) The Element content diagram at position 2; (d) The Element content diagram at position 3](image)

Figure 6 (a) shows the cross-section of the sample. Figure 6 (b), (c), and (d) represent the energy spectrum analysis diagrams of positions 1, 2, and 3 in the figure respectively. The concentrations of Ni Cr Mo Cu at site 1 were 54.13%, 29.86%, 11.24% and 5.07%, respectively. The concentrations of Ni Cr Mo Cu at site 2 were 53.62%, 29.86%, 11.38% and 5.14%, respectively. The concentrations of Ni Cr Mo Cu at site 3 were 53.38%, 29.27%, 12.32% and 5.03%, respectively. It can be seen from Figure 6 that the pore size of the second order gradient porous material is distributed in a gradient manner, and the results of energy spectrum analysis are roughly the same as the test results of the 1# sample. The difference of powder content in these three places is very small, indicating the uniformity of element distribution.

4. CONCLUSION

Ni-Cr-Mo-Cu porous matrix was prepared from Ni, Cr, Mo and Cu powders by activation reaction sintering method. The porous material with second order gradient pore size was obtained by coating the powders with different sizes. The following conclusions are reached.

A gradient pore size porous material was prepared by brushing coating. Ni, Cr, Mo and Cu powder with particle size of 20 μm were used as raw materials for gradient porous film layer. The gradient pore size porous material was obtained by vacuum sintering at 1100°C and held for 2 h. The membrane structure is complete, and the maximum pore size is 10.4 μm. The permeability is 85.7 m⁻¹·h⁻¹·KPa⁻¹.

Using 20 microns Ni, Cr, Mo, Cu mixed powder modified matrix as a transition layer, 7 microns of Ni, Cr, Mo, Cu mixed powder preparation of the second order gradient pore size as surface film layer Ni, Cr, Mo, Cu porous materials, film has good integrity, zero defect, membrane between layers. That can be observed clearly metallurgical combination, With good bonding strength, the filtration accuracy is improved, the maximum pore size is 7.2 μm, and the permeability is 78.9 m⁻¹·h⁻¹·KPa⁻¹.

Due to the excellent filtration performance of second order gradient pore size Ni-Cr-Mo-Cu porous materials, it provides hope for industrial development and practical application, and provides ideas for the filtration direction of metal and ceramic porous materials.

REFERENCES


